Adolescent medicine and general pediatrics I
Concurrent session
12:45 PM
Thursday, January 20, 2022

#1 QUALITY IMPROVEMENT INCREASES COMMUNITY HOSPITAL SMOKE EXPOSURE SCREENING, EDUCATION, AND REFERRAL

Daniel*, Hamline. Sutter Health, Sacramento, CA; UC Davis Children’s Hospital, Sacramento, CA; Adventist Health Lodi Memorial, Lodi, CA

Purpose of Study Tobacco use starts young and is the leading cause of preventable disease, disability, and death in the United States. Secondhand smoke increases ear and respiratory infections, asthma attacks, and risk of Sudden Unexpected Infant Death. Few smoking cessation studies in inpatient pediatrics are formal quality improvement projects and most are at academic institutions. We sought to increase smoke exposure screening, smoking cessation education, and referrals in our community hospital pediatric population. By improving screening and documentation, we anticipate increased provider awareness and smoking cessation interventions.

Methods Used All pediatric ward, newborn nursery, and Level II nursery admissions were eligible. Interventions were education on smoke exposure screening and Helpline referrals, standardizing documentation for screening and discharge instructions, visual reminders, and Helpline wallet cards.

The primary outcome measure was monthly percentage of pediatric inpatients screened for smoke exposure. Secondary outcomes were percentage of pediatric inpatients screening positive for smoke exposure who received discharge instructions or who received a Helpline referral (self or family member). Length of stay (LOS) was monitored as a balancing measure.

Outcome measures were analyzed with statistical process control in SPC for Excel. Baseline and intervention periods for LOS were compared with t-tests.

Summary of Results We increased baseline average smoke exposure screening rates from 14% to 73%, meeting criteria for special cause variation (figure 1). Education on smoke exposure avoidance increased from 5% to 57%. Helpline referrals increased from 0% to 21%. There was no significant change in length of stay.

Conclusions Pediatrician-led smoking cessation interventions are feasible and effective in community hospital pediatric units with no significant impact on length of stay.

#2 PILOMATRICOMA IN A CHILD WITH TURNER SYNDROME: A RARE ENTITY

Horowitz, Tappin*. Valley Children’s Healthcare, Madera, CA

Purpose of Study Pilomatricoma is a rare, skin neoplasm that is often confused with dermoid or brachial cleft cysts. Julian

Abstract #1 Figure 1 Vertical lines are timing of interventions. 1) Monthly Pediatrician education started 2) EMR documentation standardized, visual reminders posted 3) Helpline wallet cards available.

UCL= Upper Control Limit, Avg= Average, LCL= Lower Control Limit
et al., reported that pilomatricomas are commonly misdiagnosed pre-operatively in up to 75% of cases.

We report the case of a child with Turner’s syndrome with a pilomatricoma that was diagnosed on biopsy. We review the histopathologic features and emphasize its association with Turner’s syndrome.

Methods Used Case Report.

Summary of Results 2 year old female with Turner Syndrome presented with a progressive mass above her right upper lip for 6 months.

On exam, she was well-appearing with phenotypical features of Turner’s syndrome. A 0.5 x 0.5 cm erythematous, verrucous, well circumscribed, nontender, mobile lesion was noted above her lip.

She underwent complete excision of the mass without complication. Excisional biopsy revealed the presence of viable basaloïd cells and shadow cells confirming the diagnosis of pilomatricoma.

Conclusions A pilomatricoma, otherwise known as pilomatrixoma, are benign subepidermal tumors of the hair follicle matrix. The lesions occur on the face and neck with a mean age of onset between 5.8-7 years old.

Lesions are usually asymptomatic but, inflammation and ulceration can occur. The most common clinical presentation is a firm, subcutaneous lesion with an irregular surface. The overlying skin may be red, blue, or display the tent sign. Studies have reported the initial development of pilomatricomas in children with Turner’s syndrome. The exact cause of this is unknown although animal studies suggest a genetic component.

Histopathologically, a pilomatricoma appears as a mass composed of viable basaloïd cells, shadow cells, calcification, and ossification. The mainstay of treatment of a pilomatricoma is complete surgical excision as the lesions do not regress spontaneously. Early excision within 12 months of diagnosis is associated with better cosmetic outcomes. Recurrence and malignant transformation is rare.

Abstract #2 Figure 1

This case highlights the importance of considering pilomatricoma as a cause of solitary skin nodules, especially when on the head, neck or upper extremities. Additionally, physicians caring for children with Turner syndrome should be aware of the prevalence of pilomatricoma in this population.

#3 MENTAL HEALTH HOPSCOTCH: IMPROVING ADOLESCENT MENTAL HEALTH ONE HOP AT A TIME

1AS Rodriguez*, 1M Flores, 1T Fildes, 1J Charbonnet, 1K Colwell, 1R Kinman. 1Fresno High School, Fresno, CA; 1University of California San Francisco Fresno, Fresno, CA

10.1136/jim-2022-WRMC.3

Purpose of Study Fresno High Women’s Alliance students continue to collaborate with UCSF Fresno pediatrics to create community action research projects on topics of adolescent concern using a ‘youth as partner’ approach. Given the social isolation and increasing depression students noticed amongst themselves and their peers due to COVID19, the Women’s Alliance teens chose to focus this last year on improving mental health amongst their peers. Mental Health Hopscotch was chosen for its simplicity and ease of use. The fact that it was created by an adolescent in response to the COVID19 pandemic provided further impetus for its use.

Methods Used Students collaborated with the school’s Social Emotional Wellness and Support team, choosing to do their mental health intervention during National Mental Health Month. They created a Mental Health ‘Sunshine’ at their school entrance, chalk- ing positive affirmations in a sun-shaped diagram for all to see, and chalked a Mental Health Hopscotch on the sidewalk in the front of the school. Silicone bracelets with motivational quotes and mental health awareness pencils, stickers, and mini buttons were handed out to those who completed the Hopscotch. A QR code linked to Google Forms was used to survey students who completed the Hopscotch.

Summary of Results 42 students were surveyed. 12% of students reported their average stress level was ‘just right’, 38% reported they could ‘handle’ their stress, 21% felt that they were ‘getting stressed’, 19% reported they were ‘starting to lose it’, and 10% described their stress as ‘getting out of control’. Students primarily dealt with stress by listening to music (31%), exercising (19%), and sleeping (14%). Half of students surveyed reported difficulty sleeping at night, while 90% of students felt that doing Mental Health Hopscotch helped boost their mood.

Conclusions Although only a limited number of students were surveyed due to restricted numbers of students present on campus, the majority of students felt stressed with half the students reporting difficulty sleeping at night. Mental Health Hopscotch provided a simple, quick, yet no-cost approach to boost students’ mood, thus empowering teens concerned about the mental health of their peers to stage a mental health intervention on their own school grounds.

#4 DAYLIGHT SAVINGS AND PEDIATRIC EMERGENCIES

1,2Y Wong*, 1,2R Enarson, 1,2J Lee. 1The University of British Columbia Faculty of Medicine, Vancouver, BC, Canada; 2BC Children’s Hospital, Vancouver, BC, Canada

10.1136/jim-2022-WRMC.4

Purpose of Study Daylight Savings Time (DST) is a biannual time change where during ‘spring forward,’ clocks are set forward one hour, potentially resulting in sleep deprivation for much of society. During ‘fall back,’ the opposite occurs. Circadian rhythm disruption has been shown to have effects on
Abstract #4 Figure 1

cardiovascular, neuropsychiatric, metabolic, immune-related and accidental events in adults. A 2018 study showed increased emergency department (ED) visits for adults in the time after DST. These findings have not been verified in pediatric populations and if extant, may have implications on managing ED patient volume and expectations. We hypothesized that the large-scale sleep deprivation following spring time change would result in increased ED presentations, particularly among certain presentations (neurologic, psychiatric, accidental/traumatic) that may be especially susceptible to sleep deprivation, and that the fall time change would have an opposite effect.

Methods Used We retrospectively collected and analyzed the primary medical complaint of all children (0–16 years) presenting to BC Children’s Hospital ED in the 2 weeks before and 3 weeks after the biannual DST time changes during 2011 to 2019. Incidence ratios (IR) of ED presentations were calculated over day 0 (day of time change) to day 7. IRs were calculated for all presentations and broken down by specialty.

Summary of Results After excluding infectious presentations, the IR was increased during the first week following spring time change: Monday by 6%, Tuesday by 7% and Wednesday by 6%, though the results were statistical insignificant (p>0.05). There were significant decreases (p<0.05) after the fall time change: Monday by 12%, Tuesday by 13% and Wednesday by 8%.

Conclusions Following the seasonal time change in the spring, there were increases in IR, though this did not reach statistical significance. There were significant decreases in IR during the three days following the fall shift. Together these findings suggest that the widespread sleep deprivation at spring time change results in adverse health effects among children, while extra sleep in the fall time change may be protective.

Circadian rhythm disruption from DST in the children shows potentially important effects on emergency visits and further study can lead to better patient care and ED preparation. This may be informative in developing policy regarding the need for DST.

Abstract #5

THE RELATIONSHIP BETWEEN SELF-PERCEIVED HEALTH STATUS AND CHRONIC CONDITION, ACCESS TO CARE, AND AWARENESS OF CHRONIC CONDITION AMONG ADULTS WITH HISTORY OF ADVERSE CHILDHOOD EXPERIENCES

L Trinh*, S Teklehaymanot, M Shaheen. Charles Drew University of Medicine and Science, Los Angeles, CA

10.1136/jim-2022-WRMC.5

Purpose of Study Adults with a history of adverse childhood experiences (ACEs) are at increased risk for chronic disease and, thus, poor health outcomes. Yet, the effect of chronic disease awareness on health outcomes in adults with ACEs has not been examined. The objective of this study was to determine the relationship between general health status and chronic disease, access to care, and awareness of chronic condition among adults with history of ACEs.

Methods Used Data from the 2019 Behavioral Risk Factor Surveillance System were analyzed. Descriptive statistics were used to determine the prevalence of ACEs, chronic disease, healthcare access, chronic disease awareness, general health status, and population characteristics. Bivariate analysis using Chi-squared test was performed on history of ACEs by all independent variables and general health status by all independent variables. Multivariable logistics regression was used to determine the relationship between general health status and ACEs, adjusting for chronic disease, healthcare access, chronic disease awareness, and demographics.

Summary of Results Of the 78,112 respondents, 63% reported being exposed to at least one ACE. History of ACEs was associated with high prevalence of chronic diseases (p<0.002), lower healthcare coverage (p<0.0001), lower chronic disease awareness (p<0.006), and report of fair/poor general health status (p<0.0001). In adjusted analyses, adults with 2, 3, and ≥4 ACE events had 1.24, 1.22, and 1.45 times increased odds of reporting fair/poor health status compared to those reporting no ACEs.

Conclusions Adults with history of ACEs face barriers to achieving good health. There is a need to expand ACEs’ screening in primary settings so that early intervention can improve general health outcomes.
outpatient (n=35) populations. Inpatients were more likely to experience Social Connection risk (p=0.046), Financial risk (p=0.0006), and Food Insecurity (p=0.0077) than outpatients.

When comparing between patients living in the upper quartile (n=40) and the lower three quartiles (n=34) of SVI, there was little difference in BMI, Financial risk, Food Insecurity risk, Transportation risk, Physical Activity risk and Stress risk. There was similarly little difference when comparing age groups of younger versus older children.

Conclusions There are significant disparities in risk between pediatric inpatient and outpatient populations. Steps could be taken to identify inpatients with reduced resources to improve food insecurity, social connection risk, and financial risk. Although SVI is important for understanding the context of each patient, every family has unique social determinants and risks that could be addressed by physicians.

#7 SUBJECTIVE SOCIAL STATUS AND FOOD INSECURITY RELATIONSHIP WITH WEIGHT-SPECIFIC QUALITY OF LIFE AMONG LATINO ADOLESCENTS AND BETWEEN SEXES


Purpose of Study Social status and food insecurity (FiS) may contribute to health disparities among youth. This study aimed to evaluate whether subjective social status (SSS) and FiS are associated with weight-specific quality of life (wQoL) among Latino youth with obesity. We further explored whether relationships differed by sex.

Methods Used One-hundred and forty-one Latino youth (47% male; mean age: 15.3 ± 0.9) with obesity completed surveys to assess SSS, perceived FiS, and wQoL (self, social, and environment). Separate linear regression models were performed to examine the relationship between SSS & FiS with wQoL, after controlling for sex and BMI percentile. Data were then stratified by sex to determine Pearson’s r for wQoL & SSS and wQoL & FiS.

Summary of Results Mean total (64.1±24.9), self (57.3±29.3), social (69.5±25.6), and environmental (60.5±26.3) wQoL, with males reporting higher total, self, and environmental wQoL as compared to females (all p<0.05) after controlling for BMI. Over one-third of the cohort indicated very low food insecurity (36.2%) or marginal food insecurity (34%), which did not differ by sex. Despite no sex differences in SSS society (mean diff=-0.074, p=0.77) or SSS school (mean diff=-0.354, p=0.28), there was a positive relationship between SSS school and all scales of wQoL regardless of sex (all p<0.01). SSS school was a significant predictor of total wQoL (β=4.24, p<0.001) and the self (β=3.49, p=0.008), social (β=4.40, p<0.001), and environment (β=4.57, p<0.001) subscales. SSS school explained 9% of the variance in total, social, and environment wQoL, and 4% of the variance in self wQoL. There was an inverse relationship between FiS and all scales of wQoL, particularly for those experiencing marginal and high FiS (all p<0.03). This correlation was stronger for males for all wQoL scales at all levels of FiS except high FiS. Marginal FiS was a significant predictor of total (β=-12.94, p=0.006), self (β=-12.48, p=0.029), social (β=-12.06, p=0.013), and environment (β=-14.66, p=0.003) wQoL after controlling for sex and BMI percentile. High FiS was a significant predictor of total (β=-22.46, p<0.001), self (β=-18.56, p=0.016), social (β=-24.16, p<0.001), and environment (β=-22.76, p<0.001) wQoL after controlling for sex and BMI percentile. Marginal and high FiS explained 8% of the variance in total, social, environmental wQoL, and 3% of the variance in self wQoL.

Conclusions Among Latino youth with obesity, social status is associated with wQoL while food insecurity is inversely associated with wQoL.
Purpose of Study The prevalence of pediatric obesity continues to increase. In Nevada, approximately 40% of youth are overweight or obese and 70% of these children will remain overweight as adults. Traditional medical school curriculum does not adequately prepare students to counsel families on this subject. The purpose of this study was to evaluate the progression of third year medical student knowledge base involved in managing obesity in children and adolescents over the course of the pediatric clinical curriculum.

Methods Used 63 medical students in the third clinical clerkship curriculum at the University of Nevada, Reno School of Medicine were given a survey to evaluate their knowledge on pediatric obesity and perceptions surrounding treatment before a six-week pediatric clerkship. During the clerkship, students received instruction on the diagnosis and treatment of obesity. At the conclusion of the clerkship, students took a post-survey to assess knowledge gained during the experience. Paired samples T-test and Chi-square tests were used to assess differences between pre- and post-surveys.

Summary of Results After the clerkship, there was an increase in the mean score in knowledge and comfort level in recommending a treatment program for overweight/obese children between pre- (M=1.97, SD=0.91) and post-tests (M=3.55, SD=0.89) (t(62)=10.25, p<0.0001). There was also an increase in the mean score in their ability to effectively counsel overweight/obese children between pre- (M=2.40, SD=0.90) and post-tests (M=3.86, SD=0.77) (t(62)=10.33, p<0.0001). In addition, there was an increase in students believing in the overall efficacy of counseling in the treatment for overweight/obese children and adults between the pre- (M=2.91, SD=0.75) and post-tests (M=3.34, SD=0.76) (t(62) =3.23, p=0.0019).

Conclusions The curriculum improved student knowledge and understanding of pediatric obesity. In particular, perceived comfort and ability to counsel patients and families about obesity prevention and treatment increased. As obesity continues to be a challenge, expansion of medical student education in this area is imperative to address this problem.

Abstract #9 Table 1

<table>
<thead>
<tr>
<th>Patients Transplanted During COVID Pandemic (n=50)</th>
<th>Patients Transplanted Prior to COVID Pandemic (n=482)</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-Year Survival</td>
<td>94.0%</td>
<td>90.5%</td>
</tr>
<tr>
<td>1-Year Freedom from CAV</td>
<td>100.0%</td>
<td>96.5%</td>
</tr>
<tr>
<td>1-Year Freedom from NF-MACE</td>
<td>98.0%</td>
<td>86.9%</td>
</tr>
<tr>
<td>1-Year Freedom from Any Rejection</td>
<td>90.0%</td>
<td>84.2%</td>
</tr>
<tr>
<td>1-Year Freedom from Cellulare Rejection</td>
<td>90.0%</td>
<td>92.1%</td>
</tr>
<tr>
<td>1-Year Freedom from Antibody-Mediated Rejection</td>
<td>94.0%</td>
<td>94.8%</td>
</tr>
</tbody>
</table>

Cardiovascular I
Concurrent session
12:45 PM
Thursday, January 20, 2022
Conclusions Despite necessary changes being made to post-transplant care to mitigate the spread of COVID-19 and protect an immunosuppressed population, heart transplantation during the COVID-19 pandemic appears safe with 1-year outcomes comparable to years prior.

#10 COVID PANDEMIC AND SOCIAL MITIGATIONS IMPROVE OUTCOMES FOR HEART TRANSPLANT PATIENTS
C Washington, S Kim*, N Patel, T Singer-Englar, M Hamilton, J Kobashigawa. Cedars-Sinai Smidt Heart Institute, Los Angeles, CA
10.1136/jim-2022-WRMC.10

Purpose of Study The COVID pandemic affected how our medical staff treated our heart transplant (HTx) patients during this period of time. Patients were seen virtually via telemedicine and patients self-isolated at home. What we do not know is the impact of this treatment during the COVID pandemic on HTx outcomes. As patients were self-isolating, it is possible that medication and medical compliance were increased and there may have been a decrease in non-COVID infection rates as exposure was minimized due to patients self-isolating. None of these factors have been assessed prior and, thus, we reviewed our large, single center patient population for this study.

Methods Used Between March 2020 and September 2020, we assessed 55 HTx patients who were transplanted during this period of time and followed for 6 months. Patients were self-isolating and had every other clinic visit changed to a virtual visit to minimize exposure to COVID. Endpoints for this study included 6-month survival, re-hospitalization, number of non-COVID infections (defined as the need for intravenous antibiotics), any treated rejection (ATR), and maintenance of therapeutic immunosuppressive blood levels. The study patients were then compared to a control group of the previous three years, averaging each year at the same time points of March 13, 2017, March 13, 2018, and March 13, 2019, and followed for 6 months. Each group was averaged and then compared to the study group.

Summary of Results The study group (during the COVID pandemic) demonstrated a significant decrease in re-hospitalization in the first 6 months following HTx compared to the control group. There was a numerical decrease in non-COVID infectious complications. There was no difference in survival and freedom from any-treated rejection episodes between the two groups. Reasons for rehospitalization included infections, various cardiac and renal issues, malaise, and fever.

Conclusions The COVID pandemic demonstrated that self-isolation and virtual visits resulted in fewer hospitalizations possibly due to less infectious complications. This implies that perhaps stricter restrictions for community exposure might benefit HTx patients in the 6 months following transplantation.

Abstract #10 Table 1

<table>
<thead>
<tr>
<th>HTx Patients Transplanted between Mar.-Sep. 2020</th>
<th>HTx Patients Transplanted between Mar.-Sept. 2017-2019</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>(n=55)</td>
<td>(n=169)</td>
<td></td>
</tr>
<tr>
<td>6-Month Survival</td>
<td></td>
<td>0.733</td>
</tr>
<tr>
<td>92.7%</td>
<td>94.1%</td>
<td></td>
</tr>
<tr>
<td>6-Month Freedom</td>
<td></td>
<td>0.981</td>
</tr>
<tr>
<td>90.9%</td>
<td>91.1%</td>
<td></td>
</tr>
<tr>
<td>from ATR</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rehospitalization within 6 Months</td>
<td></td>
<td>0.003</td>
</tr>
<tr>
<td>16.4% (9)</td>
<td>37.9% (64)</td>
<td></td>
</tr>
<tr>
<td>Non-COVID Infection within 6 Months</td>
<td></td>
<td>0.323</td>
</tr>
<tr>
<td>10.9% (6)</td>
<td>18.9% (32)</td>
<td></td>
</tr>
</tbody>
</table>

#11 SEX DISPARITIES IN HEART TRANSPLANT WAITLIST TIME FOLLOWING THE DONOR HEART ALLOCATION POLICY CHANGE

1I Sindha*, 2T Singer-Englar, 5S Kim, 2N Patel, 2M Hamilton, 2J Kobashigawa. 1University of California Los Angeles, Los Angeles, CA; 2Cedars-Sinai Smidt Heart Institute, Los Angeles, CA
10.1136/jim-2022-WRMC.11

Purpose of Study There are many reports in organ transplantation that demonstrate that there are sex discrepancies in waitlist urgent status, time on the heart transplant waitlist, waitlist mortality. There are no differences in men versus women in terms of heart disease and in terms of mortality for these two sexes. It would be expected that both men and women would have similar percentages as urgent status, especially after the new donor heart allocation policy change took place in October 2018. We chose to assess our male and female patients to establish whether there exists a difference in patients listed as urgent status on the HTx waitlist.

Methods Used Between November 2018 and December 2020 (after donor heart allocation change in October 2018), we assessed 276 patients on the HTx waitlist. Patients were followed for 6 months and censored after they were transplanted or removed from the waitlist. Percent of patients of each sex listed as urgent status (status 1, 2, 3) was recorded. Mortality on the waitlist, waitlist time, and removal from the waitlist due to being too sick were secondary endpoints.

Summary of Results After the donor heart allocation policy change in October 2018, women were significantly less likely to be listed as urgent status compared to men. For those patients listed as urgent status, there was no significant difference in mortality for women versus men on the HTx waitlist. The waitlist time was shorter for men compared to women (see table 1).

Conclusions There appears to be a sex disparity for women being less likely to be listed as urgent status on the HTx waitlist. Further studies are needed to determine whether this difference has a biologic mechanism or whether there is selection bias and/or treatment bias present in their care.

Abstract #11 Table 1

<table>
<thead>
<tr>
<th>Endpoints</th>
<th>Female patients (n=73)</th>
<th>Male patients (n=203)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>6-month survival</td>
<td>96.6%</td>
<td>96.6%</td>
<td>0.249</td>
</tr>
<tr>
<td>6-month freedom from severe illness</td>
<td>96.8%</td>
<td>96.1%</td>
<td>0.189</td>
</tr>
<tr>
<td>Urgent status at listing</td>
<td>28.8%</td>
<td>49.3%</td>
<td>0.003</td>
</tr>
<tr>
<td>Urgent status at removal</td>
<td>37.0%</td>
<td>59.1%</td>
<td>0.002</td>
</tr>
<tr>
<td>Average time on waitlist (days)</td>
<td>124.4 ± 164.7</td>
<td>108.3 ± 181.9</td>
<td>0.546</td>
</tr>
</tbody>
</table>
Purpose of Study In heart transplantation (HTx), donor-to-recipient size matching has been done by height and weight. More recently, predicted heart mass (PHM) has been found to be more clinically useful to reflect outcome. Using PHM, it has been demonstrated that under-sizing a donor heart for a larger recipient with high pulmonary artery pressures leads to increased mortality. It has recently been noted in the International Society for Heart and Lung Transplantation (ISHLT) registry that there may be increased risk in placing an oversize donor heart using weight into a smaller recipient. This clinical outcome has not yet been established using PHM. We sought to address this question in our large, single center experience using PHM.

Methods Used Between January 2010 and June 2020, we assessed 586 donor-to-recipient donor heart matches. We used PHM to assess whether there were outcome differences when the donor hearts were oversize. We divided the donor-to-recipient PHM ratio into two categories: normal (90–110%, n=524), and markedly oversize (greater than 140%, n=64). Outcomes included 1-year survival, freedom from 1-year rejection (acute cellular rejection [ACR], antibody-mediated rejection [AMR]), freedom from cardiac allograft vasculopathy (CAG: stenosis ≥30%), freedom from cardiac dysfunction (defined as LVEF ≤40%), and freedom from non-fatal major adverse cardiac events (NF-MACE: myocardial infarction, new congestive heart failure, percutaneous coronary intervention, implantable cardioverter defibrillator/pacemaker implant, stroke).

Summary of Results Markedly oversized donor hearts using PHM compared to normal matching showed no difference in 1-year survival, freedom from 1-year ACR, freedom from CAV, freedom from NF-MACE, and freedom from cardiac dysfunction. There was a significantly lower 1-year freedom from AMR in the markedly oversized donor heart group which is due to more women recipients (sensitized due to previous pregnancies) in this group (71% oversize group vs. 20% normal group).

Abstract #12 Table 1

<table>
<thead>
<tr>
<th></th>
<th>Oversized PHM [&gt;140%] (n=64)</th>
<th>Normal PHM [90–110%] (n=524)</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean PHM</td>
<td>154.86 ± 15.44</td>
<td>99.95 ± 5.53</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>1-Year Survival</td>
<td>93.8%</td>
<td>90.6%</td>
<td>0.420</td>
</tr>
<tr>
<td>1-Year freedom from CAV</td>
<td>96.9%</td>
<td>96.6%</td>
<td>0.852</td>
</tr>
<tr>
<td>1-Year freedom from NF-MACE</td>
<td>95.3%</td>
<td>87.2%</td>
<td>0.085</td>
</tr>
<tr>
<td>1-Year freedom from ACR</td>
<td>92.2%</td>
<td>91.8%</td>
<td>0.899</td>
</tr>
<tr>
<td>1-Year freedom from AMR</td>
<td>82.8%</td>
<td>94.8%</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>1-Year freedom from Right</td>
<td>100.0%</td>
<td>100.0%</td>
<td>-</td>
</tr>
<tr>
<td>Heart Failure</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1-Year freedom from Cardiac</td>
<td>98.4%</td>
<td>99.0%</td>
<td>0.669</td>
</tr>
<tr>
<td>Dysfunction</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1-Year freedom from Pulmonary</td>
<td>93.8%</td>
<td>95.0%</td>
<td>0.706</td>
</tr>
</tbody>
</table>

Conclusions Markedly oversize donor-to-recipient matching using PHM does not result in poor outcomes after heart transplantation. This has potential to expand the donor pool, particularly for smaller patients.
#14 **THE IMPORTANCE OF NATURAL HISTORY DATA COLLECTION AS DEMONSTRATED IN DANON DISEASE ECHOCARDIOGRAM ANALYSIS**

**G Storm**, 1K Boynton, 1A Lamard, 1E Esraghanian, 1K Hong, 1L Mestarini, 1E Alder, 1M Taylor. 1CU SOM, Aurora, CO; 2UCSD, La Jolla, CA

Purpose of Study With over 7,000 rare diseases affecting 1 in 10 Americans, longitudinal data describing the clinical course of rare diseases is essential to understand disease natural history and prepare for studies of novel treatments. Traditionally, published data associated with these diseases is limited to cross-sectional times of diagnosis and major events making inferences on the progression and trajectory of various phenotypes difficult. A prime example is Danon Disease, a rare genetic cardiac disease with a malignant outcome of death or need for heart transplantation in most males.

Methods Used To mitigate the lack of longitudinal natural history knowledge in Danon patients, a retrospective clinical database was developed using the REDCap database program. All current and past medical history for Danon patients enrolled internationally was collected and entered into the database by two research sites. To date, this registry has enrolled over 100 patients, with roughly equal representation of males and females. This includes the collection of data on over 550 echos.

Summary of Results To demonstrate the power of this natural history study, echocardiogram data were collected over time and examined for trends in ejection fraction (EF) of the patients’ hearts over the age during that time of their illness. Data on the eight patients with the largest number of longitudinal EF data points, prior to transplant, were extracted and graphed. The data revealed that EF drops off much earlier, at almost half the age in males compared to females and that substantial variation in EF is present for subjects, rather than a smooth, gradual decline.

Conclusions As this natural history data continues to be analyzed, further analysis will be done that aims to look at the progression of heart disease in Danon patients prior to transplant. These trends will be imperative in not only understanding the progression of the disease to drive best clinical practices, but also to utilize as controls in clinical trials for potential treatments of Danon Disease.

Abstract #14 Figure 1

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#15 **LIFESTYLE COUNSELING FOR VETERANS WITH SYMPTOMATIC HEART FAILURE**

1C Taylor*, 1A Edmeade, 1CE Walters, 2J Patel, 1Loma Linda University School of Medicine, Loma Linda, CA; 2VA Loma Linda Healthcare System, Loma Linda, CA

Purpose of Study Lifestyle counseling has shown to be effective in modifying health behaviors and reducing cardiovascular risk in healthy patients. However, data supporting effectiveness of lifestyle counseling in patients with symptomatic heart failure is limited. This quality improvement study hypothesizes that lifestyle counseling focusing on sleep, activity, nutrition, medication adherence, and self-care will be associated with improvements in health behaviors in veterans with heart failure.

Methods Used This study screened a selected cohort of patients with symptomatic heart failure from the Loma Linda VA Heart Failure Program. The study included 5 counseling sessions over a 9 week interval. In week 1, pre-intervention surveys were completed using the following validated surveys: Pittsburgh Sleep Quality Index, Veterans Specific Activity Questionnaire, Mini Nutritional Assessment, Eight-Item Morisky Medication Adherence Scale, and Self-Care of Heart Failure Index. During weeks 2 through 8, three counseling sessions focusing on nutrition, sleep, exercise, and self-care were tailored to each patient following best practices from LIVHA Heart Failure Program. In week 9, post-intervention surveys were completed.

Summary of Results Out of 112 screened, 49 (44%) agreed to participate. Each patient was given a baseline score during the pre-intervention surveys during week 1. Baseline Pittsburgh Sleep Quality Index average score was 8.74 with post-intervention survey average of 7.68. Lower scores on the Pittsburgh Sleep Quality Index demonstrate higher quality sleep habits and sleep quality. For the Veterans Specific Activity Questionnaire baseline average score was 4.36 and post-intervention was 4.55. This questionnaire ranked activities according to metabolic equivalent of task from 1 to 13. At baseline in the Mini Nutritional Assessment patients reported an average score of 11.38. After lifestyle counseling, patients reported an average score of 12.40. The Mini Nutritional Assessment scoring system had a maximum of 14 with normal nutritional status ranging from 12 to 14, at risk of malnutrition from 8 to 11, and malnourished scores with score under 7. The Eight-Item Morisky Medication Adherence Scale had baseline and post-interventions average scores of 5.51 and 6.30, respectively.

Conclusions All survey data indicated positive changes in lifestyle from pre to post surveys. Lifestyle counseling may improve health behaviors in patients with symptomatic heart failure.
Abstracts

#16 ATRIAL FIBRILLATION, ELECTROCONVULSIVE THERAPY, AND STROKE RISK
M Kapadia*, P Jagadish, M Hutchinson, H Lee. The University of Arizona College of Medicine Tucson, Tucson, AZ
10.1136/jim-2022-WRMC.16

Purpose of Study To review the literature on stroke risk in patients with atrial fibrillation (AF) undergoing electroconvulsive therapy (ECT) as well as anticoagulation recommendations.

Methods Used Two authors independently performed a literature review of PubMed, searching for ‘electroconvulsive therapy and atrial fibrillation.’ The resulting articles and their references were reviewed for relevance to AF and stroke risk.

Summary of Results Rozig, et al. (2018) found that ECT is not associated with an increased risk of new or recurrent stroke. Among 23 studies, we found that post-ECT cardioversion of AF to normal sinus rhythm occurred in 2 cases. Neither was associated with stroke, though this finding may limited by low sample size. However, ECT has induced AF in at least 6 cases without stroke.

ECT requires fewer joules than does cardioversion for atrial fibrillation, and current is applied to the brain for ECT versus the heart for synchronized cardioversion. The mechanism of AF induction/cardioversion is a catecholamine surge and varied hemodynamic changes.

Because of the rarity of stroke in ECT patients, routine anticoagulation prior to ECT is controversial. Furthermore, direct electrical stimulation of the brain risks hemorrhagic stroke after ECT.

In patients with existing AF who are not anticoagulated, an alternative means of reducing cardioversion risk may involve modulation of post-ECT hemodynamic changes by beta-blockade. Beta-blockade may exacerbate bradycardia in some while reducing reflex tachycardia in others and may even decrease in seizure length, thus, lowering the efficacy of ECT.

Conclusions The risk of inducing stroke by cardioversion of AF in the setting of ECT is very low despite a documented risk of cardioversion. Routine anticoagulation prior to ECT remains controversial. Imaging modalities such as echocardiography or mitigation of hemodynamic effects may further reduce the risk of stroke in these patients. Areas for further study are precise assessment of cardioversion and stroke risk in AF patients and the effect of routine beta-blockade on cardioversion risk.

#17 CRITICAL ROLE OF SPECIALIZED PRO-RESOLVING MEDIATORS IN ATHEROSCLEROSIS
S Ranganajan*, P Rangchaikal, MM Radwan. Western University of Health Sciences, Pomona, CA
10.1136/jim-2022-WRMC.17

Purpose of Study Coronary artery disease is the leading cause of death worldwide with over 17.9 million deaths per year according to WHO. There has been a vast amount of research done in understanding the causes, and therefore the treatments of this disease. Our body’s inflammatory processes have been identified as a nidus of the elaborate process that ultimately leads to life-threatening cardiovascular events. However, research around understanding how the body puts an end to such naturally occurring inflammation i.e., resolution of this inflammation, is gaining traction and has shed light into new avenues for future management of CV diseases. In this narrative review we discuss the pathophysiological and molecular mechanisms of atherosclerosis including inflammation, apoptosis and efferocytosis, the recent development in the understanding of a new class of molecules called Specialized Pro-resolving Mediators (SPMs), and the impact of such findings in the realm of cardiovascular treatment options.

Conclusions

Endocrinology and metabolism I
Concurrent session
12:45 PM
Thursday, January 20, 2022

#18 CORONARY ARTERY CALCIFICATION SCORING IN TRANSGENDER INDIVIDUALS
T Duro*, E Choi, P Kapsner. The University of New Mexico, Albuquerque, NM
10.1136/jim-2022-WRMC.18

Purpose of Study To assess risk factors and the prevalence of coronary artery disease (CAD) using coronary artery calcium (CAC) scoring in transgender individuals receiving gender affirming hormone therapy (GAHT).

Transgender individuals are treated with cross sex hormone therapy. These hormones alter metabolic profiles and may be
associated with risk factors for CAD. Little data is available on the atherosclerotic vascular risk due to GAHT in transgender individuals. CAC Scores have been validated as a noninvasive method to assess risk for cardiovascular events in the general population.

Methods Used This is a pilot study assessing feasibility of obtaining baseline risk profiles and CAC scores in transgender patients over the age of 18, in order to establish a baseline risk assessment and the prevalence of CAD in this population. Patients with risk factors other than smoking and family history were excluded. Difference in CAC Scores were compared to those in The Coronary Artery Risk Development in Young Adults (CARDIA) study. Baseline characteristics were compared using t-tests.

Summary of Results Out of 25 transwomen recruited, 24 completed CAC. 3/24 (12.5%) had CAC >0. One had CAC >100. Out of 22 transmen recruited, 16 completed CAC. 2/16 (12.5%) had CAC >0. None had CAC >100. CAC scores did not correlate with the presence of risk factors. Patient data in table 1.

Conclusions Overall, 12.5% of transgender people on GAHT had positive CAC scores. This is similar to the findings of the CARDIA study where the prevalence of positive CAC scores in the general population was 11.7%. There was a higher percentage of current smokers in transgender women who also had higher triglycerides, but a lower LDL. Transmen had a higher family history of CAD, percentage having ever smoked, and a higher hsCRP. This cross-sectional study shows that obtaining CAC scores in transgender patients is feasible and that the prevalence of positive CAC scores appears to be similar to that of the general population. A larger, longitudinal study will be performed to expand on these findings.

Abstract #18 Table 1 Baseline characteristics

<table>
<thead>
<tr>
<th></th>
<th>Transwomen n=25</th>
<th>Transmen n=22</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>40 (14)</td>
<td>38 (11)</td>
</tr>
<tr>
<td>History of smoking</td>
<td>14 (56%)</td>
<td>16 (73%)</td>
</tr>
<tr>
<td>Current smoker</td>
<td>7 (28%)</td>
<td>2 (9%)</td>
</tr>
<tr>
<td>GAHT Duration, yr</td>
<td>3.2 (11.9)</td>
<td>5.5 (10.5)</td>
</tr>
<tr>
<td>Family history of CAD</td>
<td>2 (8%)</td>
<td>6 (27%) *</td>
</tr>
<tr>
<td>BMI</td>
<td>24.8 (7.3)</td>
<td>29.2 (6.2)</td>
</tr>
<tr>
<td>SBP</td>
<td>119.0 (18)</td>
<td>118.0 (16)</td>
</tr>
<tr>
<td>DBP</td>
<td>78.0 (15)</td>
<td>77.5 (9)</td>
</tr>
<tr>
<td>HbA1c</td>
<td>5.3 (0.3)</td>
<td>5.3 (0.4)</td>
</tr>
<tr>
<td>Triglycerides</td>
<td>126.0 (92)</td>
<td>102.5 (110)</td>
</tr>
<tr>
<td>Cholesterol</td>
<td>182.0 (38)</td>
<td>182.0 (41)</td>
</tr>
<tr>
<td>HDL</td>
<td>57.0 (34)</td>
<td>47.5 (30)</td>
</tr>
<tr>
<td>LDL</td>
<td>91.0 (29)</td>
<td>105.5 (36)</td>
</tr>
<tr>
<td>hsCRP</td>
<td>1.1 (1.5)</td>
<td>1.7 (2.5)</td>
</tr>
<tr>
<td>Total testosterone</td>
<td>26.0 (131)</td>
<td>623.5 (741) *</td>
</tr>
<tr>
<td>Estradiol</td>
<td>91.2 (167) *</td>
<td>35.5 (22.5)</td>
</tr>
</tbody>
</table>

*Statistically significant as determined by p value < 0.05

#19 BEYOND BINDING: KEY QUESTIONS REGARDING THE ROLE OF SHBG IN MALE REPRODUCTION

K Schuppe*, K Roberts. Washington State University Elson S Floyd College of Medicine, Spokane, WA

Purpose of Study Human Sex Hormone Binding Globulin (SHBG) is a homodimeric glycoprotein encoded by the SHBG gene on chromosome 17p13.1. The gene is predominately expressed in the liver and SHBG is secreted into the blood whereas, in murine species where much of our understanding comes from, the ortholog of SHBG is expressed primarily in Sertoli cells. SHBG binds both steroidal and nonsteroidal ligands with high affinities and has been traditionally viewed as functioning as a reservoir and means of transport for steroids in serum. SHBG has also been implicated in androgen uptake by cells and as a signaling molecule with a cell surface receptor on target cells. More recently a germ cell-specific form has been identified in the human testis, raising questions about a role for SHBG in germ cell function and fertilization. The objective of this review was to determine aspects of SHBG function that require further investigation.

Methods Used A systematic search of the literature, using NCBI and other databases compiling peer-reviewed publications, was conducted using keywords: Sex Hormone Binding Globulin; SHBG; SHBG Receptor; SHBG Isoform; SHBG Expression; Megalin; Androgen Binding Protein; ABP. Additional papers were found via citations, both forward and back. The focus of the review was on literature published in the last 25 years, prioritizing the most recent research. Studies were excluded where SHBG function, structure, genetic regulation was not a topic of investigation, such as studies focused on genome-wide analysis, SNP’s, and SHBG in the context of an upstream pathology.

Summary of Results In total 95 publications were identified and kept for producing either new experimental evidence or analysis of prior data that provided new insights into the role of SHBG on reproduction. The literature search converged on studies investigating SHBG’s steroid-binding properties,
protein-protein and protein-receptor interactions, novel germ cell forms of SHBG, and clinical findings from both normal and natural mutant phenotypes. The findings from these studies were summarized and gaps in our knowledge of SHBG’s role in reproductive function were identified.

**Conclusions**

Four unanswered questions regarding SHBG/ABP function emerged from review of the literature. 1) Is SHBG essential in the development and regulation of the reproductive system? 2) What is the identity of the cell surface receptor for SHBG? A number of studies have demonstrated specific binding and subsequent signaling, yet no receptor has been identified. 3) Does the site of SHBG synthesis and secretion matter? 4) What is the function of the germ cell form of SHBG? Germ cells have the second-highest level of expression of SHBG, but we know very little about it. These questions seem key to future research on the role of SHBG in reproduction.

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**#20 SUPRASELLAR AND SELLAR MENINGIOMA IN PREGNANCY PRESENTING WITH WORSENING VISUAL DEFICITS AND PANHYPOPITUITARISM**

1SM Laies*, 2J Fuji, 3C Lovato. 1The University of New Mexico, Albuquerque, NM; 2University of California Los Angeles Health System, Montecito, CA

**Case Report**

Meningiomas account for 13% of CNS tumors during pregnancy and are predominantly low grade WHO I tumors. Case series have reported rapid growth of these tumors during pregnancy, particularly during the second and third trimester, likely due to factors including intra-tumoral hypervascularity and tumor growth due to high sex hormone levels and the presence of progesterone receptors on tumor cells. As a result, patients can present with neurologic and visual field (VF) deficits and in rare cases, herniation and coma. We report a case of a pregnant patient with worsening of VF deficits and panhypopituitarism due to a meningioma in the suprasellar and sellar region. A 36-year-old woman G4P3 at 20 weeks, was transferred to our facility for neurosurgical intervention for acute worsening of VF deficits. She initially presented to her ophthalmologist with a 4-month history of headaches and right-sided vision loss and was found to have a sellar mass on imaging. She then began to have nausea and vomiting, weakness, and orthostasis which necessitated admission. The patient was given i.v. hydrocortisone (HC) due to a suspicion for adrenal insufficiency with improvement in symptoms. However, after receiving steroids, she developed polydipsia and polyuria. On arrival to our facility, the patient was noted to have urinary output (UO) of up to 6 liters per day with a low urine osmolality of 53 mosm/Kg (50–600 mosm/Kg) and serum Na as high as 147 mmol/L (134–144 mmol/L). She was given desmopressin (DDAVP) with improvement in serum Na and UO. Biochemical testing also revealed secondary hypothyroidism and L-thyroxine (L-T4) was initiated. An MRI of the brain on admission showed a 3.6 cm lobulated suprasellar and sellar mass with mass effect on the optic chiasm. Formal VF testing showed complete VF loss on the right and temporal loss on the left. After review of the patient’s case by a multidisciplinary team consisting of Neurosurgery, MFM, and Endocrinology it was decided to proceed with surgery. The patient underwent left pterional craniotomy and resection of the suprasellar component of the mass with decompression of the optic chiasm. Pathology showed a meningioma, WHO Grade I. Post-operatively, VF deficits improved and on discharge from the hospital, she was continued on DDAVP, L-T4 and HC. This is a rare case of a meningioma resulting in hormonal deficiencies in the anterior and posterior pituitary along with VF deficits which likely developed due to the location and rapid growth of the tumor in the setting of pregnancy. The decision for surgery during pregnancy should be based on clinical presentation and should involve a multidisciplinary team to determine the best management that limits complications to the mother and fetus.

**#21 UNRAVELING THE MYSTERY OF THYROTOXICOSIS IN A PATIENT WITH PANHYPOPITUITARISM**

AT Chuang*, BW Hager, M Bouchonville. University of New Mexico Health Sciences Center, Albuquerque, NM

10.1136/jim-2022-WRMC.21

**Case Report**

In the setting of panhypopituitarism, hypothyroidism develops as a result of TSH deficiency rather than a primary insult to the thyroid gland. We describe a case of a patient with acquired panhypopituitarism who developed severe thyrotoxicosis, which persisted despite discontinuation of replacement thyroid hormone.

A 24-year-old Hispanic female with panhypopituitarism following a craniopharyngioma resection in 2010 presented with severe thyrotoxicosis. She initially presented with anxiety and palpitations and was observed to have a mild free T4 (FT4) elevation, which prompted a reduction in her replacement dose of levothyroxine. With progression of her FT4 elevation, the levothyroxine was discontinued and she was ultimately referred to adult endocrinology when clinical presentation worsened. Upon further evaluation, she reported weight loss of more than thirty pounds, palpitations, insomnia, fatigue, muscle weakness, bilateral hand tremors, nausea, vomiting, and dizziness. She confirmed a period of several months without levothyroxine. Heart rate was 109 bpm, no propitosis was noted, thyroid examination revealed no thyromegaly, no tremors were appreciated, and there were no findings suggestive of heart failure. FT4 at that time was 7.6 ng/dL (reference 0.7–1.6 ng/dL). Thyroid peroxidase antibody and TSH receptor antibody levels were negative. Thyroid ultrasound demonstrated an atrophic left thyroid lobe; no nodules were identified. Thyroid uptake was low at 1.8% at 24 hours. This unexpected finding prompted a thyroglobulin (TG) level to distinguish between subacute thyroiditis or an ectopic source of thyroid tissue versus a low TG state such as factitious thyrotoxicosis. TG was low at 3.3 ng/ml, prompting a pharmacy query, which revealed that she was continuing to fill levothyroxine prescriptions at three pharmacies.

A more comprehensive review of her behavioral health history revealed persistent depressive disorder, PTSD, psychogenic non-epileptic seizures, and numerous recent psychosocial stressors. We approached her care in a non-confrontative manner by presenting a range of possible explanations for her clinical and laboratory findings and sharing our recommended treatment.
This case highlights an unexpected presentation of factitious thyrotoxicosis secondary to surreptitious use of levothyroxine in a patient with acquired panhypopituitarism, and the step-wise evaluation that led to this conclusion. We discuss the strategies implemented in managing this patient and review approaches to patients with factitious disorders.

#22 FEASIBILITY OF EXTENDING SLEEP IN A PILOT SAMPLE OF TEENS WITH OBESITY AND SHORT SLEEP: THE REM STUDY

A.E. Bowen*, S Simon, M.Cree-Green, K.J. Nadeau, J Kaar. Children’s Hospital Colorado, Aurora, CO; University of Colorado Anschutz Medical Campus, Aurora, CO

Purpose of Study Insufficient sleep duration is common among adolescents and may contribute to insulin resistance, dysglycemia, and precursors to type 2 diabetes (T2D). Youth-onset T2D has devastating long-term effects, and thus prevention strategies for adolescents at risk for T2D are needed. Increasing total sleep time (TST) may be one such strategy. We tested the feasibility of a 4-week sleep extension intervention for adolescents treated within an outpatient weight management clinic.

Methods Used High school students aged 14–19 years with insufficient sleep (<8 hours) on school nights were recruited during the academic year. Exclusion criteria included T2D, medications that affect sleep, and a schedule that precluded participants from adhering to sleep extension (e.g., night shift employment). Following sleep monitoring at home for 1 week, a revised sleep schedule was collaboratively created with a target of increasing time in bed (TIB) by 2 hours. Participants followed this schedule for 2 weeks (W2), returned to the clinic to discuss barriers to adherence to the prescribed schedule, and then followed the revised schedule for another two weeks (W4). Feasibility was assessed by adherence to wearing sleep watch and submitting sleep diaries, improvement in TIB and TST, and self-reported barriers to following prescribed sleep schedule. Data are reported as averages ± sd or medians (min, max).

Summary of Results A total of 6 participants have been recruited for the study to-date; however, 2 were withdrawn due to COVID-19 school closures when shifted to online learning. Participants (n=4) were aged 17.0 ± 0 years, 100% Hispanic, with a BMI percentile of 98.4±0.9 kg/m². All participants completed the intervention; 99.4% adhered to Actigraphy and 65.6% to the sleep diary. At baseline, the average TIB was 7.3±1.6 hours and TST was 5.8 ±1.5 hours. Compared to baseline, W2 hours of TIB [1.2 (-0.1,2.6) and TST [1.2 (-0.4,2.1)] increased and W4 hours of TIB [0.1 (-1.8,5.3)] and TST [0.7 (-1.5,4.1)] increased. Increased TIB was primarily achieved by shifting bedtimes earlier, while waketimes remained relatively consistent. Barriers to adherence included homework, extracurricular activities, and parents’ and youths’ variable work schedules.

Conclusions Initial findings indicate a 4-week sleep extension intervention in adolescents with short sleep seeking treatment for weight management is feasible. Participants increased TIB and TST by a median of 1.2 hours. However, additional strategies are needed to maintain such improvements. School and community efforts to delay high school start times may benefit youth at risk for T2D by enabling them to increase TST.

#23 RELATIONSHIP BETWEEN HIGH CARBOHYDRATE AND VITAMIN D INTAKE ON SLEEP QUALITY AMONG ADULT POPULATION OF UNITED STATES

C. Brown*, M. Shaheen, K. Schrode. Charles Drew University of Medicine and Science, Los Angeles, CA

Purpose of Study Quality of sleep depends on diet as carbohydrate or vitamin D. Sleep quality is challenging to study due to cost, resources, and availability of research subjects. There is a gap in the literature examining the impact of carbohydrates and vitamin D intakes on sleep quality. We aim to examine the association between carbohydrate and vitamin D intakes and sleep quality among adult US population.

Methods Used We analyzed data from the National Health and Nutrition Examination Survey 2007–2014. Carbohydrate and vitamin D intakes were categorized into three groups based on the distribution. Sleep quality was assessed using hours of sleep, have sleep problem and doctor diagnosed sleep disorder. We used chi square and multiple logistic regression to analyze the data considering the design and sample weight.

Summary of Results Of the 16,415 adults, 35% had high carbohydrate intake (>283 gm), 31% had low vitamin D intake, 36% slept <7 hours/day, 27% reported sleep problem, and 9% had sleep disorder. There was no relation between the high carbohydrate intake and low vitamin D intake and the hours of sleep (p>0.05). However, high vitamin D intake was associated with sleep disorders and troubled sleep (p<0.05) adjusting for the confounding variables. Minority, divorced/widowed, overweight/obese, smokers with diabetes, kidney disease, depression were more likely to have low sleep hours relative to the other groups (p<0.05).

Conclusions Our study indicated no association between carbohydrate and vitamin D intakes and sleep quality. Longitudinal prospective studies are needed to examine factors associated with quality of sleep and their mechanisms.

#24 AUTOIMMUNE HEPATITIS PRESENTING AS FULMINANT HEPATIC FAILURE

N. Trang*, A. Gill, K. Radick, N. Karapetians, S. Mishra. Kem Medical Center, Bakersfield, CA

Purpose of Study To present an interesting case with an atypical presentation of a rare disease.

Methods Used A single patient case report was conducted after IRB approval.

Summary of Results Autoimmune hepatitis (AIH) is a chronic inflammatory disease of the liver that typically presents with the presence of AIH-related antibodies. There are two types of AIH: type 1 is associated with anti-smooth muscle antibody (anti-SMA) and type 2 associated with anti-liver/kidney microsome type 1 antibody or anti-liver cytosol type 1 antibody. About 25% of patients with AIH are asymptomatic. Rarely, patients with AIH present with features of fulminant hepatic failure with rapidly progressive liver impairment, coagulopathy, and hepatic encephalopathy or coma.

We present a case of a 63-year-old female who presents to the hospital with altered mental status (AMS) of one day duration. Her labs were significant for acute kidney injury (AKI), elevated liver function tests, and urinalysis consistent with acute kidney injury. She is found to be anicteric with a normal chemistry. The patient is found to have decompensated liver disease with a MELD score of 40. The patient is initially treated with immunosuppressive therapy, but she fails to respond. A liver transplant is performed, but she succumbs to multiorgan failure. The post-mortem results reveal the presence of autoimmune hepatitis with features of fulminant hepatic failure.
with urinary tract infection (UTI). Computerized tomography (CT) abdomen and pelvis showed bilateral pyelonephritis. Urine cultures grew pan-sensitive Escherichia coli, and she was treated with 10 days of Ceftriaxone. During this course, renal function worsened with uremic-range BUN and she became oliguric, thus hemodialysis was initiated with good response in both renal indices and mental status. Patient was also found to have coagulopathy with a worsening PT/INR of 32.2.2.3.24. Further workup of AKI revealed proteinuria and positive atypical p-ANCA and anti-SMA, raising concern for autoimmune disease in both kidneys and liver.

Conclusions Our patient’s symptoms originally raised concerns for pyelonephritis secondary to a UTI. Despite a full course of antibiotics, her kidney function continued to decline until receiving dialysis. She was also determined to have fulminant hepatitis with coagulopathy followed by positive autoimmune antibodies. This course leads us to believe that the etiology of her encephalopathy was secondary to autoimmune liver and kidney disease.

The main interest of this case report lies in autoimmune hepatitis secondary to atypical P-ANCA and anti-SMA presenting as fulminant hepatic failure in the setting of AKI. Atypical P-ANCA seems to be more specific for autoimmune hepatitis than the typical P-ANCA [1]. Anti-SMA antibodies found in about 50% of type 1 AIH [1]. Both nephrology and gastroenterology recommended renal/liver biopsy for definitive diagnosis which will be done outpatient.

#25 DIABETIC NEUROPATHIC CACHEXIA – AN UNCOMMON MIMIC

1JS Gilbert,  2K McCluskey,  3P Gulani,  4A Outon. 1University of Colorado, Denver, CO; 2Beth Israel Deaconess Medical Center, Boston, MA; 3Jacobi Medical Center, Bronx, NY

10.1136/jim-2022-WRMC.25

Case Report A 52-year-old man with poorly controlled type 2 diabetes presented with four months of watery diarrhea. During this period, he also noticed an unintentional 80-pound weight loss, a ‘burning’ sensation on his anterior thighs, and new onset depression and anxiety. He had no recent history of fever, chills, or night sweats. He was not prescribed insulin, though glipizide was added to his diabetes regimen about six months prior. Physical examination revealed a cachectic man with a BMI of 18.4 kg/m², normal vital signs, and an unremarkable rest of exam. Laboratory evaluation was notable for a white blood cell count of 21.8 x 10^3 cells/mm³, an anion gap of 26, and a glucose level of 280 mg/dL. He also had a CRP of 147.1 mg/dL and an HbA1C of 19.3%. Serology for HIV-1, HIV-2, HbsAg, and HCV were all negative, and a fecal fat test was normal. Chest x-ray was clear, a transthoracic echocardiogram showed no valvular vegetations, and CT scans of his chest, abdomen, and pelvis were completely normal.

This patient’s concerning cluster of symptoms provoked an extensive workup to rule out cancer, infection, and malabsorptive syndromes. However, this patient’s presentation matched an uncommon neuropathy syndrome found in diabetes. Diabetic neuropathic cachexia (DNC) is an extremely rare neuropathy – only 36 cases have been reported in the literature – but its unique cluster of symptoms often incites a search for a hidden malignancy or insidious infection. DNC most commonly presents in a type-2 diabetic after initiation of an oral anti-hyperglycemic medication with profound weight loss, mood symptoms, symmetric peripheral neuropathy, and painful limb paresthesias. Management of this neuropathy is directed at improving glycemic control, as most patients recover within one to two years with improvement in A1C, though some can suffer residual neurologic deficits.

#26 UPPER EXTREMITY DEEP VENOUS THROMBOSIS SECONDARY TO THYROTOXIC PERIODIC PARALYSIS WITH ACCOMPANYING HYPERPHOSPHATEMIA AND SEVERE HYPMAGNESEMIA

J Raheesh*, 1AI-Sukhni, 1B Quraishi, 1T Naguib. Texas Tech University Health Sciences Center, Lubbock, TX

10.1136/jim-2022-WRMC.26

Case Report A 20 year old male with history of drug abuse and osteochondritis presented to the emergency department with bilateral leg weakness and feet numbness since the morning when he woke up unable to move his legs or his back and fell out of bed, after which he noticed that he had feeling in his legs but not his feet. He was found to be profoundly hypokalemic at 1.6 mmol/L and hypomagnesemic at 1.5 mg/dL. Phosphate levels were not taken but calcium was 9.0 mg/dL. These values were verified by a redraw and repeat chemistry. The patient denied nausea, vomiting, and diuretic use, but did report lose bowel movements for long months due to self-diagnosed IBS. He also reported alcohol use of 4–6 beers 1–2 times a week for years. The patient was admitted for potassium and magnesium intravenous replacement. Chest X Ray, MRI of the cerebral spine, CT scan of the brain, and drug panel were all negative.

On day 1 of hospital stay the patient’s electrolyte levels improved, with a potassium of 3.6 mmol/L a magnesium of 1.8 mg/dL, a phosphate of 4.0 mg/dL, and a calcium of 8.9 mg/dL. On day 2, he reported improved strength after working with physical therapy but did not feel at baseline. His labs indicated a potassium of 3.6 mmol/L, a magnesium of 1.8 mg/dL a phosphate of 5.0 mg/dL and a calcium of 9.3 mg/dL. His SARS-CoV2 antibody, blood, and urine cultures were all negative. A negative TTTG IgA test ruled out Celiac disease.

On day 3, potassium was 4.4 mmol/L, and magnesium was 1.7 mg/dL, and a newly elevated phosphate of 5.7 mg/dL was noted. Calcium was 9.0 mg/dL. Right arm edema and pain at the PICC line site prompted removal of the line but an ultrasound showed an occlusive DVT. A heparin drip was ordered, and a V/Q scan was negative. An asymptomatic run of ventricular tachycardia prompted an echocardiogram and troponin evaluation, but both were negative.

On day 4 his labs revealed a low TSH of <0.01 an hypomagnesemia of 1.8 mg/dL. Potassium was within normal limits. Phosphate was still elevated at 5.5 mg/dL and calcium was 10.0 mg/dL.

On day 5 a low TSH of <0.01 was confirmed once again, elevated T3 9.99 pg/ml and T4 2.86 ng/dl were also noted. Potassium at this time was 4 mmol/L, magnesium was within normal limits, phosphate was elevated at 7.0 mg/dL, and calcium was 9.5 mg/dL. A PTH was also measured at this time and was found to be normal at 36.7 pg/ml. Ventricular tachycardia was attributed to hyperthyroidism, and a diagnosis of thyrotoxic periodic paralysis was made. An ultrasound of the neck showed a hypervascular thyroid consistent with Graves’
disease or thyroiditis. Thyroid stimulating immunoglobulin was drawn, and the patient began a regimen of 10 mg methimazole 3 times daily.

On day 6 the patient had a potassium of 3.8 mmol/L, phosphate of 6.2 mg/dL, calcium of 9.6 mg/dL, and a magnesium of 2.3 mg/dL. On day 7 the patient was discharged.

Healthcare delivery research I
Concurrent session
12:45 PM
Thursday, January 20, 2022

#27 DIABETES PREVENTION AT WORK: 10-YEAR FOLLOW-UP OF GLYCEMIC CONTROL AND SERUM LIPOPROTEINS IN HEALTHY WORKERS WITH AND WITHOUT HEALTH COACHING AND MONETARY INCENTIVES

1,2LW Raymond*, 1Atrium Health, Charlotte, NC; 2University of North Carolina System, Chapel Hill, NC

10.1136/jim-2022-WRMC.27

Purpose of Study Knowing of the success of lifestyle changes in the Diabetes Prevention Program, we wondered if similar results could be achieved in a workplace setting with the collaboration of an employer in the Charlotte, NC region.

Methods Used We compared diabetes incidence and serum lipoprotein concentrations in two groups. Cohort 1 comprised 504 workers (mean age 49, 88% males). Cohort 2 comprised 131 workers (age 52, 83% males). Cohort 1 received health coaching by a physician assistant or nurse practitioner who encouraged regular exercise, healthy weight, carbohydrate limitation, smoking abstinence, and blood pressure control. They compared baseline values of HbA1c with those associated with prediabetes and diabetes, ≥5.7 and ≥6.5, respectively. Cohort 1 workers also received up to $800 per year, based on the above lifestyle choices as well as HbA1c and lipoprotein levels. The latter were also measured in Cohort 2, but no incentives or health coaching were provided, other than individual letters containing their blood test results.

Summary of Results Diabetes developed over 10 years in 59 Cohort 1 participants compared to 93 expected (chi-squared = 8.56, p = 0.003) on the basis of initial HbA1c values (Zhang X et al., Diabetes Care 2010;33:1665). Workers with prediabetes decreased from 192 to 141. In Cohort 2, five participants were diagnosed with diabetes, the same number as expected. However, workers with prediabetes increased from 28 to 36.

Conclusions Preventing diabetes benefits individual workers and their families. Employers who share in the health care costs of their workforce also stand to benefit substantially from diabetes prevention, as these costs are $9,601 per year higher in persons with this condition. Health coaching and monetary incentives were associated with improved glycemic control as well as lower lipoproteins. Only the latter improvement was found in the group of workers not provided either of the above interventions in this retrospective analysis. Further prospective observations may identify the respective roles of coaching and monetary incentives. Whether these improvements are associated with better outcomes in cardiovascular end points in these workers would also be of great importance.

#28 MEDICAL STUDENT PERCEPTIONS OF INTIMATE PARTNER VIOLENCE SCREENINGS IN IN-PERSON VS TELEMEDICINE SETTINGS

1 J Lai*, 1A Cheng, 1AL Nelson, 1M Fraix, 1M Hudson, 1Western University of Health Sciences, Pomona, CA; 2University of California Los Angeles, Los Angeles, CA

10.1136/jim-2022-WRMC.28

Purpose of Study Intimate Partner Violence (IPV) is a public health crisis that impacts 25% of women and 10% of men in the US, totaling 43 million women and 38 million men. IPV screenings traditionally occurred at doctor offices, which was to be a safe space, but with the transition to telemedicine, screening is done at home. This study sought to identify potential educational and practice gaps in care surrounding IPV screenings in different settings. We aimed to understand medical students’ general experiences (personal and professional), attitudes, and perceptions of IPV screening.

Methods Used To assess student experiences with IPV screenings as both patients and clinicians-in-training, two separate IRB-approved surveys were created and beta-tested for Western U COMP/COMPNW medical students. One for those who had clinical rotations and the other for those who had not. Questions included personal experiences with in-person and telemedicine IPV screenings, how screenings were conducted, and their perceived importance. Fourth-year students were asked additional questions regarding their experiences observing patients being screened. We used descriptive analysis of the responses to determine the frequency of IPV screening and the modalities in which IPV screening was conducted in in-person and telemedicine environments.

Summary of Results 170 students participated in the study for a response rate of 13%: 140 students from the non-clinical cohort and 30 students from the clinical cohort. Overall, 36.9% of students, who had been seen for an in-person appointment, reported they had been screened for IPV, while only 12.5% of students seen in telemedicine appointments were screened. Among those with in-person appointments, screening was by written survey (31.7%), online survey (10%), and via verbal screen (58.3%). However, among those seen via telemedicine, screening was conducted by online survey (30%) and by verbal screening (70%). Healthcare staff who administered IPV screens shifted from medical assistants (MAs) (35.6%), doctors (35.6%) and nurses (26.7%) for in-
person screenings to MAs (28.6%), doctors (28.6%) and receptionists (28.6%) in the telemedicine screenings. In the clinical cohort, 56% of students observed in-person IPV screenings during rotations, compared to 8.33% of students with telemedicine experience who observed IPV screenings via telemedicine.

Conclusions Medical students reported IPV screening was decreased in the telemedicine setting, which could increase the risk of under detection. Virtual screening was more frequently done verbally and conducted by less trained personnel. This study is unique because it provides the perspective of medical students as both patients and healthcare providers in training and demonstrates both educational and practice gaps in this new environment.

#29 TDAP VACCINE UPTAKE AND ATTITUDES AMONG ENGLISH AND SPANISH SPEAKING PREGNANT PATIENTS

1E Jimenez*, 2H Stoil, 3A Nelson. 1Western University of Health Sciences, Pomona, CA; 2Harbor-UCLA Medical Center, Torrance, CA; 3University of California Los Angeles David Geffen School of Medicine, Los Angeles, CA.

Purpose of Study Despite the recommendations of The American College of Obstetrics and Gynecology, and the health benefits that administering the Tdap vaccine can have on prenatal patients and their newborns, nearly 45% of prenatal patients do not receive the vaccine. We surveyed English and Spanish speaking prenatal patients, to measure uptake and assess patient knowledge and attitudes of the Tdap vaccine. We compared the responses of English and Spanish speaking patients to identify if inconsistencies existed.

Methods Used Using an IRB approved protocol, we surveyed low-income patients at Harbor UCLA Medical Center Obstetrics clinic in Torrance, CA for 7 weeks in Summer 2021. Patients were at least 18 years of age and ≥ 32 weeks gestational age. Upon obtaining verbal consent, we administered a 31-question beta-tested survey in their preferred language, English or Spanish.

Summary of Results The response rate achieved by the survey was 97%. There was a total of 98 participants, 80 of which answered the survey in English and 18 in Spanish. 49% of subjects were Hispanic/Latino, 35% African American, 3% Caucasian and 13% were of other ethnicities. Most of our patients (67%) were between the ages of 20-30. 69% of English-speaking patients received the vaccine and 79% of Spanish speaking patients also received the vaccine. Among all our 98 participants there was an overall Tdap vaccine uptake of 70%. The most common reason for refusal of the Tdap vaccine among English speaking patients was due to safety concerns for their baby (47%), concerns for themselves (47%) or because they believed that they did not need the vaccination (47%). While, only 4% of English-speaking patients denied the vaccine because they were unaware they required it, this was the most common reason for refusal of the vaccine by Spanish speaking patients (66%).

Conclusions Overall, 70% of patients received the Tdap, higher than national averages, but still short of the goal, leaving substantial numbers of women and their newborns unprotected. Both uptake and reasons for refusal of Tdap vaccine differed among English and Spanish speaking patients. This may suggest a need to look more closely at the how Tdap vaccine information is presented in both languages to ensure that the same information or information more relevant to different groups, is being relayed to all prenatal patients.
Purpose of Study: Traumatic brain injury (TBI), is a common injury amongst veterans who have served in Iraq and Afghanistan. With the number of veterans from these conflicts now approaching 3 million, it is estimated that approximately 20% have suffered at least one TBI. Beyond the structural trauma, TBI may also lead to transient, or even permanent, pituitary insufficiency. Of particular interest are the consequences of TBI-induced hypogonadotropic-hypogonadism (HG) on short and long-term health. TBI clearly presents economic implications for the nation, related both to direct medical expenses and indirect costs. The purpose of this review is to summarize the knowledge about post-TBI hypopituitarism, its screening and treatment recommendations, costs, with a special focus on the potential impacts of post-traumatic hypogonadism on naval special operators.

Methods Used: We utilized DoD, CDC, and NIH datasets on TBI’s, alongside Endocrine Society and AUA Guidelines, and finally NCBI and Google Scholar searches of the following key terms: pituitary dysfunction, traumatic brain injury, hypogonadotropic hypogonadism, hypogonadism treatment, TBI screening, hormone therapy, fertility, special operations, special forces. The aforementioned sources were used to roughly predict prevalence of post-TBI Hypogonadism in special operators, it’s potential costs and consequences, and carry over its standards for screening and treatment.

Summary of Results: According to recent estimates, the range of TBI amongst all servicemen serving in Iraq or Afghanistan, taken together from 2000 to 2016, runs from 11–23%. Persistent hypogonadotropic hypogonadism following TBI meanwhile, amongst the general population, is predicted to fall within the wide range of 8–41%. It was found that hypogonadism was associated with incident of PTSD among other physiologic consequences such as sexual dysfunction, osteoporosis, and neurodegeneration. Recent literature has proposed post-TBI pituitary dysfunction screening and therapy among the general public, but little for post-deployment special forces or veterans specifically.

Conclusions: While we can crudely postulate from comparisons to other groups, there is a distinct lack of recent data on TBI’s in naval special forces. But, what is clear is the connection between TBI’s and subsequent pituitary dysfunction, in which hypogonadotropic hypogonadism is likely the second most common subtype. Taken together, further retrospective and prospective studies are needed to further investigate hypogonadotropic pituitary dysfunction after TBI’s acquired in the line of duty by naval special forces servicemen with the purpose of establishing screening guidelines for these servicemen and ultimately provide appropriate treatment algorithms for the preservation of their quality of life, fertility, and protection against comorbid disease.
thoracic cancer. Competency guidelines for residents and fellows to properly perform FNAs are lacking, leading to these essential skills being taught on the job. Simulation-based trainings offered by professional societies are effective but often require travel and are expensive. We created a brief hands-on module designed to be 1–2 hours in length to introduce trainees to basic thyroid ultrasound (US) and US-guided FNA to improve trainees’ comfort with the procedures. This study evaluated whether participating in this module improves the comfort of resident and fellows with thyroid US and FNA prior to performing the procedure on patients while being cost- and time-effective.

Methods Used A hands-on training module for US-guided FNAs was developed and offered yearly for 6 years to residents and fellows at Harborview Medical Center. The models used were purchased from Northwestern Medical Center for $25 each and one model was used for each session. 40 pre-surveys and 26 post-surveys were collected directly before and after the module. Participants were primarily otorhinolaryngology residents (n= 15, 11) and endocrinology fellows (n=13, 8) with varying experience. The surveys assessed their comfort level performing US-guided FNAs on a scale of 1 to 5, with 5 being most comfortable and able to perform independent of supervision. The surveys also assessed their comfort with interpreting thyroid US, and long- and short-axis US-guided FNA. Effectiveness was assessed by calculating the change in comfort using population averages, regardless of specialty on the post-survey compared with the baseline level. Significance was determined using a permutation test.

Summary of Results On average, participants’ comfort with US-guided FNA increased by 1.19 (p=0.0006) on the 1–5 scale, comfort level with long- and short-axis US-guided FNA increased by 1.54 and 1.51, and comfort with interpreting thyroid US improved by 0.97. For endocrinology fellows and otorhinolaryngology residents specifically, their change in comfort level for performing US-guided FNA increased by 1.02 and 1.17.

Conclusions Overall, residents and fellows showed an improvement in comfort level after completion of the module. This improvement was not only evident in performing US-guided FNA but also with FNA technique and US interpretation. Although comfort does not equate skill, at $25 per session our module is a promising alternative to costly and time-consuming simulations courses, which often cost upwards of $700 per individual. These training modules can be executed in most residency and fellowship training programs to provide accessible training of these important skills.

#34 PRE-OPERATIVE ISTOP HUDDLE FOR IMPROVED TRANSITION OF CARE OF PEDIATRIC AND NEONATAL CRITICAL CARE PATIENTS

Pre-operative (pre-op) checklists have been shown to improve patient safety. The iSTOP checklist, currently used for surgical patients, has been utilized for non-surgical areas in the hospital. In ICUs, a similar checklist would reduce the number of surgical delays and need for postoperative intensive care.

Purpose of Study Patients admitted to the pediatric or neonatal intensive care units (PICU or NICU) at Helen DeVos Children’s Hospital in Grand Rapids, MI, prior to their surgery are taken directly from the PICU or NICU to the operating room (OR). Therefore, these patients do not undergo routine pre-operative (pre-op) checklists in the pre-op holding area. A critical care pediatric patient underwent a wrong-sided surgery, highlighting the need for a standardized approach to improve completion of the pre-op checklist and communication between the ICU and surgery teams in the perioperative period.

Methods Used Using quality improvement methodology, the NICU, PICU, pediatric surgery, and pediatric hospital medicine teams completed an A3 form and performed a gap analysis. To address the concerns identified in the root cause analysis, we developed a bedside team huddle composed of ICU, surgery, and anesthesia teams to be performed in the ICU prior to the patient being taken to the OR. We created the acronym iSTOP to outline the components of the pre-op huddle: (i) introductions; (S) surgical procedure to be performed; (O) any tubes, lines, or drains; (T) ongoing plan/intra-operative plan; and (P) post-operative care and pain management plan.

Summary of Results Over the course of 90 days, 24 pre-op bedside iSTOP huddles were convened for ICU patients requiring surgery. All team members were present and all key elements of iSTOP were reviewed in over 90% of instances. Surgical site was appropriately marked 100% of the time, and pre-op checklist was completed at least 80% of the time. During this time period, there were zero serious safety events for ICU-to-OR patients.

Conclusions The iSTOP huddle improved completion rate of the pre-op checklist and enhanced care team communication and patient safety surrounding care transitions between ICU and surgical departments. This huddle format can be extended to incorporate other bedside procedures within the ICUs and other areas of the hospital.
cases were excluded. Surgeries that were delayed due to COVID, or those that deviated from the established care pathway were also excluded. Impact of the checklist on the frequency of peri-operative delays and need for postoperative intensive care were investigated. Chi-square analysis was used to interpret these data.

Summary of Results Of 235 patients scheduled for complex spine surgery, 193 met our criteria. Checklist-directed surgical optimization did not significantly reduce surgical delays, with 19.0% of surgeries experiencing a delay in the historical control group compared to 15.7% in the study group (p = 0.38). However, patients in the study group were less likely to require postoperative intensive care (11.1%) compared to the control group (25.3%) (p = 0.031).

Conclusions Checklist directed pre-surgical optimization was instituted at a single, high-volume spine surgery center. Although this intervention did not reduce the number of surgical delays, it does have the potential to increase patient safety, as use of the checklist was associated with reduced need for postoperative intensive care. Further research on ways to improve interdisciplinary coordination for preoperative optimization to reduce surgical delays is needed to maximize patient safety and minimize AEs.

Immunology and rheumatology I
Concurrent session
12:45 PM
Thursday, January 20, 2022

#36 SARS-COV-2 IMMUNE COMPLEXES ELICIT TISSUE FACTOR EXPRESSION ON HUMAN PERIPHERAL BLOOD MONOCYTES

1,2 J Plagenz*, 2 T Peters, 1 S Pincus, 1 N Meissner. 1 University of Washington School of Medicine, Seattle, WA; 2 University of California, San Francisco, CA

Purpose of Study In innate and adaptive immune responses may play a role in severe complications of SARS-CoV-2 infection (COVID-19). The formation of virus-antibody immune complexes may result in aberrant activation of innate immune cells, including circulating monocytes. Thromboembolic complications are a hallmark of severe COVID-19, in which a hypercoagulable state is observed. Currently, the mechanism of this is poorly understood. Tissue factor, also known as Coagulation Factor III, is key to activating the clotting cascade; it is constitutively expressed extra-vascularly but can be upregulated on circulating monocytes during inflammation. Conditions that predispose patients to severe COVID-19, such as metabolic syndrome, are associated with elevated plasma levels of endotoxin. We postulate that aberrant inflammatory activation of monocytes via SARS-CoV-2/antibody immune complexes, in tandem with endotoxin, can upregulate tissue factor and induce hypercoagulability.

Methods Used Immune complexes were formed by mixing inactivated SARS-CoV-2 with Bamlanivimab (Bam), a therapeutic monoclonal antibody specific for the spike receptor-binding domain of SARS-CoV-2. Antibodies to a different domain of the spike were used to capture immune complexes, which were then detected using biotin/avidin. Effects on monocyte cell-surface expression of tissue factor were investigated using flow cytometry. Human peripheral blood mononuclear cells were cultured with SARS-CoV-2, Bam, endotoxin, and combinations of the three. Monocytes were identified by forward/ side scatter and CD14 expression.

Summary of Results SARS-CoV-2 immune complexes were readily detectable by immunoassay. Immune complexes were also stable under different storage conditions. These complexes increased endotoxin-induced tissue factor expression on monocytes to a greater degree than did endotoxin alone. Incubation with neither Bam nor SARS-CoV-2 alone induced tissue factor expression.

Conclusions Antibody-mediated mechanisms are key in clearing SARS-CoV-2 infection. Our results show that the formation of virus-antibody immune complexes may also result in aberrant activation of innate immune cells, including circulating monocytes, leading to tissue factor upregulation. These results may aid in understanding the hypercoagulable state seen in SARS-CoV-2 infection. The next step is to evaluate the effect of immune complexes on in-vitro coagulation by using tissue factor-induced Factor Xa activity assays.

#37 HOW THE TFH RESPONSE IN NEONATES FED BREAST MILK DEVOID OF MATERNAL ANTIBODIES IMPACTS THE COMPOSITION AND FUNCTION OF INTESTINAL MICROBES

1 IT Vantreu*, 2 B Wang, 3 S Torres. 1 University of Washington School of Medicine, Seattle, WA; 2 Fred Hutchinson Cancer Research Center, Seattle, WA

Purpose of Study Breast milk is essential to the health and development of children. In addition to nutrients, breastmilk contains immune-modulating factors, including antibodies that protect infants from infections. However, breastfeeding is not available for all women and children. While infant formula is designed to meet growing infants’ basic nutrition needs, it does not contain factors like antibodies. We discovered that in the absence of breastmilk antibodies, mice develop increased levels of CD4 T follicular helper (Tfh) cells and germinal center (GC) B cells in gut draining lymphoid tissues. However, in germ-free mice that were also deficient in maternal antibodies (matAbs), the Tfh and GC B cell levels closely reflected antibody-sufficient neonates’ levels. These results suggest the significance of the microbiota in the role of Tfh and GC B cell responses. Tfh cells are essential for maintaining host-microbe homeostasis, and dysregulated increases in Tfh cells can alter the microbiota composition, potentially causing colitis. An indispensable role of the microbiota is to prevent the spread of pathogenic infections through colonization resistance. We hypothesize that the increase in Tfh and GC B cells in pups lacking maternal antibodies will alter the intestinal microbiota and its function.

Methods Used To understand how the Tfh cells can alter the intestinal microbiota, we treated half of the mice that are sufficient and deficient in maternal antibodies with anti-ICOSL, which effectively dampens the Tfh cell expression. We then infected all the mice with Salmonella typhimurium, a bacterial pathogen. Fecal samples were collected daily for 5 days. On day 5 post-infection, fecal samples, cecum, and liver were harvested to determine infection burden.
Abstracts

Summary of Results Our data indicates no significant difference in colony-forming units (CFU) between the matAb sufficient and deficient groups, signifying that the presence of maternal antibodies does not change the susceptibility to S. typhimu-
rium infection. In addition, there was no difference seen between the groups treated with anti-ICSOL versus control, indicating that early life Tfh cells giving rise to GC B cells producing T-dependent antibodies do not play a role in con-
ferring differential resistance to S. typhimurium. This trend was seen when examining both localized and systemic infection across all three organs.

Conclusions Although neonates that do not receive matAbs in breastmilk have increased numbers of Tfh and GC B cells which have the potential to produce antibodies and change the microbiota composition of the gastrointestinal tract, there was no difference in S. typhimurium infectivity between mice transiently devoid of Tfh cells versus control. To continue exploring the role of colonization resistance, further research is needed to determine how Tfh cells influence host-microbe interactions and subsequently its change in infection susceptibility.

#38 CONSERVATIVE MANAGEMENT OF KNEE OSTEOARTHRITIS UTILIZING BONE MARROW ASPIRATE CONCENTRATE

1B Leiby*, 2GB Foremny, 2J Hanley, 2J Galloway, 2J Willford, 2,3JC McGinley. 1University of Washington School of Medicine, Seattle, WA; 2The McGinley Clinic, Casper, WY; 3WWAMI Medical Education Program, Laramie, WY; 4University of Washington School of Medicine, Seattle, WA

Purpose of Study Bone marrow aspirate concentrate (BMAC), along with conservative patient management, offers a mini-
mally invasive option in treating chronic pain from knee osteoarthritis. Knee osteoarthritis affects 35% of adults aged 65 years and older. BMAC has been shown to decrease inflammation and improve cartilage signal on MRI. We hypothesize BMAC injections, along with conservative care, will provide short- and long-term relief of pain associated with knee osteoarthritis.

Methods Used A retrospective chart review was conducted to identify patients with knee osteoarthritis who received BMAC injections and conservative care in our clinic from November 2013 to November 2019. Under CT and ultrasound guidance, 60cc of bone marrow was aspirated from the posterior iliac crest. Each 60cc sample of aspirate was centrifuged, concen-
trated to 10cc, and injected into the knee joint under sono-
graphic guidance. Patients were non-weight bearing utilizing crutches and a compartment specific off-loading brace for 3 weeks after the procedure and partial weight bearing with just the brace for an additional 3 weeks. All NSAIDs were held 10 days prior and 3 months following the procedure. A 0–10 patient self-reported pain scale was used as the primary outcome. Secondary outcomes included adverse events and additional treatments. Pain scores were collected on the day of treatment and fixed timepoints up to 3 years post-injection. A 2-tailed Wilcoxon signed rank test with a .05 alpha level was used to determine statistical significance between differences in reported pain level at each follow-up compared to baseline.

Summary of Results Forty-seven patients (71 knees, 26 males, 21 females) with an average age 64±9 years, received BMAC injections and conservative management. These patients were followed for 3 years post treatment (mean follow-up 30.1 ± 11.0 months). Reported pain level was significantly reduced 3 weeks post-injection compared to baseline (47 patients, 71 knees; mean Δ -2.0 points; p<.001; table 1). Pain continued to decrease up to 3 years post-injection compared to baseline (37 patients, 54 knees; mean Δ -3.9 points; p<.001). No adverse events were reported. Thirtyseven patients (17 knees) subsequently received additional treatments including injections (10 patients, 12 knees). Three patients (5 knees) underwent knee arthroplasty. Zero patients underwent repeat BMAC injections during the 3-year follow-up period.

Conclusions BMAC injections, along with conservative manage-
ment, represents a safe, effective, and minimally invasive treat-
ment option for treating knee osteoarthritis pain for up to 3 years. Few patients in our study progressed to knee arthro-
plasty suggesting this approach to be a viable alternative to surgery.

#39 PREVALENCE OF FRAILTY AND ASSOCIATED FACTORS IN A NATIONAL OBSERVATIONAL COHORT OF RHEUMATIC DISEASES

1C Chang*, 1N Singh, 1J Andrews, 2K Wipfker, 3S Lieber, 4SE Sattui, 5JP Baker, 1P Katz, 1K Michaud, 1K Wysham, 1UW, Seattle, WA; 2FORWARD, Wichita, KS; 3HSS, New York, NY; 4UPitt, Pittsburgh, PA; 5UPenn, Philadelphia, PA; 6VA, Philadelphia, PA; 7UCSF, San Francisco, CA; 8VHA, Seattle, WA

Purpose of Study Frailty is associated with disability and early mortality and may be reversible. It is accelerated in patients with certain rheumatic musculoskeletal diseases (RMDs). The prevalence of and disease-specific factors associated with frailty across multiple RMDs is unknown.

Methods Used Data were acquired from FORWARD, The National Databank for Rheumatic Diseases, an observational longitudinal US registry with biannual patient questionnaires. Frailty was measured by self-reported measure: the FRAIL scale, which queries 5 items: 1) fatigue, 2) resistance (climbing stairs), 3) ambulation, 4) illnesses, and 5) loss of weight and categorizes those with ≥3 items as frail. Those with missing RMDs or frailty variables were excluded (N=117). Prevalence of frailty across RMDs was described. Multivariable logistic regression was performed to identify variables independently

<table>
<thead>
<tr>
<th>Abstract #38 Table 1</th>
<th>Mean reported pain scores</th>
</tr>
</thead>
<tbody>
<tr>
<td>Time</td>
<td>Baseline</td>
</tr>
<tr>
<td>n = 71</td>
<td>n = 71</td>
</tr>
<tr>
<td>Mean Pain Score</td>
<td>5.6</td>
</tr>
</tbody>
</table>

*p < .001
associated with frailty in the entire cohort and stratified by RMDs.

**Summary of Results** 3,348 individuals were included and 1,084 were frail (32%). RMDs evaluated were rheumatoid arthritis (71%), osteoarthritis (OA) (16%), fibromyalgia (5%), systemic lupus erythematosus (SLE) (4%), other connective tissue diseases (CTDs) (2%), spondylarthropathy (1%), and vasculitis (1%). Frail participants were older (69.8±10.6) compared to non-frail (66.3±11.7) and had a higher prevalence of obesity (52% vs. 31%). The distribution of frailty was equal across RMDs (~33%) except vasculitis and CTDs, which had a lower prevalence of frailty (20% and 26%, respectively). Ambulation and fatigue were the most common frailty components across RMDs. In the primary multivariable model evaluating the entire cohort, increasing age (OR=1.05 [95%CI 1.04–1.06]), female sex (OR=1.74 [95%CI 1.57–1.95]), overweight (OR=1.49 [95%CI 1.17–1.89]) and obesity (OR=3.04 [95%CI 2.42–3.82]), prior fracture (OR=1.87 [95%CI 1.56–2.26]), increased disease activity (OR=1.24 [95%CI 1.18–1.30]), and pain (OR=1.11 [95%CI 1.07–1.16]) had significant independent associations with frailty (table 1). Biologic use was associated with lower odds of frailty (OR=0.78 [95%CI 0.64–0.96]). Among the RMDs, SLE was associated with an increased odds of frailty with OA as the reference (OR=1.70 [95%CI 1.02–3.03]). Overall, disease-specific associations were similar to the primary multivariable model with obesity and disease activity maintaining statistical significance in most models.

**Abstract #39 Table 1 Multivariable logistic regression evaluating factors associated with frailty in the entire cohort (N=2947)**

<table>
<thead>
<tr>
<th>Variables</th>
<th>OR</th>
<th>95%CI</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>1.05</td>
<td>1.04–1.06</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Male sex</td>
<td>0.74</td>
<td>0.57–0.95</td>
<td>0.017</td>
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<td>BMI:</td>
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<tr>
<td>-Underweight</td>
<td>1.60</td>
<td>0.87–2.93</td>
<td>0.131</td>
</tr>
<tr>
<td>-Normal weight</td>
<td>ref</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>-Overweight</td>
<td>1.49</td>
<td>1.17–1.89</td>
<td>0.001</td>
</tr>
<tr>
<td>-Obese</td>
<td>3.04</td>
<td>2.42–3.82</td>
<td>&lt;0.001</td>
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<td>Primary Rheumatic Diagnoses:</td>
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<tr>
<td>-Rheumatoid arthritis</td>
<td>1.18</td>
<td>0.91–1.54</td>
<td>0.217</td>
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<tr>
<td>-Systemic lupus</td>
<td>1.76</td>
<td>1.02–3.03</td>
<td>0.042</td>
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<td>-Erythematos</td>
<td>ref</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>-Osteoarthritis</td>
<td>ref</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>-Fibromyalgia</td>
<td>0.98</td>
<td>0.65–1.48</td>
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<td>-Connective Tissue</td>
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</tr>
<tr>
<td>Diseases*</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fracture ever</td>
<td>1.87</td>
<td>1.56–2.26</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Disease duration (years)</td>
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<td>1.00–1.01</td>
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<td>Disease severity</td>
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<td>1.18–1.30</td>
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</tr>
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<td>Pain Scale</td>
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<td>1.07–1.16</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Medication use</td>
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<tr>
<td>-DMARD use</td>
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<td>0.77–1.13</td>
<td>0.452</td>
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<tr>
<td>-Biologic Use</td>
<td>0.78</td>
<td>0.64–0.96</td>
<td>0.019</td>
</tr>
<tr>
<td>-Prednisone dose</td>
<td>1.01</td>
<td>1.00–1.03</td>
<td>0.142</td>
</tr>
</tbody>
</table>


**Conclusions** Frailty is common among RMDs affecting nearly 1 in 3 participants. Obesity, prior fracture and a diagnosis of SLE had the highest associations with frailty. Future work is needed to identify factors that predict frailty onset and potential interventions to treat frailty within RMDs.

**Abstract #40 ADJUNCT THERAPIES FOR PEMPHIGUS DISEASE: A SYSTEMATIC REVIEW**

1K Nguyen*, 2S Worwick, 1Western University of Health Sciences, Pomona, CA; 2University of Southern California, Los Angeles, CA

10.1136/jim-2022-WRMC.40

**Purpose of Study** Pemphigus vulgaris (PV) and IgA Pemphigus are mucocutaneous autoimmune diseases that commonly present as painful blisters eroding the skin of the face, trunk, scalp, groin, and axillae in affected patients. The pathogenesis of pemphigus disease stems from autoantibodies against desmosomal cadherin proteins essential to maintaining keratinocyte adhesion. A histopathologic exam may reveal a reduction in desmosomal cadherin proteins and epidermal acantholysis. Currently, there is no cure for pemphigus, though corticosteroids and steroid-sparing agents are commonly used to control the proliferation of lesions and prevent disease progression. Frequently used non-steroidal agents include mycophenolate mofetil, azathioprine, IVIG and rituximab. Despite these treatment options, patients often succumb to long-term corticosteroid complications. Less oft-used therapies include dapsone and sulfasalazine for PV, and colchicine for IgA pemphigus which offer potential steroid-sparing alternatives with fewer adverse effects, however, their efficacies has not been clearly established. The objective of our systematic review is to investigate the use of dapsone, sulfasalazine, and colchicine in the treatment of PV and IgA pemphigus.

**Methods Used** We searched the PubMed database using the search terms: ‘dapsone’ ‘sulfasalazine’ ‘pemphigus vulgaris’ ‘colchicine’ ‘IgA pemphigus disease.’ Our inclusion criteria included published articles written in English between 1970–2021 exploring the use of dapsone, colchicine or sulfasalazine for pemphigus, and included case series, retrospective studies, and randomized control trials. Our exclusion criteria eliminated reports with fewer than three patients, and review articles. 275 articles were identified, of which 27 relevant studies were eligible. 15 studies were excluded after screening, resulting in 12 remaining studies.

**Summary of Results** 46 (63%) out of 73 patients responded to dapsone, suggestive of its efficacy as either a monotherapy for or as a part of combination therapy for PV. In 65 patients receiving sulfasalazine adjunct therapy, 61 (94%) achieved clinical remission. Adequate data is lacking regarding colchicine therapy for pemphigus as the current literature only reports four IgA pemphigus patients treated with this agent.

**Conclusions** More research is required to elucidate an effective and safe therapy for individuals burdened with pemphigus disease. Certainly, the rarity of this condition and the difficulty in finding adequate control groups present a major barrier for holding clinical trials on alternative therapies. Going forward, dermatologists may consider the use of dapsone or sulfasalazine adjuvant therapy in PV patients to slowly lower corticosteroid use as lesions begin to diminish and to prevent relapse of cutaneous flare ups for patients in remission.

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### Abstracts

#### #41 THE ADJUNCTIVE THERAPY FOR MYCOBACTERIUM TUBERCULOSIS INFECTION IN TYPE 2 DIABETES MELLITUS

C Sidiyan, A Beever, N Khouz, J Owens, K Sasaminia, A Kalloli, W Khamsa, S Subbian, V Venketaraman. Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA; Western University of Health Sciences, Pomona, CA; Rutgers New Jersey Medical School, Newark, NJ; Western University of Health Sciences, Pomona, CA

**Purpose of Study** Type 2 Diabetes Mellitus (T2DM) is an inflammatory disease that can alter the immune response resulting in several physiological manifestations. Glutathione (GSH), a thiol required to maintain intracellular redox state homeostasis, is classically deficient among individuals with T2DM. Glutathione also appears to be pertinent in the immune response against *Mycobacterium tuberculosis* (*Mtb*) infection. In our previous studies, we have identified L-GSH’s direct opposing effects against oxidative damage as well as its immune enhancing effects in HIV+ patients. We explored for similar effects in T2DM which also involves inflammatory and infectious states that could potentiate the replication of *Mtb* and further diminish the immune response. Specifically, our study aims to further elucidate GSH’s role in the granulomatous effector response. In this study, we attempted to uncover whether GSH deficiency in diabetic mice (db/db) impairs the formation of granulomas and the granulomatous effector response to further the understanding of the detailed mechanism of *Mtb* pathogenesis and the potential for novel therapies against the disease brought on by the infection.

**Methods Used** Db/db mice were infected with *Mtb* and treated with one of 3 regimens, either: 1) an optimal dose of rifampicin (RIF), 2) a suboptimal dose of RIF, or 3) a suboptimal dose of RIF in addition to reduced form of GSH encapsulated in liposomes (L-GSH). 3 male and 3 female mice were sacrificed for each group over the span of 3h, 2, 4 wks, 6 wks, and 8 wks post-treatment to study the collective effects of L-GSH and RIF in *Mtb* infection. Granuloma samples from each group were formalin-fixed and analyzed accordingly. We are currently measuring the survival of *Mtb* along with the levels of cytokines, free radicals and GSH in untreated, RIF-treated and RIF+GSH-treated db/db mice.

**Summary of Results** We expect to obtain the data from the aforementioned assays shortly.

**Conclusions** If our data shows a marked elevation of immune defensive cytokines and granuloma formation with concurrent reduction in *Mtb* survival and free radical production in L-GSH treated mice, then we can support our hypothesis that GSH enhances the granulomatous effector response against *Mtb* infection in T2DM. In addition, if we observe greater immune responses in RIF+GSH treated db/db mice, we may further explore the use of GSH as an adjunct therapy in *Mtb* infection in T2DM.

#### #42 ACTINIC GRANULOMA: A RARE CASE OF SUN DAMAGE

K Nguyen, C Wong, E Nguyen. Western University of Health Sciences, Pomona, CA; Riverside Community Hospital, Riverside, CA

**Case Report** A 53-year-old female with a past medical history of anxiety presented to the dermatology clinic with a pruritic eruption for six weeks. She had been gardening without gloves the day prior to the onset of the eruption. A review of systems was inconclusive. Her examination revealed diffuse red papules coalescing into plaques with mild scales involving the scalp, face, neck, torso, and upper and lower extremities including palms and soles, and sparing the ears, bilateral axillae, elbows, and knees. Her biopsy revealed solar elastosis and abundant multinucleated foreign body giant cells with ingested elastic fibers. The patient’s clinical presentation and histopathology was consistent with a diagnosis of actinic granuloma (AG). Her treatment included 20 mg of prednisone PO QAM for one month along with fluticasone 0.05% face cream BID and triamcinolone 0.1% cream BID applied to the affected skin on the body. After one month, all lesions flattened except for post-inflammatory erythema macules. Sun avoidance and daily sunscreen use was also recommended. At the most recent follow up, her lesions resolved demonstrating the efficacy of corticosteroid treatment.

#### #43 COMPLETE CONGENITAL HEART BLOCK IN A NEONATE

S Berke, S Sukumaran. Valley Children’s Hospital, Madera, CA

**Case Report** A 5-day-old female presented with heart block. She was born at term with a birth weight of 3000 g, with an APGAR score of 7/9. She was born via normal ventouse assisted delivery. She was discharged home on day 2 with a 2:1 atrioventricular block. On day 5, she was brought to our neonatal intensive care unit with tachycardia and tachypnea. A chest x-ray revealed cardiomegaly, and an echocardiogram showed a non-conducting ventricular septum defect. She was diagnosed with complete congenital heart block and transferred to the pediatric cardiology unit. She was treated with intravenous injection of aminophylline to promote ventricular depolarization, resulting in stable cardiac rhythm. She was subsequently discharged home on day 10 with normal ventricular rhythm.
Case Report Congenital heart block (CHB) in neonates is associated with high morbidity and mortality. CHB generally occurs due to the presence of maternal autoantibodies of the Ro/La family or cardiac defects.

We describe a neonate born with CHB who was found to have neonatal lupus erythematosus (NLE).

Methods Used Case Report

Summary of Results A term female infant was born by cesarean delivery at 37 weeks to a 24-year-old healthy primigravida. At delivery, the baby’s heart rate was 55 beats per minute. The patient was admitted to the neonatal intensive care unit for further evaluation and management of fetal bradycardia.

Electrocardiogram demonstrated third-degree atrioventricular (AV) block and fetal echocardiogram showed a ventricular rate of 60–65 beats per minute and an atrial rate of 116–128 beats per minute. There was good ventricular function without evidence of hydrops.

Physical exam revealed a term, well-appearing infant female with bradycardia but normal S1 and S2 without murmurs. The remainder of her examination was within normal limits. Laboratory evaluation of the infant and the mother showed positive anti-SSA/Ro and anti-SSB/La antibodies.

The baby was diagnosed with NLE and CHB. The infant’s heart rate was monitored closely but she maintained a heart rate greater than 60 beats per minute and hence was discharged home. A pacemaker was scheduled for placement as an outpatient.

Conclusions NLE is a rare acquired autoimmune disorder that occurs due to passive placental transfer of maternal autoantibodies to SSA/Ro and/or SSB/La. Anti-SSA/Ro autoantibodies are found in about 85–90% of mothers of neonates with CHB, and studies of pregnancies in anti-SSA/Ro positive mothers estimated the risk of CHB to be 1–5%.

Cardiac involvement in NLE is usually irreversible and characterized by second- or third-degree CHB. A ventricular rate of less than 35 beats per minute, hydrops fetalis, or atrioventricular valve regurgitation indicate poor fetal prognosis.

This case emphasizes the importance of considering NLE in infants with fetal bradycardia, congenital AV block or arrhythmias and evaluating the mother and infant for autoantibodies to SSA/Ro and/or SSB/La. We also highlight the need for early referral to cardiology and possible pacemaker implantation in infants who do not respond to medical therapies alone.

#44 OBJECTIVE CHARACTERIZATION OF OSTEOPATHIC LYMPHATIC PUMP TECHNIQUE EFFICACY USING SERUM COVID-19 ANTIBODY LEVELS FOLLOWING VACCINATION

A Comer *, E Lee, E Martinez, B Lovelless, P Croone, S Fuchs, H Szumant, J Sanchez. Western University of Health Sciences, Pomona, CA

10.1136/jim-2022-WRMC.44

Purpose of Study Lymphatic Pump Technique (LPT) is an Osteopathic Manual Medicine technique involving external pressure to various lymphatic structures with the goal of improving lymph drainage. Because of the leukocyte content of lymph, LPT is often indicated for use as an adjuvant therapy in patients with acute or chronic infections. Previous studies have primarily characterized LPT efficacy in terms of clinical or symptomatic outcomes, typically with rather small cohorts of subjects, which has been a criticism of the technique. To address the limitations of past studies, we present a study in which the serum concentration of anti-spike protein COVID-19 antibodies are measured in 100 subjects following treatment with LPT (experimental group) and 100 subjects without LPT treatment (control group) in conjunction with COVID-19 vaccination. The ongoing study is designed to follow the subjects for one year after the first COVID-19 vaccine.

Methods Used Subjects were split into treatment or control groups in a double-blinded randomized process. Participants returned for blood draws with the following schedule based on the day of their first vaccination: day 0 (1st vaccine), day 7, day 21 (2nd vaccine), days 28, 35, 90, 182 and, 365. In the treatment branch, LPT was performed on the day of each vaccination and the following day for a total of 4 treatments. Blood draws were performed immediately prior to both vaccine administration and treatment with LPT. Blood samples were processed, and serum biobank is created. Serum anti-spike antibody levels are to be determined using quantitative ELISA. All recruited participants were over the age of 18 and were not vaccinated. The study was approved by the Western Institutional Review Board.

Summary of Results Recruitment has been successful and is ongoing. Currently, 96 participants have been recruited. 12 participants have dropped out for various reasons, leaving 84 continuing participants. To address attrition, compensation has been changed from a total of $100 to $200, dispersed as $25 per blood draw. Attrition rate before the compensation change was 30% (6/20) but reduced to 7.9% (6/76) after the change. Total retention rate is 87.5% with 53.6% (45/84) of participants enrolled. Recruitment has been successful and is ongoing.

Conclusions The ongoing study is still recruiting participants, however significant progress has been made, with 84 participants currently on board. Additionally, retention and the diversity of the participant population is promising with significant representation of Latinx individuals (55.9%) and females (58.3%).

Infectious diseases I

Concurrent session

12:45 PM

Thursday, January 20, 2022

#45 ANTI-DEPRESSANTS AND COVID-19 SEVERITY: A RETROSPECTIVE STUDY OF HOSPITALIZED ADULT PATIENTS

1SH Rauchman*, 2S Mendelson, 3C Rauchman, 4A Pinkhasov, 5U Kasselman, 3AB Reiss. 1Fresno Institute of Neuroscience, Fresno, CA; 2Providence Holy Cross Medical Center, Mission Hills, CA; 3NYU Long Island School of Medicine, Mineola, NY

10.1136/jim-2022-WRMC.45

Purpose of Study The SARS-CoV2 virus continues to have devastating consequences worldwide. Though vaccinations have
helped to reduce the impact of the virus, new strains still pose a threat to the unvaccinated, and to a lesser extent vaccinated, individuals. Therefore, it is imperative to identify treatments to reduce the severity of Covid-19. Recently, acute use of selective serotonin reuptake inhibitor (SSRI) antidepressants in COVID+ patients has been shown to reduce the severity of symptoms compared to placebo. Since SSRIs are a widely used anti-depressant, the aim of this study was to determine whether COVID+ patients already on SSRI treatment upon admission to the hospital had reduced mortality compared to COVID+ patients not on chronic SSRI treatment.

Methods Used A retrospective observational study design was used. Electronic medical records of 9,044 patients with a laboratory-confirmed diagnosis of Covid-19 from 03/2020 to 03/2021 from six hospitals were queried for demographic information, admission date; discharge date and disposition; length of stay; admission diagnoses; medications on admission; comorbidities; age; gender; ethnicity; admission to ICU; ventilator use; supplemental oxygen; oxygen saturation; discontinuation of antidepressant medications upon ICU admission.

Using R, a logistic regression model was run with mortality as the outcome and SSRI status as the exposure. An adjusted logistic regression model was run to account for R age category, gender, and race. All tests were considered significant at p of 0.05 or less.

Summary of Results In this sample, no patients admitted on SSRIs had them discontinued. This is consistent with current recommendations. There was no significant difference in the odds of dying between COVID+ patients on chronic SSRIs vs COVID+ patients not taking SSRIs, after controlling for age category, gender, and race. The odds of COVID+ patients on chronic SSRIs dying was 0.90 (95%CI: 0.74, 1.09; n=832) compared to COVID+ patients not on SSRIs (p=0.29; n=8211).

Conclusions In times of pandemics due to novel infectious agents it is difficult, but critical to evaluate safety and efficacy of drugs that might be repurposed for treatment. This large sample size of 9,044 patients suggests that there will be no significant benefit to use of SSRIs to decrease mortality rates for hospitalized patients with Covid-19 who are not currently on SSRI medications. This study shows the utility of large clinical databases in addressing the urgent issue of determining what commonly prescribed drugs might be useful in treating COVID-19.

#46 T-CELL AND ANTIBODY RESPONSES TO MRNA VACCINATION IN SARS-COV-2-CONVALESCENT SUBJECTS

1S Warner, 1E L Campbell, 1S Selke, 1DM Koelle. 1University of Washington School of Medicine, Seattle, WA; 2Fred Hutchinson Cancer Research Center, Seattle, WA

Purpose of Study Natural and vaccine-induced immunity are important for SARS-CoV-2 control. We evaluated SARS-CoV-2 specific T cell-mediated immune responses in COVID-19 survivors followed through vaccination. We compared T cell tests from Oxford Immunotec (OI) with an in-house laboratory-developed test (LDT). Each used peptides covering Spike (S) and non-vaccine proteins within SARS-CoV-2. We hypothesized that T cell responses to S will increase after mRNA vaccination. We compared vaccine immune boost in persons previously hospitalized vs. non-hospitalized for COVID-19, and the relationship between T cell and neutralizing antibody (nAb) responses.

Methods Used 20 subjects (median age 62.7, 50% female, most White) with PCR-confirmed SARS-CoV-2 infection donated plasma and peripheral blood mononuclear cells (PBMCs). Samples were from a median of 49 days after recovery from COVID-19 (V0), just prior to the 1st vaccination (E01), and 2-4 weeks after each mRNA dose (E02 and E03). T cell responses were measured by interferon-gamma enzyme-linked immunospot assays (ELISPOT). We compared our LDT assay using the S and nucleocapsid (N) proteins with the OI assay including the S, N, and matrix (M) proteins. nAb levels were measured by fluorescence inhibition. Linear regression was used to assess correlation between tests. Wilcoxon matched-pairs signed rank tests were used to evaluate differences in immune responses over time. Mann-Whitney tests were used to examine differences between hospitalized and non-hospitalized groups.

Summary of Results Strong correlation was noted between LDT and OI results for S protein at each time point (rho = 0.88, 0.85, 0.60, and 0.77, respectively). Between V0 and E02 (median of 327 days) and V0 and E03 (median of 345.5 days), there were significant increases in S-specific T cell responses (p = 0.0005 and 0.0006, respectively). No additional boost between E02 and E03 (p = 0.54) was observed. Low level (V0) responses to N and M were not boosted with vaccination. No significant difference in S-specific T cell responses between hospitalized and non-hospitalized groups were noted. For both hospitalized and non-hospitalized persons, nAb levels increased significantly after 1st dose of vaccine (p < 0.0001), with no additional nAb increase after the 2nd dose. No correlation between nAb and S-specific T cell responses at either V0 or E03 was noted.

Conclusions The OI assay is suitable for assessing T cell responses to SARS-CoV-2 mRNA vaccines. T cell responses to N and M did not boost, as expected. In this cohort, primary infection severity did not impact vaccine responses 9 months later. nAb and T cell response increases were complete after one dose. This indicates that a second dose may not be needed, at least if given 3-4 weeks after the first in persons recovering from COVID-19 in the prior 9 months.
vulnerability. Limited data describes how COVID-19 severity shapes maternal & infant ab responses. Stratification by severity can help characterize the protection to the newborn. The purpose of this study is to investigate maternal SARS-CoV-2 ab concentrations during L&D by disease severity, and compare infant ab responses at birth when exposed to varying severity in utero.

Methods Used This project is part of the prospective observational cohort study COVID19 Outcomes in Mother-Infant Pairs, analyzing mother-infant dyads in the US & Brazil. Serology of 101 pregnant women in Los Angeles (delivery: April 15, 2020-May 28, 2021) were analyzed and confirmed SARS CoV-2 PCR+ during pregnancy. Maternal blood at L&D, cord blood, and infant blood at birth were analyzed by ELISA for IgA, IgG & IgM (anti-spike receptor binding domain).

Summary of Results For 101 women, 72 had matched cord blood & 86 infant specimens. 76% of women produced all 3 anti-SARS-CoV-2 IgG, IgM, and IgA; 93% had at least one positive ab class; 5% had no detectable abs. Infant serum at birth contained only IgG and no IgM or IgA. With increased duration between onset of infection & delivery, maternal IgG levels waned, and conversely, transplacental transfer ratios increased (R²=0.27). Maternal IgG levels increased with disease severity. A significant increase in infant IgG levels was observed in children born to symptomatic mothers vs asymptomatic mothers (p<0.0001). A trend towards more robust ab responses was observed in infants with severe/critical COVID-19 exposure in utero (p=0.07).

Abstract #47 Table 1 Demographics and clinical characteristics of mother-infant dyads infected with SARS-CoV-2 during pregnancy

<table>
<thead>
<tr>
<th>Maternal Demographics and Medical History</th>
<th>All Women (N = 101)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, Median (Range)</td>
<td>33 (16–42)</td>
</tr>
<tr>
<td>Race/Ethnicity</td>
<td>No. (%)</td>
</tr>
<tr>
<td>Latina</td>
<td>46 (45.5)</td>
</tr>
<tr>
<td>White</td>
<td>27 (26.7)</td>
</tr>
<tr>
<td>Black/African American</td>
<td>8 (7.9)</td>
</tr>
<tr>
<td>Asian/Other</td>
<td>20 (19.8)</td>
</tr>
<tr>
<td>Insurance</td>
<td>No. (%)</td>
</tr>
<tr>
<td>Public</td>
<td>36 (35.6)</td>
</tr>
<tr>
<td>Private</td>
<td>65 (64.4)</td>
</tr>
<tr>
<td>Gravidity, Median (Range)</td>
<td>2 (1–10)</td>
</tr>
<tr>
<td>COVID-19 Severity</td>
<td>No. (%)</td>
</tr>
<tr>
<td>Asymptomatic</td>
<td>14 (13.9)</td>
</tr>
<tr>
<td>Mild/Moderate</td>
<td>76 (75.2)</td>
</tr>
<tr>
<td>Severe/Critical</td>
<td>11 (10.0)</td>
</tr>
<tr>
<td>Gestational Age at Diagnosis</td>
<td>No. (%)</td>
</tr>
<tr>
<td>First Trimester</td>
<td>10 (9.9)</td>
</tr>
<tr>
<td>Second Trimester</td>
<td>39 (38.6)</td>
</tr>
<tr>
<td>Third Trimester</td>
<td>52 (51.5)</td>
</tr>
<tr>
<td>Diagnosis Date-to-Delivery Interval, Median (IQR), Days</td>
<td>62 (32–120)</td>
</tr>
<tr>
<td>Medical History Prior to Pregnancy</td>
<td>No. (%)</td>
</tr>
<tr>
<td>Any Comorbidities</td>
<td>52 (51.5)</td>
</tr>
<tr>
<td>Obesity (Pre-Pregnancy BMI &gt;30)</td>
<td>32 (31.7)</td>
</tr>
<tr>
<td>Diabetes Mellitus (Not Gestational)</td>
<td>3 (3.0)</td>
</tr>
<tr>
<td>Congenital Heart Disease</td>
<td>5 (5.0)</td>
</tr>
<tr>
<td>Asthma</td>
<td>12 (11.9)</td>
</tr>
</tbody>
</table>

Conclusions Our findings demonstrate how altered maternal responses across distinct COVID-19 disease severity categories influence neonatal protection against SARS CoV-2.

Abstracts

#48 BIOMARKERS PREDICTIVE OF MORTALITY IN COVID-19 PATIENTS WITH DIAGNOSED HEART FAILURE

1T Nguyen*, 1JY Hwang, 1C Lee, 2T Buck, 2K Mun, 2IT Vanteru, 2A Lu, 2D Tirschwell, 1A Kim. 1Washington State University. 2Elson S Floyd College of Medicine, Spokane, WA; 2University of Washington School of Medicine, Seattle, WA

10.1136/jim-2022-WRMC.48

Purpose of Study The purpose of this study is to identify common biomarkers and biosignals in COVID-19 patients with heart failure that are associated with increased risk of in-hospital mortality. COVID-19 is associated with worse outcomes in patients with pre-existing comorbidities, such as heart failure (HF). Biomarkers such as B-type natriuretic peptide, troponin, and interleukin-6 have been elevated in patients with HF and COVID-19 and may provide insight on the severity of disease but may not be collected in all patients. The exact association between patients with prior HF and the biomarkers commonly utilized is limited and should be evaluated further. Our study evaluates biosignals and biomarkers that may be predictive of mortality in COVID-19 patients with history of HF.

Methods Used All patients included were 18 years of age or older, diagnosis of COVID-19 was confirmed by PCR test or hospital clinical criteria and were hospitalized in the University of Washington (UW) Medicine hospital systems between February 2020 to December 2020. The data was collected as part of a national effort for the American Heart Association COVID-19 CVD Registry. The biosignals that were analyzed include temperature, heart rate, respiratory rate, diastolic blood pressure, and systolic blood pressure. The biomarkers include admission white blood cell (WBC) count, platelets, serum creatinine, AST (u/L), ALT (u/L), and lymphocyte count. Patient data also tracked previous medical history and disposition at discharge. A LASSO multivariate regression model was used to identify the variables most predictive of mortality among patients with heart failure.

Summary of Results The study included 54 of the 393 COVID-19 patients (13.7%) with previous diagnosis of heart failure (46% male, mean age 77). Our model estimates that for each standard deviation unit above average (z-score), patients with previously diagnosed heart failure were 13% more likely to die due to COVID-19 (p = 0.021). Among patients with prior heart failure, each z-score increase for WBC count and serum creatinine increased risk of mortality by 3.5% (p = .043) and 5.7% (p = .046), respectively.

Conclusions Our data suggests that there may be significance in monitoring WBC count and serum creatinine levels among COVID-19 patients with prior heart failure. The WBC count and serum creatinine have a stronger relationship to mortality in patients with prior heart failure compared to those without heart failure. Immune response may be reduced in heart failure patients which can account for the decreased WBC count, but further studies are needed to elucidate the exact mechanism and relationship. The results of this study may provide a roadmap to triage heart failure patients based on admission lab values in the COVID-19 environment.
**Purpose of Study** The national rate of congenital syphilis (CS) has dramatically increased recently. It remains unknown if the children of the agricultural worker population (AWP) are more susceptible to CS in California. Identifying subpopulations vulnerable to transmitting CS may inform the design of intervention efforts. Thus, this study set out to determine whether CS incidence rates are associated with the female AWP in California.

**Methods Used** Data from all 58 California counties were retrospectively obtained from the California Department of Public Health and United States Department of Agriculture regarding CS incidence per 100,000 live births and female AWP from December 2014 and December 2018. Female AWP per county was estimated according to the national proportion of female to male agricultural workers provided by the Department of Agriculture. Data was analyzed using geographical information systems mapping and Pearson’s correlation coefficient (r) tests.

**Summary of Results** The average statewide CS incidence was 68.2 cases per 100,000 live births in 2018. CS incidence and female AWP were concentrated heavily in California’s agricultural Central Valley, with a few coastal exceptions (figure 1A-C). CS incidence and female AWP were moderately but significantly correlated (r = 0.343; 95% confidence interval = 0.093–0.552; p < 0.001) (figure 1D).

**Conclusions** Our findings provide evidence that California counties with a higher incidence of CS tend to be home to a greater number of female agricultural workers than counties with low incidence of CS. Given these findings, this study suggests the urgent need to implement culturally appropriate and enduring prenatal healthcare interventions that prioritize treatment of maternal syphilis and prevention of CS in female AWP.
strategies may be developed to prevent human infection. Pathogen recognition receptors (PRRs) are part of the first line of defense against pathogens. They are hypothesized to be under balancing selection due to selection pressure on pathogens to evolve novel epitopes to evade immune recognition and on host receptors to detect pathogens. Thus, PRRs and other immune loci are expected to be among the most diverse regions of the genome. Using whole genome data from an important African snail vector, Biomphalaria sudanica, we hypothesize that diverse regions of the genome will be enriched with immune related loci, and that we can identify novel PRRs through annotation of these regions.

Methods Used Five B. sudanica strains (collected: Lake Victoria, Kenya) were sequenced using the PacBio and Illumina-paired-endreads. Mean inter-line diversity was calculated across the genomes and segmented into smaller windows (10–100kb). Each window with a mean inter-line diversity value >1%, had up to 1Mb of surrounding nucleotides annotated and trans-membrane domains (TMDs) identified using predictive software. To determine if TMD peptides were over-represented in high-diversity regions of the genome, we compared the proportion of TMD peptides against the proportion of those in 30 randomly assigned contig regions.

Summary of Results 67 of 6815 windows met our nucleotide diversity threshold of 1% divergence. 421 of 818 immune-suspected peptides were identified to have TMDs, over-represented in regions of high diversity when compared to randomized control regions, supporting our hypothesis. Immune-related genes associated with Schistosoma resistance in other species including PTC2 and GRC were also identified using this bioinformatic approach.

Conclusions Our findings support the diversity-based approach to identifying PRRs which successfully identified known B. glabrata PRRs and novel PRRs in B. sudanica genomes. Our established list of candidate genes for pathogen recognition will provide a foundation guiding resistance studies, gene knockout and GWAS with Biomphalaria species.

Abstract #52 Table 1 Studies evaluating the association between vitamin D level and COVID-19 disease severity

<table>
<thead>
<tr>
<th>Author, year, and location</th>
<th>Controls: number (N) and mean or median age (ys)</th>
<th>Subjects: number (N) and mean or median age (ys)</th>
<th>Serum 25(OH)D concentration in controls vs. subjects, p-value</th>
<th>Outcome based on comorbidities and vitamin D levels, p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Radujkovic, 2020, Germany</td>
<td>Controls: COVID-19 outpatients, N=92, Median age=56</td>
<td>Subjects: COVID-19 inpatients, N=93, Median age=63</td>
<td>Serum 25(OH)D concentration in controls vs. subjects, p-value</td>
<td>Outcome based on comorbidities and vitamin D levels, p-value</td>
</tr>
<tr>
<td>Luo, 2020, China</td>
<td>Controls: Non-severe COVID-19, N=261, Median age=54.0</td>
<td>Subjects: Severe COVID-19, N=74, Median age=62.5</td>
<td>Remove 25(OH)D levels significantly predictive of in-hospital mortality</td>
<td>Multivariate analysis: OR 0.927, 95%CI 0.875–0.982, p=0.01.</td>
</tr>
<tr>
<td>Karahana, 2021, Turkey</td>
<td>Controls: Moderate COVID-19, N=47, Mean age=56.1</td>
<td>Subjects: Severe-critical COVID-19, N=102, Mean age=67.0</td>
<td>Serum 25(OH)D concentration in controls vs. subjects, p-value</td>
<td>Outcome based on comorbidities and vitamin D levels, p-value</td>
</tr>
<tr>
<td>Jain, 2020, India</td>
<td>Controls: Asymptomatic COVID-19, N=91, Mean age=42.34</td>
<td>Subjects: Severe COVID-19, N=63, Mean age=51.4</td>
<td>Serum 25(OH)D concentration in controls vs. subjects, p-value</td>
<td>Outcome based on comorbidities and vitamin D levels, p-value</td>
</tr>
<tr>
<td>Macaya, 2020, Spain</td>
<td>Controls: Non-severe COVID-19, N=49, Mean age=63</td>
<td>Subjects: Severe COVID-19, N=31, Median age=75</td>
<td>Serum 25(OH)D concentration in controls vs. subjects, p-value</td>
<td>Outcome based on comorbidities and vitamin D levels, p-value</td>
</tr>
<tr>
<td>Ye, 2020, China</td>
<td>Controls: Mild/moderate COVID-19, N=50, Mean age=39</td>
<td>Subjects: Severe-critical COVID-19, N=10, Mean age=65</td>
<td>Serum 25(OH)D concentration in controls vs. subjects, p-value</td>
<td>Outcome based on comorbidities and vitamin D levels, p-value</td>
</tr>
<tr>
<td>Kerget, 2020, Turkey</td>
<td>Controls: COVID-19 without ARDS, N=53, Mean age=38.3</td>
<td>Subjects: COVID-19 with ARDS, N=35, Mean age=67.9</td>
<td>Serum 25(OH)D concentration in controls vs. subjects, p-value</td>
<td>Outcome based on comorbidities and vitamin D levels, p-value</td>
</tr>
<tr>
<td>Campi, 2021, Italy</td>
<td>Controls: Severely symptomatic COVID-19 hospital ward admits, N=49, Mean age=68.82</td>
<td>Subjects: Severely symptomatic COVID-19 ICU admits, N=54, Mean age=63.67</td>
<td>Serum 25(OH)D concentration in controls vs. subjects, p-value</td>
<td>Outcome based on comorbidities and vitamin D levels, p-value</td>
</tr>
</tbody>
</table>

Units for vitamin D levels in two studies (Luo et al. and Ye et al.), originally reported in nmol/L, were converted to ng/mL in order to achieve consistency in the units across all studies. 25(OH)D=25-hydroxyvitamin D, VDD=Vitamin D Deficiency, HR=Hazard Ratio, OR=Odds Ratio, RR=Relative Risk, NR=Not Reported
Abstracts

Summary of Results Eight studies satisfied our inclusion criteria (see table 1 below). A majority of studies showed significantly lower vitamin D levels in the more severe COVID-19 subjects compared to controls with less severe COVID-19. Most studies that assessed COVID-19 outcomes based on vitamin D deficiency (VDD) found VDD to be associated with worse outcomes, including more severe disease and increased mortality. The limitations of this review include inter-study variability in the co-morbidities included in multivariate analysis, variability in the definition of VDD among different studies, and a lack of information on vitamin D supplements and other treatments before infection or during hospitalization in several studies. Finally, a causal relationship could not be assessed because all studies were observational, and information on the vitamin D levels before hospital admission, during healthy state, was not available in a majority of studies.

Conclusions There may be an association between lower vitamin D levels and more severe COVID-19 disease. However, larger longitudinal studies that not only measure vitamin D levels pre-COVID-19 disease but also take into account all variables, such as comorbidities and treatments that could affect disease severity, are warranted.

#53 THE USE OF VAGINAL PROBIOTIC SUPPOSITORIES IN PREVENTION OF RECURRENT URINARY TRACT INFECTIONS IN ADULT WOMEN: A LITERATURE REVIEW

D Devineni*, 1B Bade, 1C Diyakonou, 1E Lee, 1N Lulla, 1K Parang, 1S Soni, 1B Alghani. 1University of California Irvine School of Medicine, Irvine, CA; 2Children’s Hospital of Orange County, Orange, CA

Purpose of Study Urinary tract infections (UTIs) are one of the most common bacterial infections in women. Concerns over the effectiveness of antibiotics in preventing recurrent UTIs, due to antibiotic resistance and the adverse effects of antibiotics on healthy microbiota, have raised the necessity to investigate reliable non-antibiotic treatments for preventing recurrent UTIs. It has been proposed that probiotics or lactobacilli may be effective in preventing infections by restoring the normal vaginal flora. The purpose of this study is to investigate the effectiveness of vaginal probiotic suppositories for prevention of recurrent UTIs in adult women.

Methods Used A systematic literature review was conducted through databases such as PubMed and Google Scholar. Only studies that were published after 1990, and compared use of vaginal probiotics with a control group in adult women with recurrent UTI were included. Studies with follow-up period of <6 months were excluded.

Conclusions We found 5 studies that fit our inclusion criteria (See table 1). In majority of the studies, the frequency of recurrent UTIs was lower in patients who received probiotic vaginal suppositories when compared to controls. However, there was great variability among the studies with respect to the probiotic formulation as well as treatment dose and frequency. The applications were intermittent and varied from daily to weekly to monthly. The probiotic species used in different studies included L. Rhamnosus, L. Fermentum, and L. Crispatus. The sample sizes were small and did not divide the patients into different categories based on risk factors or co-morbidities. In addition, the bacterial cause of UTI was not mentioned in majority of the studies. Mild side effects were noted in both probiotic and control group, and included increased vaginal discharge, vaginal odor, mild irritation and dysuria.

Conclusions Our review suggests a promising role for use of intermittent vaginal probiotic suppositories for prevention of recurrent UTIs in adult women. Larger prospective studies with longer follow-up period are needed to determine the optimal probiotic dosage and frequency in different groups of patients with recurrent UTI.

Neonatology general I

Concurrent session

12:45 PM

Thursday, January 20, 2022

#54 NUCLEATED RED BLOOD CELL EMERGENCE-TIME IN NEWBORN LAMBS FOLLOWING A DOSE OF DARBEPOETIN ALFA

1,2D Bahr*, 1,2R Albertine, 1,2D Christensen, 1MI Dahl, 1A Rebentsch, 1E Dawson, 2E Major, 2H Foreman, 2D Headden, 2V Vordos, 2A Nabi, 1L Pettet, 2P Badrov, 1C Addison, 2D Christensen. 1,2Intermountain Healthcare, Salt Lake City, UT; 2University of Utah Health, Salt Lake City, UT

Purpose of Study A high nucleated red blood cell (NRBC) count in a neonate at birth has been suggested as a biomarker for fetal hypoxia. However, it is not clear if it indicates when

Abstract #53 Table 1 Studies on the efficacy of intravaginal probiotics in preventing recurrent UTIs

<table>
<thead>
<tr>
<th>First author’s last name, Year of Publication, Location of study</th>
<th>Name of Probiotic</th>
<th>Duration and frequency of usage (intravaginal)</th>
<th>Number of Subjects and controls</th>
<th>Frequency of UTIs in subjects vs controls</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Reid, 1992, Canada</td>
<td>L casei rhamnosus and L fermentum</td>
<td>Twice a week for 2 weeks then once monthly</td>
<td>19 vs 21</td>
<td>21% vs 47%</td>
<td>p&lt;0.05</td>
</tr>
<tr>
<td>Baehrheim, 1994, Norway</td>
<td>L casei rhamnosus</td>
<td>Twice a week for 26 weeks</td>
<td>25 vs 22</td>
<td>Incidence ratio=1.4</td>
<td>95% CI=0.88 to 1.98 (p&lt;0.05)</td>
</tr>
<tr>
<td>Reid, 1995, Canada</td>
<td>L. rhamnosus and L. fermentum</td>
<td>Once a week 1 year</td>
<td>55 subjects, baseline was control</td>
<td>Epidemics decreased from 6 to 1.6yr</td>
<td>p&lt;0.001</td>
</tr>
<tr>
<td>Uehara, 2006, Tokyo</td>
<td>L. Cnipatus (Lactin-V)</td>
<td>Every 2 days for 1 year</td>
<td>9 Subjects, baseline was control</td>
<td>Recurrences decreased from 5.0yr to 1.3yr</td>
<td>p=0.0007</td>
</tr>
<tr>
<td>Stapleton, 2011, USA</td>
<td>L. Cnipatus (Lactin-V)</td>
<td>5 days, then weekly for 10 weeks</td>
<td>50 vs 50</td>
<td>15% vs 27%</td>
<td>p&lt;0.01</td>
</tr>
</tbody>
</table>
the hypoxia occurred. We aimed to measure the time between administering a high-dose of darbepoetin, simulating the marked rise in erythropoietin that follows a hypoxic event, and the first appearance of NRBC in the blood. Limited observations of this interval, the ‘NRBC emergence-time,’ in human neonates suggest it is greater than 24 hours.

**Methods Used**
We obtained serial blood counts on ten newborn lambs; five dosed with darbepoetin (10 μg/kg) and five placebo controls, to assess the NRBC emergence-time.

**Summary of Results**
The first appearance of NRBC was at 24 ± 72 h (16.758 ± 8.434/μL vs. 0/μL in controls), followed by fewer at 96 hours (7823 ± 7.114/μL vs. 0/μL in controls). Similarly, reticulocytes peaked at 48–72 h (113,094 ± 3210/μL vs. 10,790 ± 5449/μL in controls), with no changes in platelets or leukocytes.

**Conclusions**
The NRBC emergence time in newborn lambs is similar to reports from newborn humans. By extrapolation, if a neonate has a high NRBC at birth, the erythropoietic stimulus likely occurred within the interval 24 to 96 hours prior to birth.

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**REFERENCE INTERVALS FOR END-TIDAL CARBON MONOXIDE OF PRETERM NEONATES**

**Purpose of Study**
Every molecule of heme metabolized to bilirubin releases one molecule of carbon monoxide (CO). On that basis, hemolysis can be detected and quantified by measuring CO in exhaled breath. We constructed reference intervals for end-tidal carbon monoxide (ETCOc) levels of neonates 28–34 weeks gestation to assess the hemolytic rate. New instrumentation allows providers to non-invasively measure ETCOc in preterm neonates with low tidal volumes and breathing rates up to 70bpm. Reference intervals for term and late preterm neonates exist, but until now none have been created for preterm neonates.

**Methods Used**
Prospective four-NICU study in Bangkok, Thailand, and Utah, USA. Neonates born between 28–34 weeks and up to 28 days old were eligible. Once informed consent was received, a modified CoSense ETCOc analyzer was used to record results. Data from the CoSense devices were linked to patient charts to obtain demographic information.

**Summary of Results**
Values from days one through 28 were charted and upper (>95th percentile) reference interval limits were calculated. During the entire 28 days, the ETCOc upper reference intervals of babies in Bangkok were higher than those in Utah (p<0.01). No differences were found due to sex, or earliest vs. latest gestation at birth (both p>0.1). Preterm neonates in Bangkok and Utah had higher ETCOc values during the first 48 hours after birth than thereafter (p<0.01).

**Conclusions**
Using the reference interval chart we created, the hemolytic rate of preterm infants ≥28 weeks can be assessed. This identification allows us to focus subsequent testing to find the cause of the hemolysis, administer more intensive phototherapy, and to assure consistent in-and out-patient follow-up to those with hemolytic jaundice.
Summary of Results Of 299,927 live births 344 had severe anemia. In 153 (44.5%) the anemia was unrecognized during the first 24 hours. The lowest hemoglobin/hematocrit values were among those with hemorrhage vs. hemolysis (P<0.013) or vs. hypoproduction (P<0.001). In infants with severe anemia secondary to hemorrhage, abrupton/other perinatal event and fetomaternal hemorrhage (FMH) were the most likely etiologies. DIC was a common hemolytic cause of anemia, with 85% of DIC cases coincident with hemorrhagic anemia.

Conclusions Severe anemia at birth often went unrecognized on the first day. Earlier recognition may be facilitated by an electronic medical record-associated hemoglobin/hematocrit nomogram, with values <1st percentile clearly identified.

Purpose of Study We previously reported fetomaternal hemorrhage (FMH) in 1/9160 births, and only one neonatal death from FMH among 219,853 births. Recent reports indicate FMH is not uncommon among stillbirths. Consequently, we speculated we were missing cases among early neonatal deaths. We began a new FMH initiative to determine the current incidence.

Purpose of Study Perinatal anemia is a massive global public health burden with an estimated global prevalence of approximately 40%. Severe anemia increases the risk of maternal mortality and can adversely affect fetal development. Adequate correction of anemia is essential for a healthy pregnancy and infant, but universal screening and monitoring is not the care standard in most LMICs. In lieu of universal screening and treatment, providing access to Iron Folic Acid (IFA) tabs is considered an effective and cost-efficient intervention to prevent and treat anemia of pregnancy. However, despite widespread availability of IFA tabs, anemia prevalence continues to be high and the presence of IFA programs may falsely reassure clinicians that patients taking them have adequate hemoglobin.
Methods Used The study took place at Mota Fofalia Community Health Center (MF-CHC) in Gujarat, India operated by a public-private partnership. The University of Utah operates an academic global health program in collaboration with MF-CHC and assists the health center in sustainable capacity building in maternal-child health. As part of a community-based antenatal care (ANC) program, we recruited a cohort of pregnant women from the surrounding community to complete a standardized nutrition and health survey and participate in scheduled prenatal visits according to WHO and Indian ANC guidelines which include measurement of vital signs and ANC guideline-based interventions. At each ANC visit, a blood hemoglobin level was drawn and each participant was asked if they are currently taking IFA or Albendazole, an antiparasitic.

Summary of Results A total of 501 women were included in the study. 448 (89%) report taking IFA and 53 (11%) report not taking IFA. The average hemoglobin for those taking IFA was 10.11 g/dL (IQR 9.3–11.1) with an average gestational age at screening of 23.0 weeks while the average of those not taking was 10.41 g/dL (IQR 9.8–11.6) (p = .28) with an average gestational age of 10.9 weeks. In the group taking IFA tablets, 97% were also taking Albendazole while only 21% of mothers not taking IFA tablets were taking Albendazole.

Conclusions In areas with a high prevalence of anemia, patient compliance with standard IFA antenatal therapy is not an adequate indicator of intervention. While it appears many mothers begin taking IFA as they become pregnant, the presence of readily available IFA therapy to the community is not sufficient in addressing perinatal anemia.

Abstract #59 Table 1 Correlation between CBC HB and paired POC/blood gas HB

<table>
<thead>
<tr>
<th>Sample size</th>
<th>Mean Difference</th>
<th>Standard Deviation of Mean Difference</th>
<th>Pearson Coefficient (r)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>CBC vs POC Gas within 12 hours</td>
<td>488</td>
<td>1.1</td>
<td>1.0</td>
<td>0.89, p &lt; 0.05</td>
</tr>
<tr>
<td>CBC vs POC Gas between 12–24 hours</td>
<td>243</td>
<td>1.3</td>
<td>1.0</td>
<td>0.83, p &lt; 0.05</td>
</tr>
<tr>
<td>CBC vs POC Hgb within 12 hours</td>
<td>477</td>
<td>1.0</td>
<td>1.0</td>
<td>0.87, p &lt; 0.05</td>
</tr>
<tr>
<td>CBC vs POC Hgb between 12–24 hours</td>
<td>290</td>
<td>1.3</td>
<td>1.2</td>
<td>0.75, p &lt; 0.05</td>
</tr>
</tbody>
</table>

Methods Used This is a retrospective study of patients admitted to the LAC+USC Medical Center NICU between January 2020 and April 2021 with paired HB measurements from the laboratory-run CBC and either POC HB from HemoCue B 201 or blood gas HB from Gem Premier 5000. Qualifying data was divided into groups based on time between different blood draws, the first comprising of measurements collected within 12 hours of each other, the second of measurements collected between 12 and 24 hours of each other. POC or blood gas HB and CBC HB measurements collected over 24 hours from each other were excluded. T-tests were used for analysis of continuous, normally distributed variables. Regression analysis was performed to determine the relationship between paired HB measurements. Statistical significance was set at p < 0.05.

Summary of Results We identified 250 subjects with qualifying paired HB values from the CBC and POC HB or blood gas HB. There were 488 paired CBC and blood gas HB samples in the <12 hour group, and 243 paired samples in the 12–24 hour group. There were 479 paired CBC and POC HB samples in the <12 hour group, and 290 paired samples in the 12–24 hour group (table 1). Correlation coefficient (R) for the CBC-blood gas HB groups were 0.89 in the <12 hour group and 0.83 in the 12–24 hour group. Correlation coefficient for the CBC-POC HB groups were 0.87 in the <12 hour group and 0.75 in the 12–24 hour group (figure 1).

Conclusions There was a strong correlation between paired POC HB or blood gas HB and CBC HB values obtained within 12 hours of one another. Our results show that POC HB or blood gas HB should be considered as alternatives for CBC HB. The patients in the NICU would benefit in prevention of iatrogenic anemia. Prospective studies with age- or weight-based grouping and planned pairing within pre-defined time periods would be beneficial in determining whether the correlation persists between POC and CBC HB measurements in the NICU.
**Abstracts**

**#60** ALTERNATIVE WAYS OF ACQUIRING BILIRUBIN MEASUREMENT IN PRETERM AND TERM INFANTS ADMITTED TO NEONATAL INTENSIVE CARE UNIT

D Cho*, K Ramm, L Barton, R Ramanathan, M Biniwale. LAC+USC Medical Center, Los Angeles, CA

10.1136/jim-2022-WRMC.59

**Purpose of Study** Measuring bilirubin levels in infants admitted to neonatal intensive care unit is done to avoid hyperbiliru- biniemia and bilirubin toxicity. Practitioners strive to minimize tests to reduce patient discomfort and iatrogetic anemia. To minimize blood draws in the monitoring of bilirubin levels, we assessed the accuracy of alternative methods of measurement via blood gas analyzers and transcutaneous bilirubin monitoring.

**Methods Used** Using serum bilirubin as a gold standard, we analyzed the accuracy of simultaneous measurements from blood gas analyzer and transcutaneous monitoring. The accuracy of blood gas and transcutaneous bilirubin measurements was ascertained with correlation coefficient and by calculating mean differences between the serum bilirubin levels and the two alternative methods.

**Summary of Results** Study consisted of 86 patients with gesta- tional ages 24 to 41 weeks. The correlation coefficient for serum bilirubin vs transcutaneous measurements was \( r = 0.893 \) (\( p<0.00001 \)). The correlation coefficient was \( r = 0.9283 \) (\( p<0.00001 \)) for preterm infants, and \( r = 0.8392 \) (\( p<0.000013 \)) for term infants. The mean difference between serum bilirubin vs blood gas bilirubin was 0.45 with a standard deviation of 1.55 mg/dL. The correlation coefficient for serum bilirubin vs blood gas bilirubin was \( r = 0.959 \) (\( p=0.00001 \)). The correlation coefficient was \( r = 0.9291 \) (\( p=0.00001 \)) for preterm infants, and \( r = 0.9742 \) (\( p=0.00001 \)) for term infants. The mean difference between serum bilirubin vs blood gas bilirubin was 0.21 with a standard deviation of 0.87 mg/dL.

**Conclusions** Both transcutaneous and blood gas analyzer bilirubin levels had a strong correlation with serum levels, with bilirubin using blood gas analyzer being slightly more accurate. We plan to continue collecting bilirubin data for a total of 6 months. If accuracy of transcutaneous or blood gas analyzer bilirubin measurements are acceptable, we plan to pursue these alternative methods of bilirubin measurements over the following 6 months and assess to what extent we were able to minimize blood draws.

**Abstract #60 Figure 1**

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**#61** PREDICTORS OF HOME OXYGEN THERAPY IN VERY LOW BIRTH WEIGHT INFANTS

Y Shao*, A Hisey, N Nanduri, K Ramm, C Marquez, L Barton, R Ramanathan, M Biniwale. Los Angeles County University of Southern California Medical Center, Los Angeles, CA

10.1136/jim-2022-WRMC.60

**Purpose of Study** Very low birth weight (VLBW) infants with prolonged respiratory morbidity such as bronchopulmonary dysplasia (BPD) may need to be discharged home while receiving oxygen therapy. The risk factors causing prolonged respiratory support vary for these infants. The study was performed to characterize factors associated with VLBW infants who require oxygen therapy at discharge.

**Methods Used** Data on all VLBW infants was gathered from the electronic medical record between the years of 2009 and 2021 retrospectively with IRB approval obtained prior. Oxygen therapy was statistically analyzed using SPSS Version 28 statistical software against early neonatal outcomes such as ventilation, intubation, chest compressions, or surfactant, or against common neonatal morbidities, including IVH, BPD, ROP, and patent ductus arteriosus (PDA).

**Summary of Results** Of 560 VLBW infants, 144 (25.7%) were discharged home on oxygen. Significant maternal risk factors

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**Abstract #61 Table 1** Home oxygen therapy requirement against common neonatal outcomes

<table>
<thead>
<tr>
<th></th>
<th>Infants requiring home oxygen</th>
<th>Infants not requiring home oxygen</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean Birth Weight (g)</td>
<td>795</td>
<td>1077</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Mean Gestational Age (weeks)</td>
<td>26</td>
<td>29</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Histologic Chorioamnionitis (%)</td>
<td>19.5</td>
<td>6.5</td>
<td>0.003</td>
</tr>
<tr>
<td>Intubation in Delivery (%)</td>
<td>56.0</td>
<td>30.8</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Room (%)</td>
<td>59.0</td>
<td>30.8</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Chest Compressions (%)</td>
<td>18.0</td>
<td>7.0</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Surfactant Therapy (%)</td>
<td>50.7</td>
<td>39.4</td>
<td>0.024</td>
</tr>
<tr>
<td>Invasive Ventilation at 24 hours (%)</td>
<td>70.3</td>
<td>24.9</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Duration of Invasive Ventilation (days)</td>
<td>33</td>
<td>9</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>PDA (%)</td>
<td>84.7</td>
<td>52.8</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>PDA Requiring Surgery (%)</td>
<td>41.9</td>
<td>6.5</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Severe IVH (%)</td>
<td>8.1</td>
<td>1.5</td>
<td>0.014</td>
</tr>
<tr>
<td>BPD (%)</td>
<td>91.4</td>
<td>35.5</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Length of Stay (days)</td>
<td>81</td>
<td>59</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

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142 J Investig Med 2022;70:110–345
included histologic chorioamnionitis, intubation, and chest compressions at delivery. These infants had lower gestational age as well as lower birth weight. These infants were more likely to require surfactant, invasive ventilation at 24 hours, and receive additional ventilatory support including high-frequency oscillatory ventilation or jet ventilation (table 1). Infants who were discharged with home oxygen therapy were significantly more likely to have BPD and hemodynamically significant patent ductus arteriosus (PDA) requiring surgery. Associated comorbidities included retinopathy of prematurity (ROP) requiring treatment, severe intraventricular hemorrhage (IVH), and increased length of stay in NICU (table 1). Regression analysis revealed lower birth weight, longer duration of invasive ventilation, PDA requiring surgical intervention, and BPD to be the most significant predictors.

Conclusions The need for high-frequency ventilation such as jet and oscillatory ventilation, as well as chest compressions, is associated with home oxygen need at discharge. Additionally, VLBW infants who required home oxygen therapy were more likely to have needed invasive ventilation, delivery room intubation, and surfactant therapy. They were also more likely to have common neonatal morbidities such as BPD, severe IVH, severe ROP, and PDA.

**Abstracts**

## #62 CONSEQUENCES OF MATERNAL BREAST MILK ANTIBODIES ON THE ABUNDANCE OF INTESTINAL MICROBES IN NEONATES

**1H. Wolf**, 2S. Wang, 3M. Koch, 1University of Washington School of Medicine, Seattle, WA; 2Fred Hutchinson Cancer Research Center, Seattle, WA

**Purpose of Study** Breast milk is an important contributor of the neonatal microbiome. Studies have associated breastfeeding with a decreased risk of acquiring inflammatory bowel disorders later in life. As breastfeeding is not possible for all mothers and children, gaining a mechanistic understanding of this process can lead to the development of early-life interventions that foster beneficial host-microbiota relationships.

We have previously shown that breast milk antibodies are important for maintaining mucosal homeostasis. Mice deficient in breast milk antibodies exhibit perturbations in mucosal immunity, including elevated T follicular helper (Tfh) cell and germinal center (GC) B cell responses in the gut-associated lymphoid tissues. We hypothesize that the Tfh and GC B cell response generated by the neonate in the absence of breast milk antibodies target resident mucosal bacteria and lead to long-term alterations in the abundance of gut-microbiota.

**Methods Used** We extracted DNA from intestinal-wall associated microbes and quantified the abundance using quantitative-PCR of the bacterial 16s rRNA. We looked at both the small intestinal wall microbes and large intestinal wall microbes at ages 3, 5 and 11 weeks in maternal antibody sufficient or deficient pups. To explore the effects of Tfh cells and GC B cells, we treated half of each group with anti-Inducible T-cell Co-Stimulator Ligand (anti-ICOSL) antibody, which blocks Tfh and GC B cell formation. We used a paired T-test to determine the significance of our results.

**Summary of Results** We found no significant difference in the abundance of wall-associated microbes across all four groups in the small or large intestine. These data suggest that the breast milk antibody dependent response as well as the Tfh-cell dependent response do not alter the abundance of wall-associated microbes.

**Conclusions** Studies are ongoing to determine if breast milk antibodies affect intestinal microbe composition. Limitations to this study may include the drinking water our mice were exposed to throughout the duration of this experiment. Due to mouse husbandry standards operations within our institution our mice were given acidified water which we have recently discovered to significantly alter their microbial composition. Our goal is to repeat this experiment with non-acidified water as we suspect any potential differences in the microbial abundance may have been masked by the effects of acidified water.

## Neonatology pulmonary I

### Concurrent session

**12:45 PM**

**Thursday, January 20, 2022**

## #63 EFFECTS OF GESTATIONAL LONG-TERM HYPOXIA ON THE COLLAGEN MATRIX OF FETAL SHEEP PULMONARY ARTERIES

**1RC Torres Chavez, 2C. Gheonoge, 1L. Zhang, 3M. Yellon S, 2S. Wilson, 1Loma Linda University School of Medicine, Loma Linda, CA; 2Loma Linda University Department of Basic Sciences, Loma Linda, CA**

10.1136/jim-2022-WRMC.63

**Purpose of Study** Gestational-long term hypoxia (gLTH) is a significant stressor that leads to multiple diseases including pulmonary hypertension. Evidence indicates that gLTH causes oxidative stress and inflammation, which changes cell structure and function. These effects are driven by changes in cellular metabolism, protein expression, and transcriptional regulation. Our proteomic data show that gLTH leads to vascular remodeling and specifically to reduction of collagen 1A1, 1A2, and 3A1, though the data do not delineate where in the arterial wall these changes are occurring. We hypothesized that gLTH causes loss of collagen in all arterial layers which was tested by visualizing and quantifying the collagen content in different layers of fetal pulmonary arteries.

**Methods Used** Fetal sheep pulmonary arteries from normoxic and gLTH environments were obtained and stained with picrorius red dye (PSR) to visualize collagen in captured images of arterial biopsies by assessing the optical density (OD) of birefringence from polarized light. Fluorescence microscopy was used to capture images of the arterial samples. Data from Image J analysis of OD birefringence, inversely related to crosslinked collagen, of the various vascular layers and treatments were evaluated by analysis of variance.

**Summary of Results** The image analysis showed a significant decrease in optical density, and therefore enhanced crosslinking, in the adventitia compared to the media for all samples belonging to either normoxic or gLTH groups. However, there was no significant difference in optical density of the adventitial versus medial layers between vessels from normoxic and gLTH fetuses.

**Conclusions** The results indicate that optical density quantification can be used to detect substantial differences in collagen...
and crosslinked structure between the medial and adventitial layers. The findings also raise the possibility that neither collagen nor its crosslinked structure may be affected by gLTH. The data provides evidence that this technique needs refinement to properly visualize the locations where modest changes in expression may occur. Secondarily, the inability of PSR stain to distinguish among certain types of collagen subtypes leaves open the possibility that a shift in the type of collagen may affect biomechanical processes that are associated with pulmonary vascular development or gLTH. The PSR red staining method may not have been definitive, but this study is an important steppingstone towards developing an experimental strategy of visualizing modifications in vascular collagen isoform expression that complement contemporary analytical quantification techniques that provide unique insight into vascular structure and function.

#64
MESENCHYMA STROMAL CELL EXTRACELLULAR VESICLES IMPROVE ALVEOLAR FORMATION IN MECHANICALLY VENTILATED PRETERM LAMBS

E Major*, A Rebentisch, E Dawson, H Foreman, D Headden, Z Vordos, MJ Dahl, D Null, A Mitsialis, S Kouriembanas, K Albertine. 1The University of Utah School of Medicine, Salt Lake City, UT; 2University of California Davis, Davis, CA; 3University of California Davis Health System, Sacramento, CA; 4Harvard Medical School, Boston, MA

Purpose of Study Bronchopulmonary dysplasia (BPD) is histopathologically characterized alveolar simplification in preterm infants who are chronically mechanically ventilated. Mesenchymal stromal cell extracellular vesicles (MES) treatment improved alveolar formation in mouse neonatal hyperoxia models of BPD. We tested the hypothesis that MES will improve alveolar formation in chronically mechanically ventilated preterm lambs. Methods Used Preterm lambs (128d; term ~150d; ~28w human gestation) were exposed to antenatal steroids, perinatal surfactant, and resuscitated and supported by mechanical ventilation for 6–7d (Drager VN500, SIMV). Physiological targets were PaO2,60–90 mmHg, PaCO2,45–60 mmHg, O2 saturation 88–92%, pH 7.25–7.35. One group was treated with MES (60 x 106 cell equivalents; 10 mL; n=8; 4F 4M) at hours of life 6 and 78 (iv); the control group received vehicle (MEx diluent in saline; 10 mL; n=8; 4F 4M). We used morphometry and stereology to quantify structural indices of alveolar formation, and immunoblot to quantify apoptosis (cleaved caspase 3) and proliferation (PCNA). Summary of Results Radial alveolar count and secondary septal volume density were significantly greater (* p<0.05) in the MEx-treated group compared to the control group (figure 1A and B). Distal airspace wall thickness was significantly narrower in the MEx-treated group compared to the control group (figure 1C). Normalized cleaved caspase 3 protein abundance was not different between the MEx-treated and control groups (0.71±0.05 vs 0.69±0.04, respectively). Normalized PCNA protein abundance was significantly lower in the MEx-treated group versus the control group (0.43±0.05 vs 0.55±0.05, respectively). No differences were detected between males and females. Conclusions We conclude that MEx improved alveoform formation in chronically mechanically ventilated preterm lambs. We speculate that MEx may be an effective therapy to promote normal structural development of the lung in preterm infants who require mechanical ventilation and are at-risk of developing BPD.

#65
REGENERATIVE RESPONSE OF ALVEOLAR TYPE 2 CELLS TO A GENETICALLY-INDUCED MOUSE PHENOCOPY OF BRONCHOPULMONARY DYSPLASIA

GA Kohbodi*, Gao, LL Peinado, R Ramanathan, P Minoo. 1Los Angeles County University of Southern California Medical Center, Los Angeles, CA; 2University of Southern California Keck School of Medicine, Los Angeles, CA

Purpose of Study In adult lung, alveolar type II cells (AT2s) serve as facultative stem cells. They proliferate in response to injuries to regenerate and repair the alveoli. There is lack of information on whether AT2s in immature lungs undergoing alveologenesis, such as those of preterm neonates may act as resident stem cells and undergo proliferation in response to injuries that cause BPD. Genetic inactivation of both TGFβ receptors in secondary crest myofibroblasts (SCMF) arrested alveologenesis causing a BPD phenocopy. Alveolar arrest was accompanied by decreased number of SCMF and AT2s, thus suggesting cross-communication between the two cell types during alveologenesis. To determine the mechanism, we quantified AT2 cell numbers in control and mutant lungs at postnatal days 7 and 14 (PN7 and PN14) during alveologenesis.

Methods Used A total of 12 mouse lungs, (control, n=6, and mutant, n=6) were examined at PN7 and PN14 (n=3 for each control and mutant.) Immuno-histochemistry and immuno-fluorescence were performed on multiple samples of lung tissues. AT2s were identified as SPC positive cells. Proliferating AT2s (pAT2s) were identified as SPC; Ki67 double positive cell. To correct for hypoplasia in BPD samples, all results were normalized against total lung cells, identified as DAPI positive.

Summary of Results In PN7 lungs, the ratio of AT2s/total cells (SPC+/DAPI+) was higher in mutant vs control (AT2s: 10.77% vs 8.58, respectively) likely due to reduction in DAPI + cells that included reduced SCMF. The pAT2s remained unchanged (pAT2: mutant 0.47% vs control 0.54%). In contrast, in PN14 lungs, both total AT2s, and pAT2s decreased in the mutant lungs vs control (total AT2s: 8.62% vs 10.54%, respectively) and (pAT2: 0.50% vs 0.93%, respectively) indicating that proliferation of both SCMF and AT2s has decreased.

Conclusions In mutant lungs, TGFβ receptors inactivation decreases SCMF numbers, while AT2s are unaffected in early phases of BPD-like pathogenesis. As the phenotype and loss of SCMF become more established and widespread, inhibition of AT2 proliferation becomes measurable. Two conclusions are derived from these observations: 1) targeted SCMF have a regulatory impact on AT2 proliferation, which is a
regenerative response to injury. 2) Despite important differences between mouse phenocopy and human BPD, similar dynamics may occur in the lungs of preterm infants who develop BPD. Response of the endogenous stem cells (i.e. AT2s) in the lung to initial injuries may be governed by an orthologous mesenchymal cell type (SCMF-like) and their communications with AT2 stem cells.

Supported by NHLBI, NIH and the Hastings Foundation

MESENCHYMAL STROMAL CELL EXTRACELLULAR VESICLES IMPROVE RESPIRATORY SYSTEM PHYSIOLOGICAL OUTCOMES IN MECHANICALLY VENTILATED PRETERM LAMBS

1A Rebentisch*, 1E Dawson, 1E Major, 1H Foreman, 1D Headden, 1Z Vordos, 1MJ Dahl, 1D Null, 1A Missialis, 1S Kouroubalis, 1K Albertine. 1The University of Utah School of Medicine, Salt Lake City, UT; 2University of California Davis, Davis, CA; 3Harvard Medical School, Boston, MA

Purpose of Study Mesenchymal stromal cell extracellular vesicle (MEx) treatment has therapeutic efficacy in murine neonatal hyperoxia models of bronchopulmonary dysplasia (BPD). Whether MEx will be beneficial in chronically ventilated preterm neonates is unknown. We tested the hypothesis that MEx will improve respiratory system physiological outcomes in chronically mechanically ventilated preterm lambs.

Methods Used Preterm lambs (128d; term ~150d; ~28w human gestation) were exposed to antenatal steroids, perinatal surfactant, and resuscitated and supported by mechanical ventilation for 6–7d (Drager VN5000, SIMV). Physiological targets were PaO2 60–90 mmHg, PaCO2 45–60 mmHg, O2 saturation 88–92%, pH 7.25–7.35. One group was treated with MEx (60 x 10^6 cell equivalents; 10 mL; n=8; 4F 4M) at hours of life 6 and 78 (iv); the control group received vehicle (MEx diluent in sterile saline; 10 mL; n=8; 4F 4M). We report daily physiological outcomes for respiratory severity score (RSS), oxygenation index (OI), Arterial-alveolar (A-a) gradient, and oxygen saturation/FiO2 (S/F) ratio. Liver and kidney function tests were assessed.

Summary of Results MEx-treated preterm lambs were ~1d younger (* p<0.05) and weighed less (*) at delivery than control lambs (figure 1A and B). MEx-treated lambs tolerated enteral feeding and maintained weight (*) whereas control lambs were less tolerant of enteral feedings and lost weight over 7d (figure 1B). RSS, OI, and A-a gradient were lower for MEx-treated group (*) compared to the control group. S/F ratio was higher for the MEx-treated group (*) compared to the control group. Neither liver nor kidney toxicity was detected. Differences were detected between females and males.

Conclusions We conclude that MEx improved respiratory system physiological outcomes in chronically mechanically ventilated preterm lambs. We speculate that MEx may be an effective therapy for appropriate functional development of the lung in preterm infants who require mechanical ventilation.

OXIDIZED PHOSPHOLIPID NEUTRALIZING ANTIBODY AMELIORATES HYPEROXIA INDUCED LUNG INJURY

1W Tang*, 1X Sun, 1L Witt stem, 1C Glass, 1E Sajti. 1University of California San Diego, La Jolla, CA; 2The University of Texas Health Science Center at San Antonio, San Antonio, TX

Purpose of Study Oxidized phospholipids (OxPL) are formed during inflammatory processes, and they are known to induce cellular stress and apoptosis. The role OxPL play in lung inflammation is not known. OxPL are recognized by the IgM natural antibody (Ab) E06, which can bind to and block many of the pro-inflammatory properties of OxPL. To investigate the role of OxPL in hyperoxia-induced acute lung injury (HALI) and whether neutralizing OxPL using E06 would ameliorate or prevent hyperoxia induced lung injury.

Methods Used C57BL/6J (B6) sensitive and DBA/2J (DBA) resistant mice were exposed to hyperoxia for 48h to induce lung injury. We examined the content of OxPL by immunohistochemistry with E06 and examined inflammatory responses by measuring changes in immune cell composition in the lung by fluorescence-activated flow cytometry and by immunohistochemistry. We measured gene expression changes in whole lung by RNA-seq. Data were analyzed with FlowJo and HOMER. To examine the pathogenic role of OxPL, we also exposed E06-scFv transgenic mice to hyperoxia. These mice generate a high plasma level of functional E06-scFv (single-chain variable fragment of E06).

Summary of Results Using immunohistochemistry, we observed an accumulation of OxPL in the lungs of sensitive B6 mice after hyperoxia. OxPL were more abundant in lungs of B6 mice compared to resistant DBA mice. To further explore the
molecular determinants of interstrain susceptibility to oxygen, we performed transcriptomic analysis of the whole lung. Transcripts that most distinguished B6 from DBA mice were associated with apoptotic and cell death pathways. To test whether OxPL have a pathogenic role, we exposed E06-scFv mice-on B6 background-to same hyperoxic conditions. Unlike B6 mice, E06-scFv mice did not show activation of apoptosis and cell death related gene pathways.

**Conclusions** We observed significant increases in OxPL accumulation following acute hyperoxia exposure in the lungs of injury sensitive compared to resistant mice. OxPL accumulation in the lungs of B6 mice was associated with upregulation of apoptosis and cell death related genes. Blocking of OxPL by the secreted E06-scFV Ab resulted in a significant reduction of apoptosis protecting the lung from HALL. These data suggest that OxPL are not only a useful biomarker for hyperoxia induced lung injury but that an OxPL neutralizing antibody could be used to ameliorate or prevent HALL. Furthermore, the magnitude of interstrain variability in lung gene expression could form the basis for understanding human interindividual variability in susceptibility to oxygen induced injury.

#68

**ASYNCHRONOUS CHEST COMPRESSIONS WITH HIGH FREQUENCY VENTILATION IMPROVES GAS EXCHANGE DELIVERY IN ASPHYXIATED CARDIAC ARRESTED PRETERM LAMBS**

1E Giusto*, 2A Lesneski, 3H Loudi, 4M Hardie, 4L Zeinali, 4D Sankaran, 5S Lakshmiminnusinha, 2P Vakil. 1University of California Davis, Sacramento, CA; 4University of California Davis, Davis, CA

10.1136/jim-2022-WRMC.67

**Purpose of Study** Gas exchange is severely impaired during cardiopulmonary resuscitation (CPR) in the cardiac arrested lamb model despite ventilation with 100% O2. Optimizing gas exchange during neonatal CPR may improve cerebral oxygen delivery (cDO2), prevent rapid fluctuations in PaCO2, and stabilize cerebral blood flow. We hypothesize that asynchronous continuous chest compressions with high frequency percussive ventilation (HFPV) in preterm asphyxiated cardiac arrested lambs will result in improved gas exchange and cDO2 compared to 3:1 compression-to-ventilation (C:V) resuscitation.

**Methods Used** Time-dated preterm (~125d gestation; equivalent human ~25 weeks) fetal lambs were intubated, instrumented to measure cerebral blood flow and arterial blood pressure, and catheterized to collect venous and arterial blood. After instrumentation, lambs were asphyxiated by umbilical cord occlusion until asystole and delivered. Lambs were randomized to (1) 3:1 C:V resuscitation using a T-piece resuscitator following the neonatal resuscitation program (NRP) algorithm (control), or (2) asynchronous continuous chest compressions (120 compressions/min) with HFPV using a TXP-5 ventilator (intervention). First dose of epinephrine (0.03 mg/kg) was given at three minutes and repeated q3min until return of spontaneous circulation (ROSC). Lambs in the control group that achieved ROSC were managed on conventional ventilation and lambs in the intervention group were maintained on HFPV. Ventilation parameters and O2 were adjusted to maintain SpO2 at 90–95% and PaCO2 between 45–60 mmHg.

**Summary of Results** Eight lambs were studied and all achieved ROSC. Baseline characteristics, time to ROSC, and epinephrine doses were similar between groups (figure 1). Mean (SD) PaCO2 was 158 (24) mmHg and the mean (SD) PaO2 was 47 (42) mmHg 15 minutes post-ROSC despite maximum ventilation support and 100% O2 in the control group compared to a mean (SD) PaCO2 of 30 (11) mmHg and a PaO2 of 60 (24) mmHg in the intervention group (table 1).

**Conclusions** Resuscitation using asynchronous continuous chest compressions during HFPV is feasible with similar success of ROSC and improved gas exchange in an asphyxiated cardiac arrested neonatal lamb model. Further studies are required to validate our results and to assess lung injury by immunohistochemistry and biomarkers.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Control 3:1 (N = 4)</th>
<th>Intervention HFPV (N = 4)</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Weight</td>
<td>2.89 (0.21)</td>
<td>2.69 (0.46)</td>
<td>0.47</td>
</tr>
<tr>
<td>Time to Asystole (min)</td>
<td>28 (8.5)</td>
<td>18 (1.3)</td>
<td>0.09</td>
</tr>
<tr>
<td>Asphyxia pH</td>
<td>6.80 (0.08)</td>
<td>6.83 (0.04)</td>
<td>0.59</td>
</tr>
<tr>
<td>Asphyxia PaO2</td>
<td>Undetectable</td>
<td>Undetectable</td>
<td>N/A</td>
</tr>
<tr>
<td>Asphyxia PaCO2</td>
<td>145 (14)</td>
<td>137 (12)</td>
<td>0.40</td>
</tr>
<tr>
<td>Asphyxia Lactate</td>
<td>9 (1.3)</td>
<td>11 (3.9)</td>
<td>0.40</td>
</tr>
<tr>
<td>ROSC success</td>
<td>100%</td>
<td>100%</td>
<td>N/A</td>
</tr>
<tr>
<td>Time to ROSC (min)</td>
<td>4.4 (1.8)</td>
<td>4.1 (0.6)</td>
<td>0.61</td>
</tr>
<tr>
<td># of Epi Doses</td>
<td>1</td>
<td>1.25 (0.5)</td>
<td>0.39</td>
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**Abstract #68 Table 1** Comparison of blood gases at fixed timepoints

<table>
<thead>
<tr>
<th>Time Point</th>
<th>PaCO2 (mm Hg)</th>
<th>PaO2 (mm Hg)</th>
<th>CaCO2 (mL O2/ml)</th>
<th>Brain DO2 (mL O2/kg/ml)*</th>
<th>FiO2</th>
</tr>
</thead>
<tbody>
<tr>
<td>During Chest</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Compressions</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>At ROSC</td>
<td>153 (11)</td>
<td>13 (15)</td>
<td>2.38 (3.2)</td>
<td>0.05 (0.06)</td>
<td>1.0</td>
</tr>
<tr>
<td>15 minutes post-ROSC</td>
<td>158 (24)</td>
<td>47 (42)</td>
<td>4.58 (4.8)</td>
<td>0.62 (0.59)</td>
<td>1.0</td>
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<tr>
<td>Intervention Group (continuous asynchronous chest compressions with HFPV)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>During Chest</td>
<td>110 (5)</td>
<td>22 (3.1)</td>
<td>5.04 (1.8)</td>
<td>0.12 (0.11)</td>
<td>1.0</td>
</tr>
<tr>
<td>Compressions</td>
<td>At ROSC</td>
<td>85 (20)</td>
<td>57 (13)</td>
<td>3.8 (0.14)</td>
<td>1.0</td>
</tr>
<tr>
<td>15 minutes post-ROSC</td>
<td>50 (11)</td>
<td>60 (24)</td>
<td>13.7 (1.5)</td>
<td>2.42 (0.71)</td>
<td>0.48</td>
</tr>
</tbody>
</table>

**Abstract #68 Figure 1** Comparison of baseline characteristics between groups. There was no significant difference of characteristics between groups (p > 0.05)
Surgery I
Concurrent session
12:45 PM
Thursday, January 20, 2022

HEMOGLOBIN A1C AS A PROGNOSTIC INDICATOR OF POSTOPERATIVE INFECTION FOLLOWING IMMEDIATE BREAST CANCER RECONSTRUCTION
CA Czajkowsky*, S Gupta. Loma Linda University School of Medicine, Loma Linda, CA
10.1136/jim-2022-WRMC.69

Purpose of Study Hemoglobin A1c (HgbA1c) is a marker of an individual’s glycemic exposure over a preceding 2–3 month period. Minimal evidence currently exists to support increased infection risk following mastectomy. We aimed to evaluate the association of HgbA1c with the incidence of surgical site infection (SSI) in patients undergoing mastectomy and immediate breast reconstruction.

Methods Used An institutional database was queried for patients with CPT code for reconstruction AND diagnosis code for breast malignancy for patients from January 1, 2014 to June 20, 2021. We defined SSI incidence by diagnosis or procedure for SSI within 90 days following mastectomy. A one sample t-test was performed to determine if there is a significance difference in the average HgbA1c of the standard patient population and the sample SSI group. A chi-square test was used to analyze data for correlations between SSI rates in diabetics and non-diabetics. The patients were analyzed using a standard chi-square based on a 2x2 contingency table.

Summary of Results A total of 1386 patients were included in the query; 268 of which having received a pre-op HgbA1c and with 136 having received a pre-op HgbA1c and a DM diagnosis. Only 22 patients fit our defined SSI sample group criteria. The average population HgbA1c was 6.74 (N=268), the average HgbA1c of diabetics was 7.33 (N=136), the average HgbA1c of non-diabetics was 6.13 (N=132). The average sample HgbA1c was 6.94 (N=22), the average HgbA1c of diabetics was 7.34 (N=17), the average HgbA1c of non-diabetics was 5.58 (N=5). The one sample t-test of the average HgbA1c value in patients with DM vs non-diabetics in the...
Purpose of Study

The association of HgbA1c with the incidence of surgical site infection (SSI) following undergoing mastectomy is evident. We found the presence of a DM diagnosis as a better prognostic tool for SSI than HgbA1c level alone. Chi-square analysis determined the relative risk of SSI following mastectomy in diabetic patients at 2.28x that of a nondiabetic. Considering the SSI sample group, a HgbA1c threshold of 7.34 is what we propose as presenting great additional post-surgical complication risk.

### Abstract #71

**THE IMPACT OF THE COVID-19 PANDEMIC ON SPINE SURGERY PRACTICE AND OUTCOMES IN AN URBAN HEALTHCARE SYSTEM**

J Parekh, B Attaripour, SY Xiang*, M Siow, B Mitchell, B Shahidi. University of California San Diego, University of California San Diego, La Jolla, CA, San Diego, CA

10.1136/jim-2022-WRMC.70

**Purpose of Study**

To elucidate changes due to COVID-19 on patient demographics, surgical care, logistics, and patient outcomes in spine patients.

**Methods Used**

This is a retrospective study of patients who had spine surgery at UCSD from 3/1/19 to 5/31/19 (pre-COVID-19) and 3/1/20 to 5/31/20 (first COVID-19 surge). 331 subjects met the study criteria. Demographic and surgical data were collected from medical records. Pain levels at preoperative, discharge, short- (3–6 month) and long-term (9–15 month) timepoints were extracted.

**Summary of Results**

There were no significant differences in patient demographics including age, BMI, gender, race, ethnicity, ASA rating, smoking status, or diabetes status between groups (p>0.14). The diagnostic indications for surgery (spondylolysis, tumor/infection, spondylolisthesis, fracture) were not different between groups (p>0.13). There were no differences in operating room duration and skin-to-skin time (p>0.64), however length of stay was 4.7 days shorter during the COVID-19 pandemic (p=0.03) and more cases were classified as ‘urgent’ (p=0.04). Preoperative pain scores did not differ significantly between groups (p=0.58), however pain levels at discharge were significantly higher in patients operated upon during COVID (p=0.04) and trended towards remaining higher in the short- (p=0.06) but not long-term (p=0.21) after surgery (table 1).

**Conclusions**

The pandemic resulted in a greater proportion of ‘urgent’ spine surgery cases and shorter hospital length of stay. Pain levels upon discharge and at short-term timepoints were higher following surgery, however these differences did not persist in the long term.

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### Abstract #72

**MEASURING THE CLINICAL VALUE OF SECONDARY HAND SURGERY IN PATIENTS WITH COMPLEX HAND INJURIES**

1BC Goodlin*, 1CA Czaykowsky, 2S Gupta. 1Loma Linda University School of Medicine, Loma Linda, CA; 2Loma Linda University, Loma Linda, CA

10.1136/jim-2022-WRMC.71

**Purpose of Study**

The objective of this study is to assess the value of performing secondary hand surgery in a population of complex hand procedures at a quaternary referral site academic medical center.

**Methods Used**

This was done by obtaining electronic medical records (EMR) of 166 patients over a five year period using specific keywords relating to the types of surgeries generally performed as secondary procedures such as tenolysis, contracture release, and capsulotomies. Of those patients, 50 were found to fit within the parameters of the study. For these 50 remaining patients, hand therapy data was obtained and the percentage of motion that each patient had before the secondary surgery and following the secondary surgery was calculated and this data was used to calculate the overall change in motion. A paired sample t-test was performed to determine if there is a significant difference in the average motion data in the measurements taken before and after secondary hand surgeries. One sample t-tests were performed to determine if there is a significance difference in the average change in the range of motion between common comorbidities (asthma, diabetes mellitus, hypertension, obesity, age over 45 years) or gender between the patient population and each subgroup.

**Summary of Results**

During a median follow-up period of 8.5 months and an average follow-up period of 18.87 months, a total of 75 complications in 50 patients were recorded. The average percentage of full motion before surgery was 49.43% and the average percentage of full motion after surgery was 66% to give an overall change in motion to be +16.58%. Hand motion measurements following secondary hand surgery (M = 0.491, SD = 0.188) compared to the hand motion measurements preceding secondary hand surgery (M = 0.660, SD = 0.259) demonstrated a significantly better change in motion percent change, t(50) = 24.50, p = 4.177E-5.

**Conclusions**

Although we speculate that we would have seen an even greater impact from secondary hand surgeries if there had been greater adherence to post-surgical hand therapy, our p value indicates results that are statistically significant. Therefore, we conclude that secondary hand surgery performed on patients with complex hand injuries has a significant measurable impact and we believe that a similar study in a larger population would yield similar results.
Purpose of Study Malnutrition is associated with increased morbidity and mortality in patients with head and neck cancer (HNC) undergoing surgery. Despite the profound impact malnutrition has on this patient population, objective screening tools are still lacking in a clinical setting. Without a clear approach to identify malnutrition, there is currently a barrier to capturing patients with inadequate nutrition, delaying interventions that could otherwise be implemented to optimize their nutritional status. Therefore, recognizing the need for a tool, the aim of this study is to assess the ability to use the geriatric nutrition risk index (GNRI) to screen for malnutrition among HNC patients and determine if there is an association between GNRI scores and postoperative complications.

Methods Used A retrospective review of medical records was conducted for patients undergoing surgical resection at a tertiary academic hospital from June 2012 to June 2021. Patients were included if surgical excision was the primary treatment modality and if a serum albumin was obtained 6 months prior to surgery. A total of 44 HNC patients were included in the study and analysis. Preoperative body weight and serum albumin were abstracted from medical records to calculate the GNRI.

Summary of Results Of the 44 patients included in the study, there were 30 men (68%) and 14 women (32%), with a total mean age of 62 ±12 years. Malnutrition was defined by a GNRI score of <97.5 and was present in 27% of patients (n=12). Malnourished patients had significantly higher rates of postoperative complications and required discharge to a skilled nursing facility (SNF) more often compared to the control group. Conclusions A low GNRI score appears to be a predictor of increased complications after head and neck surgery. The GNRI is a simple tool that utilizes serum albumin and body weight to objectively assess nutritional status. Results from this study suggest that, in the future, the GNRI may be a clinically useful approach to screen for malnutrition and identify patients who are at high risk for complications during the postoperative course.

Abstract #74 DEMOGRAPHICS, FRACTURE CHARACTERISTICS, AND TREATMENT STRATEGIES FOR PERIPROSTHETIC DISTAL FEMUR FRACTURES COMPARED TO NATIVE DISTAL FEMUR FRACTURES

Purpose of Study The incidence of periprosthetic distal femur fractures is increasing due to the increasing number of knee arthroplasties being performed in the aging population. The purpose of this study was to analyze the demographics, fracture characteristics, and treatment strategies associated with periprosthetic distal femur fractures (PDFF) compared to native distal femur fractures (NDFF) in order to identify important clinical differences between these groups that might help guide management.

Methods Used A retrospective study was conducted of 209 patients >18 years old who underwent surgical treatment for either a native distal femur fracture (NDFF) or a periprosthetic distal femur fracture (PDFF) about a total knee arthroplasty (TKA) from January, 2006 to December, 2020. Fracture classification of CT images by the Association for Osteosynthesis/Orthopedic Trauma Association (AO/OTA) was reported. Demographics, fracture characteristics, fixation constructs, and surgical outcomes were compared between subjects with PDFF vs. NDFF.

Summary of Results Out of 70 patients with PDFF, 81.1% were female and 18.6% were male, with an average age of 80 years old (range= 49–102 yrs). PDFFs were most often isolated (80%) or comminuted (85%) injuries with AO classification 33A.3 (71.4%). Out of 139 patients with NDFF, 53.2% were female and 46.8% were male with an average age of 57 years old (range =18–96 yrs). NDFFs were commonly comminuted (92.1%) injuries with AO classification 33C.2 (28.1%) or 33A.3 (25.2%). NDFFs were extra-articular (54.0%) or intra-articular (46.0%). Nearly half of subjects with NDFF (48.2%) experienced concomitant fracture of the ipsilateral knee (14.4%) or tibial plateau (15.1%). Intramedullary nailing was the most common fixation construct for both fracture groups (42.6% PDFF;36.7% NDFF). The second...
most common fixation construct for PDFF was combined nail/plate (17.3%) and lateral locking plate (20.9%) for NDFF. Patients with PDFF experienced shorter length-of-stays (6.36 days vs. 11.4 days) but had higher complication rates compared to NDFF (5.7% vs 4.4%). Incidence of low bone density (osteopenia or osteoporosis) was higher in those with PDFF compared to NDFF (55.7% vs. 19.4%).

Conclusions PDFFs frequently occur as isolated comminuted injuries with greater complication rates compared to NDFF. Though intramedullary nailing remains the most common fixation construct for both NDFF and PDFF, stabilization via combined plate/nail is increasingly being used for PDFFs. Elderly women with TKA and poor bone quality are a high risk group for PFF. Further research should entail how physicians can improve their surgical and clinical approach for this type of fracture in the affected population.

Summary of Results A statistically significant reduction in intraoperative times (p-value 0.03) was observed in patients who were not taking any form of antithrombotic medication (cohort 5), as compared to patients in cohort 4 who were actively taking anticoagulant medications. Other classes of antithrombotic medication (cohorts 1–3) were associated with higher average intraoperative times relative to cohort 5, however the difference was not statistically significant. There was no statistically significant difference in closure size across the cohorts.

Conclusions The use of oral anticoagulants in patients undergoing Mohs and cosmetic flap surgeries results in significantly longer intraoperative times. Further investigation of this relationship and consideration of this finding may influence management of dermatologic and cosmetic procedures.

Abstract #75 Table 1

<table>
<thead>
<tr>
<th>Cohort</th>
<th>Medications</th>
<th>Number of Patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cohort 1</td>
<td>Supplements (Fish oil, krill oil, garlic, turmeric)</td>
<td>12</td>
</tr>
<tr>
<td>Cohort 2</td>
<td>COX inhibitors</td>
<td>104</td>
</tr>
<tr>
<td>Cohort 3</td>
<td>ADP inhibitors, phosphodiesterase inhibitors, glycoprotein IIb/IIa inhibitors</td>
<td>11</td>
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<tr>
<td>Cohort 4</td>
<td>Vitamin K antagonists, direct thrombin inhibitors, direct Xa inhibitors, indirect thrombin inhibitors</td>
<td>25</td>
</tr>
<tr>
<td>Cohort 5</td>
<td>No antithrombotic medications</td>
<td>91</td>
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</tbody>
</table>
patients on opioid-inclusive analgesia during RALP while decreasing post-surgical and home opioid use.

**Conclusions** By markedly decreasing post-surgical opioids prescriptions, we can reduce the risk opioid addiction and the associated harm to the patient. This study is a proof of principle that buprenorphine use for perioperative analgesia during RALP is an alternative to traditional opioid-inclusive analgesic pathways. We believe such a strategy will decrease the incidence of opioid use disorder and have benefits including less associated healthcare spending, improved patient health and reduced social harm.

**Purpose of Study** This study compares the outcomes, features, and costs of laminectomy and fusion (LEF) versus laminoplasty (LP) as surgical treatments for patients with cervical spondylotic myelopathy (CSM).

**Methods** Elective LEF and LP procedures performed at a single institution between 2014 and 2020 were identified. Included patients had no prior cervical spine surgery. All patients received pre- and postoperative outpatient evaluations in the outpatient clinic. Only procedures involving three or more spinal levels were included. Clinical data was collected from electronic medical records. SPSS 27 was used for statistical analysis. Hospital costs were obtained from hospital billing for a subgroup of patients for whom this information was available.

**Summary of Results** 135 patients were included: 76 underwent LP and 59 underwent LEF. Mean follow-up time was 14 months. Compared to LEF, LP procedures involved fewer levels (4.2 vs 4.8 levels, p < .001) and trended shorter operative time per level (47 vs 62 minutes, p < .001). Intraoperative blood loss and fluid replacement were similar between groups (p = .79 and p = .08). Patients in the LP group were discharged an average of 1.1 days earlier (p = .001). LP was not associated with higher rates of C5 palsy (p = .28).

Patients who underwent LEF were five times more likely to develop wound infection or dehiscence (risk ratio = 5.2, 95% CI: 1.1 to 23.4). Postoperative ground-level falls requiring an emergency department (ED) visit occurred more frequently in the LEF group (11.9% vs 2.6%, p = .04). The frequency of ED visits for postoperative neck pain did not differ between groups (p = .42). Likewise, rates of new-onset neck pain were similar (p = .45). Both groups reported improved VAS neck pain over the course of follow-up (p = .001). Surgery type, involvement of the C7 level, and the number of levels involved were not predictive of differences in postoperative neck pain (p = .66, p = .31, and p = .87). Opioid analgesic needs in the year before and the year after surgery were similar between groups (p = .41 and p = .33). The LP cohort had greater preoperative cervical lordosis (C2-C7 Cobb angle: 11.69 vs 6.59, p = .01) and lost more lordosis postoperatively (-7.9 vs -1.8, p = .004). LEF cases at this hospital incurred 18% and 34% greater fixed and variable costs (p = .03 and p < .001).

**Conclusions** When used to treat patients with multilevel CSM, LP does not seem to be associated with new or worsening axial neck pain compared to LEF. Neck pain may be expected to improve similarly with either surgery. When cervical deformity is not prohibitive, LP could be offered as a less morbid and more cost-efficient alternative to LEF. Modern patient-reported outcomes and randomized controlled trials are still needed to optimize the utility of both procedures.

**Purpose of Study** Diabetes Mellitus (DM) has a significant burden in the United States and results in worsening health outcomes. Patients are at risk of peripheral neuropathy, which increases the risk of lower extremity burns, delay in burn presentation, and more complications that translate to more amputations. However, there are limited reports regarding the incidence and outcomes of DM foot burns. We aim to better understand DM health outcomes, specifically lower limb amputations, in DM foot burns at national Level 1 and 2 trauma centers.

**Methods** Implementing a retrospective cohort study design, we reviewed de-identified data on 116,796 adult admissions from 2007–2015 from the National Trauma Database (NTDB) for patient age, DM, foot burn status, sex, race/ethnicity, region, burn size, and comorbidities. An exploratory logistic regression of factors associated with lower limb amputations was performed.

**Summary of Results** Of the 7,963 (7%) foot burn patients, 1,338 (17%) had DM (median age 56 years [17]) and 378 (28%) were male. Common comorbidities included alcohol use, smoking, and chronic kidney disease (table 1). Using an exploratory logistic regression analysis, when all other variables were kept the same, factors (OR, CI) linked with total lower limb amputations in DM foot burns national Level 1 and 2 trauma centers.

**Conclusions** When used to treat patients with multilevel CSM, LP does not seem to be associated with new or worsening axial neck pain compared to LEF. Neck pain may be expected to improve similarly with either surgery. When cervical deformity is not prohibitive, LP could be offered as a less morbid and more cost-efficient alternative to LEF. Modern patient-reported outcomes and randomized controlled trials are still needed to optimize the utility of both procedures.
use (2.78, [2.13, 3.61]), smoking (0.78, [0.62, 1.00]), chronic kidney disease (2.90, [1.72, 4.88]), burn size >20% (4.12, [2.96, 5.73]), African-American/Black race (1.61, [1.29, 2.01]), male sex (1.61, [1.28, 2.02]), and age >40 years. There was a higher rate of lower limb amputations in patients with DM foot burns.

Conclusions There is a higher rate of total amputations in DM foot burn patients, which indicates the need for increased patient education and treatment protocols that address the incidence of complications in this group. Future steps include confirmatory research to assess the risk of lower limb amputations in DM foot burn patients.

Cardiovascular II
Concurrent session
3:15 PM
Thursday, January 20, 2022

#79 OUTCOME OF HOSPITALIZED HEART TRANSPLANT PATIENTS WITH COVID INFECTION AT A LARGE WEST COAST CENTER

A Mohanty, K Kim, N Patel, T Singer-Englar, M Hamilton, J Kobashigawa. Santa Clara University, Santa Clara, CA; Cedars-Sinai Smidt Heart Institute, Los Angeles, CA

Purpose of Study The COVID-19 pandemic infected large portions of the US community and infected many heart transplant (HTx) patients, but in distinct geographical patterns. HTx programs have reported mortality in the range of 23–29% and in non-transplant patients in the range of 15–17%. The impact of hospitalized HTx patients with COVID infection in a large West Coast heart transplant program has not been reported. We now report our outcomes for hospitalized patients with COVID.

Methods Used Between March 2020 and March 2021, we assessed 22 HTx patients who were admitted to the Cedars-Sinai Medical Center (CSMC) for COVID infections. COVID is known to affect many systems within the body, and we report the effects on lungs, heart, and kidney. Morbidity and mortality, including risk of death, were included within 90 days post-infection.

Summary of Results Of the 22 HTx patients hospitalized at the CSMC, 7 patients died (31.8%). All patients had COVID pneumonia requiring supplemental oxygen and 5 patients required ventilatory support. The mean peak FiO2 of the patients was 79.7%. 16 of these patients also were noted to have an increase in serum creatinine, with 6 patients requiring kidney dialysis. Cardiac function was maintained in all patients with COVID-19 and no myocarditis or cardiac dysfunction was observed. 9 patients received remdesivir and 19 patients received corticosteroids. 4 patients received tocilizumab anti-inflammatory therapy.

Conclusions COVID-19 resulted in significant morbidity and mortality in hospitalized HTx patients. The immunosuppressed state appears to be a risk factor for poor outcome and is higher compared to non-transplant hospitalized patients.

#80 IS SACUBITRIL/VALSARTAN A RISK FACTOR FOR VASOPLEGIA/PRIMARY GRAFT DYSFUNCTION AFTER HEART TRANSPLANTATION?

M Oda, T Singer-Englar, N Patel, S Kim, M Hamilton, J Kobashigawa. University of California Los Angeles, Los Angeles, CA; Cedars-Sinai Smidt Heart Institute, Los Angeles, CA

Purpose of Study Primary graft dysfunction (PGD) is seen in approximately 7–29% of heart transplant (HTx) patients. Many of these patients with PGD also develop significant vasoplegia which requires high doses of intravenous vasoconstrictors. Outcomes of these patients with severe PGD is compromised within 30 days after HTx. Risk factors for the development of severe PGD have included angiotensin-convertase enzyme inhibitors (ACEi). There may be a connection between ACEi and the kallikrein-kinin system whereby bradykinin is increased, thus resulting in more vasoplegia and PGD. It is not known whether the new drug sacubitril/valsartan (S/V) is also a risk factor for the development of vasoplegia/severe PGD as bradykinin is also increased with sacubitril. Therefore, we reviewed our large HTx program to see if there is a correlation of S/V as a risk factor for this complication.
Methods Used Between 2015 and 2020, we assessed 65 HTx patients who were on S/V at the time of transplantation. Vasoplegia was defined as requiring more than 2 vasoconstricting drugs with BP systolic <90 mmHg, and PGD was defined as per the ISHLT classification scheme (within 24 hours post-transplant). These patients on S/V were compared to patients on ACEI/ARB (1:1 control group for age, sex, transplant year). Outcomes included death, cardiac dysfunction, and non-fatal major adverse cardiac events (NF-MACE: MI, new CHF, PCI, ICD/pacemaker, or stroke) in the first year after HTx.

Summary of Results Compared to ACEI/ARB, S/V had similar risk for the development of vasoplegia or severe PGD. Furthermore, 1-year survival, and 1-year freedom from cardiac dysfunction and NF-MACE were not significantly different between groups.

Conclusions Patients undergoing HTx on S/V do not appear to be at risk for vasoplegia or severe PGD.

Abstract #81 Table 1 Comparison of Entresto vs. ACEI/ARB

<table>
<thead>
<tr>
<th>Endpoint</th>
<th>Entresto (n=65)</th>
<th>ACEI/ARB control (n=65)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-year survival</td>
<td>89.2%</td>
<td>90.8%</td>
<td>0.716</td>
</tr>
<tr>
<td>1-year freedom from cardiac dysfunction</td>
<td>78.5%</td>
<td>72.3%</td>
<td>0.432</td>
</tr>
<tr>
<td>1-year freedom from NF-MACE</td>
<td>95.3%</td>
<td>87.7%</td>
<td>0.139</td>
</tr>
<tr>
<td>Development of severe PGD</td>
<td>4.6%</td>
<td>6.2%</td>
<td>1.000</td>
</tr>
<tr>
<td>Development of vasoplegia</td>
<td>29.2%</td>
<td>33.9%</td>
<td>0.706</td>
</tr>
</tbody>
</table>

Abstract #82 ACUTE ABDOMINAL COMPLICATIONS IMMEDIATELY FOLLOWING HEART TRANSPLANTATION

Purpose of Study The calcineurin inhibitors (CNIs), including tacrolimus and cyclosporine, have revolutionized heart transplantation (HTx) in terms of maintaining low rejection rates. However, CNIs have significant side effects such as nephropathy, hypertension, malignancy, and hypomagnesemia. It is this hypomagnesemia that has not been addressed as to whether this has an impact on outcome after HTx. Hypomagnesemia has been involved in muscle cramping and cardiac arrhythmias. Therefore, we reviewed our HTx patients and assessed magnesium (Mg) levels to assess outcome in the first 6 months after HTx.

Methods Used Between 2010 and 2020, we assessed 956 HTx patients and recorded their Mg levels in the first 6 months after HTx. Patients with low Mg levels less than or equal to 1.8 mg/dL were assessed for complications including muscle cramping, cardiac arrhythmias, rehospitalization, rejection episodes, and death. Patients with low Mg levels were grouped into mildly low Mg levels (1.7–1.8 mg/dL) and moderately low Mg levels (1.4–1.7 mg/dL). Patients were compared to control patients who had normal Mg levels (>1.8 mg/dL) during this period of time.

Summary of Results Patients with mildly or moderately low Mg levels compared to patients with normal Mg levels had no difference in muscle cramping, rejection episodes, cardiac arrhythmias, and use of antihypertensive medications. Kidney function was abnormal in those patients with normal magnesium levels.

Conclusions Mild-moderate hypomagnesemia did not have significant adverse effects in heart transplant patients in terms of muscle cramping, cardiac arrhythmias, cardiac rejection, or cardiac function.

Abstract #81 Table 1

<table>
<thead>
<tr>
<th>Moderate</th>
<th>Mild Low</th>
<th>Normal</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mg</td>
<td>Mg</td>
<td>Mg</td>
<td></td>
</tr>
<tr>
<td>Low</td>
<td>Normal</td>
<td>Normal</td>
<td></td>
</tr>
<tr>
<td>1.4–1.7 mg/dL</td>
<td>1.7–1.8 mg/dL</td>
<td>&gt;1.8 mg/dL</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>(n=158)</td>
<td>(n=625)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6-Month Average Magnesium</td>
<td>1.66 ± 0.06</td>
<td>1.79 ± 0.02</td>
<td>2.05 ± 0.15</td>
</tr>
<tr>
<td>Incidence of Muscle Cramping within 6-Months Post-Transplant</td>
<td>12.1%</td>
<td>8.2%</td>
<td>11.5%</td>
</tr>
<tr>
<td>Incidence of Cardiac Arrhythmias within 6-Months Post-Transplant</td>
<td>5.2%</td>
<td>10.8%</td>
<td>21.9%</td>
</tr>
<tr>
<td>Rehospitalization within 6-Months Post-Transplant</td>
<td>17.3%</td>
<td>22.8%</td>
<td>25.1%</td>
</tr>
<tr>
<td>6-Month Freedom from Any Treated Rejection</td>
<td>89.0%</td>
<td>88.6%</td>
<td>87.4%</td>
</tr>
<tr>
<td>6-Month Survival</td>
<td>100.0%</td>
<td>99.4%</td>
<td>93.6%</td>
</tr>
<tr>
<td>6-Month Average Creatinine</td>
<td>1.25 ± 0.46</td>
<td>1.28 ± 0.48</td>
<td>1.74 ± 0.78</td>
</tr>
</tbody>
</table>
surgery. Types of surgical interventions included hemicolec-
tomy, cholecystectomy, and exploratory laparoscopy. Compared
to the control group, the acute abdomen group had signific-
antly worse 30-day survival and 1-year survival. In the study
group, infectious complications occurred in an additional
36.4% of these patients who required rehospitalization with
administration of intravenous antibiotics. Rejection episodes
following these events was not different from the control
population.

Conclusions Acute abdomen immediately post-heart transplant
resulting in urgent abdominal surgery requiring hemicolec-
tomy and/or cholecystectomy has significant morbidity/mortality.
For patients awaiting heart transplant with gallstones, prophylactic
laparoscopic cholecystectomy might be considered.

#83

CLINICAL MANIFESTATION AND PROGNOSIS OF
DIFFERENT CARDIOMYOPATHY TYPES ON THE BASIS OF
GENETIC BACKGROUND

P-Aldino, P-Medo, L-Mestroni, M-Melto, T-Maylor, A-Universiti degli Studi di Trieste, Trieste, Italy, U-University of Colorado Denver School of Medicine, Aurora, CO

10.1136/jim-2022-WRMC.82

Purpose of Study Cardiomyopathies (CMP) are a heterogeneous
group of heart disease characterized by structural and electrical
abnormalities lacking secondary causative etiology and fre-
quently related to mutations in CMP genes. Recent studies in
this field have showed important phenotype overlaps between
Dilated Cardiomyopathy (DCM) and Arrhythmogenic Cardi-
omyopathy (ACM), making the diagnosis a challenging task.
The aim of this study is to assess whether a classification of
CMP patients (not hypertrophic) based on genetic character-
ization outperforms in diagnostic and prognostic accuracy the
classical, phenotype-driven, diagnostic approach.

Methods Used We analyzed a population of patients affected
by genetically determined DCM and ACM, including carriers of
‘pathogenic’ or ‘likely pathogenic’ (P/LP) variants, registered
into the Heart Disease Centers of Trieste and Denver hospi-
tals. We described the phenotype distribution in our popula-
tion with a clinical and echocardiographic evaluation based on
the different disease-related mutated genes. Then, we
examined the prognostic impact of the single gene/genetic
cluster in determining these outcomes: 1) all-cause mortality
and heart transplant; 2) heart failure-related death, heart
transplant or destination left ventricular assist device implanta-
tion (DHF/HTx/VAD); and 3) sudden cardiac death, sustained
ventricular tachycardia/ventricular fibrillation or appropriate
defibrillator shock (SCD/VT/VF/shock).

Summary of Results 281 patients carrying P/LP variants (82%
DCM) were included in the study. Titin (TTN) and sarcomeric
genes (SARC) variants were the most prevalent (TTN: 95
patients, 34% of total population; SARC: 63 patients, 22% of
total population) and almost completely related to DCM phe-
notype (TTN: 100% DCM, SARC: 95% DCM); lamin (LMNA) patients (29 patients, 10% of total population, 96%
DCM). A more heterogeneous phenotypic distribution between
DCM and ACM were noted for desmoplakin (DSP), plakoglobin (PKP2) and filamin (FLNC) variants. Patients with uncate-
gorized DCM phenotype and carriers of DSP, PKP2, FLNC
and LMNA variants (arrhythmic genes) experienced more fre-
quent SCD/VT/VF/shock events (p value=0.002 and p=0.023), compared to patients with DCM phenotype, during
follow-up (median=132 months). The analysis shows only P/
LP variants of arrhythmic genes, early age of onset and male
gender, were associated with an increased risk of SCD/VT/VF/
shock events during follow-up. Additionally, there were no dif-
ferences in terms of HF events was significantly related to
genotype.

Conclusions In a large DCM and ACM population with a
positive genetic test for P/LP variants, the classification based
on specific genotypes is a useful tool in arrhythmic prognosti-
cation. These findings support the need of extensive genetic
testing to support CMP diagnosis and prognosis.

#84

SEX DIFFERENCES IN DESENSITIZATION FOR PATIENTS
AWAITING HEART TRANSPLANTATION: IS THERE A
DIFFERENCE?

Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA, Cedars-Sinai Smidt Heart Institute, Los Angeles, CA

10.1136/jim-2022-WRMC.83

Purpose of Study For patients awaiting heart transplantation
(HTx) who have high levels of circulating antibodies (greater
than 70%), desensitization therapy may be indicated. This will
allow expansion of the donor pool for a compatible donor.
As women appear to be more highly sensitized (due to multi-
ple pregnancies), it is not clear as to whether women can ben-
efit from desensitization therapy. We sought to answer this
question with review of our large, single center database.

Methods Used Between 2008 and 2020, we assessed 49
patients awaiting HTx who underwent desensitization therapy.
These patients were divided into groups by sex for their
response to desensitization therapy. Our desensitization proto-
cols consist of regimens including intravenous immune globu-
lin, anti-CD20 monoclonal antibody, plasmapheresis, and/or
proteosome inhibitors. A response to desensitization therapy
was assessed by the decline of the dominant circulating anti-
body determined by mean fluorescence intensity (MFI). Post-
HTx data was assessed for 1-year survival and freedom from
rejections (acute cellular rejection [ACR], antibody-mediated
rejection [AMR]). Rejection episodes were compared to a
control group of non-sensitized patients transplanted during the same period (n=771).

Summary of Results Desensitization therapy in women appeared to have similar response to various desensitization protocols. There were no significant differences in waitlist mortality, time on the waitlist, 1-year post-transplant survival, or 1-year freedom from ACR or AMR between the two groups. Compared to non-sensitized patients, freedom from AMR was significantly lower in both sensitized men and women (72.7% men vs. 78.9% women vs. 96.5% control group, p<0.001).

Conclusions Sensitized women awaiting HTx compared to men appear to have similar response to various desensitization regimens. Post-HTx, there was more AMR in both groups, suggesting memory B-cells may be responsible.

Abstract #84 Table 1

<table>
<thead>
<tr>
<th></th>
<th>Male</th>
<th>Female</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>(n=11)</td>
<td>(n=38)</td>
<td></td>
</tr>
<tr>
<td>Immunodominant Antibody MFI Pre-Treatment</td>
<td>12613 ± 3642</td>
<td>12779 ± 3498</td>
<td>0.891</td>
</tr>
<tr>
<td>Immunodominant Antibody MFI Post-Treatment</td>
<td>12045 ± 3514</td>
<td>12055 ± 3925</td>
<td>0.994</td>
</tr>
<tr>
<td>Change in Immunodominant Antibody MFI</td>
<td>-568 ± 1687</td>
<td>-723 ± 1782</td>
<td>0.798</td>
</tr>
<tr>
<td>Average Time on Waitlist (years)</td>
<td>0.61 ± 0.62</td>
<td>0.65 ± 0.82</td>
<td>0.096</td>
</tr>
<tr>
<td>1-Year Survival</td>
<td>100.0%</td>
<td>89.5%</td>
<td>0.299</td>
</tr>
<tr>
<td>1-Year Freedom from Acute Cellular Rejection (ACR)</td>
<td>81.8%</td>
<td>94.7%</td>
<td>0.106</td>
</tr>
<tr>
<td>1-Year Freedom from Antibody-Mediated Rejection (AMR)</td>
<td>72.7%</td>
<td>78.9%</td>
<td>0.691</td>
</tr>
</tbody>
</table>

Summary of Results This is a 39-year-old Hispanic female with history of PCOS, hyperlipidemia, hypertension, Oral Contraceptive Pill, provoked Deep Vein Thrombosis and Pulmonary Embolism on Rivaroxaban, who presented to emergency department with 4 days of new onset intermittent severe substernal chest pain radiating down to her left arm. She had SARS-CoV-2 pneumonia the month prior to this presentation significant for cough, anosmia, and myalgias, which resolved without hospitalization. On arrival, she was hypertensive, tachycardic, and afebrile. Coagulation panel was normal, troponin-I was elevated at 6.25 with a peak of 9.27. Toxicology was negative for stimulants. She tested positive for SARS-CoV-2 but remained asymptomatic. Patient was started on dual anti-platelet therapy and anti-coagulation therapy. Repeat ECG showed no new changes. A second episode of chest pain revealed lateral ST-elevations and Q-waves in inferior leads. Troponin continued to downtrend. Left heart catheterization was performed with incidental finding of 60% stenosis of the proximal LAD with a smooth plaque. This patient clinically improved without further chest pain and was discharged with dual-antiplatelet therapy.

Conclusions Evaluation and tracking of clinically suspected myocarditis in the setting of COVID-19 infection may give insight into the pathophysiology of infection in cardiomyocytes due to SARS-CoV-2. This case report aims to illustrate the possible association between COVID-19 and myocarditis in the hopes of decreasing morbidity and mortality.

Abstracts

Abstract #84 Table 1

#85 APICAL HYPERTROPHIC CARDIOMYOPATHY MIMICKING AS MYOCARDIAL INFARCTION

1VK Narang*, 2P Chan, 1F Joolhar, 1T Win. 1UCLA-Kern Medical, Bakersfield, CA; 2Ross University School of Medicine, Miramar, FL

10.1136/jim-2022-WRMC.85

Case Report

Purpose Hypertrophic cardiomyopathy (HCM) is known to have a wide spectrum of patterns. This case highlights an uncommon form of HCM called apical hypertrophic cardiomyopathy (ApHCM) which was seen to mimic myocardial infarction.

Methods Retrospective Study.

Summary A 46-year-old Punjabi male with hypertension presented to an outside hospital with chest pain and was found to have elevated troponin levels of 0.31 ng/mL. Nuclear Lexiscan stress test at that time showed ‘reversible defect of the cardiac apex suggestive of ischemia’, cardiac catheterization was negative, and transthoracic echocardiogram (TTE) showed preserved left ventricular function and mild mitral regurgitation. Troponin trended down to 0.23 ng/mL and the patient was discharged.

Patient then comes to the medicine clinic to establish care and was complaining of palpitations that are intermittent and last about 2–3 minutes per episode. Patient reports that the episodes are initiated by physical activity such as walking about 100 feet and alleviated with rest. Patient denied any chest pain or shortness of breath. Positive history for heavy alcohol use, drinks 6–8 alcoholic beverages 2–3 times a week. Electrocardiogram (ECG) done in the clinic showed left ventricular hypertrophy and abnormal T waves in inferior leads. Repeat TTE showed left ventricular ejection fraction is estimated at >65% and apical to mild LV is unusually thickened

A CASE OF POST-COVID MYOCARDITIS IN A 39 YEAR OLD FEMALE WITH PCOS AND INCIDENTAL FINDING OF CAD

S Ratnakar*, A Mesfin, L Moasvi, F Joolhar, A Gandforoush. Kern Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.84

Purpose of Study Reports of cardiovascular manifestations in the setting of COVID-19 have included arrhythmia, pericarditis, heart failure, acute coronary syndrome, and myocarditis. Myocarditis is defined as inflammation of the heart muscle and is commonly associated with viral infection. Common symptoms of myocarditis can include chest pain, shortness of breath, as well as arrhythmia and fatigue. While endomyocardial biopsy remains the gold standard for diagnosis, clinically suspected myocarditis in low-risk patients can be established through presentation and non-invasive diagnostic findings. Here, we aim to highlight the association between coronavirus disease of 2019 infection (COVID-19) and cardiovascular complications such as myocarditis in this case report. In our case, heart catheterization demonstrated 60% stenosis of the proximal left anterior descending artery; however, this lesion was not suspected to be the culprit lesion causing myocardial injury. Etiology of injury was thought to be caused by global ischemia in the setting of post-COVID-19 infection.

Methods Used Retrospective chart review after IRB approval.
which is consistent with ApHCM. Patient was then referred to the cardiology clinic for further management. The patient will be treated with appropriate beta-blocker and cardiac monitoring for further risk stratification.

Conclusion There are many different spectrums to hypertrophic cardiomyopathy with the most common form being asymmetric septal hypertrophy (ASH). There is a more rare form called ApHCM which is more prevalent in the Asian population (25%) than in non-Asians (1% to 10%). Compared to the ASH, it is more sporadic and associated with more atrial fibrillation (AF) and different risk factors for sudden cardiac death (SCD). There are no current guideline recommendations for diagnosis, screening, or patient risk stratification available for ApHCM.

This case illustrates the importance of understanding and diagnosing patients with ApHCM since patient symptoms mimicked a myocardial infarction. The accurate and timely diagnosis may likely improve the clinical outcome and overall well-being of the patient.

Diversity, equity, inclusion I
Concurrent session
3:15 PM
Thursday, January 20, 2022

**#87** STAKEHOLDER ENGAGEMENT EFFORTS TO IMPROVE DIVERSITY IN RARE DISEASE RESEARCH: INSPIRING NEW SCIENCE IN GUIDING HEALTHCARE IN TURNER SYNDROME (INSIGHTS) REGISTRY

1,3,5 Davis*, 1,4 V Bamba, 1,3 W Brickman, 1,3,5 A Carl, 1,4,5 Dowlut-McElroy, 1,4,5 M Good, 1,2,3,5 Howell, 1,8,9,10,12 Kanakatti Shankar, 1,2,3,5 Law, 1,4,8,9,11,12 Prakash, 1,6,7,11,12 K Ranallo. 1INSIGHTS Consortium, Overland Park, KS; 2Children’s Hospital of Colorado, Aurora, CO; 3University of Colorado – Anschutz Medical Campus, Aurora, CO; 4The Children’s Hospital of Philadelphia Division of Endocrinology and Diabetes, Philadelphia, PA; 5Ann Robert H Lurie Children’s Hospital of Chicago, Chicago, IL; 6National Institute of Child Health and Human Development, Bethesda, MD; 7Turner Syndrome Global Alliance, Overland Park, KS; 8Children’s National Hospital, Washington, DC; 9University of North Carolina at Chapel Hill School of Medicine, Chapel Hill, NC; 10The University of Texas Health Science Center at Houston, Houston, TX

10.1136/jim-2022-WRMC.86

Purpose of Study Turner syndrome (TS) occurs in ~1 in 2,000 females who are born with partial or complete absence of the second sex chromosome. Like many rare disease conditions, most research in TS has been focused on specific features (particularly growth), limited to single centers, included minimal diversity, and lacked community engagement. The Inspiring New Science in Guiding Healthcare in Turner Syndrome (InsIGHTS) Registry was developed to address these limitations.

Methods Used A Steering Committee with stakeholders comprised of researchers, multidisciplinary clinicians, and patient advocates was formed to develop the goals, infrastructure, data collection tools, protocols and engagement strategies for a national, collaborative clinic-based longitudinal registry for individuals with TS. Six institutions with multidisciplinary TS clinics across geographical regions were onboarded as recruitment sites with the goal of >80% of eligible patients enrolling with diversity in age, race, ethnicity, payer status, and timing of diagnosis. The team identified patient-centered multidisciplinary outcomes obtainable through medical records and optional additional study procedures.

Summary of Results To date, 154 participants representing all regional centers have enrolled in InsIGHTS with an average enrollment rate of 15 per month. The average age at enrollment of 11.9 ± 11 years (range 0-67, 16.9% ≥18 at enrollment). 18.5% identify as Hispanic/Latinx ethnicity and racial distribution includes 6.2% Asian, 13.7% Black, 71.9% White, and 11.0% Other Race. TS was identified prenatally in 30.3% of participants. The majority of participants agreed to be contacted for future studies (89%), complete annual surveys (83%) and contribute to the biobank (61%).

Conclusions Stakeholder engagement for the development of a national clinic-based registry for the rare genetic condition of TS has successfully led to a diverse cohort representative of the US population. Additional engagement strategies to increase enrollment while prioritizing diversity are underway.

**#88** PARENTAL PERSPECTIVES TOWARDS COVID-19 VACCINES AND RETURN TO SCHOOL: FOCUSING ON HEALTH DISPARITIES

1K He*, 2M Beria, 2M Kung, 2A Afghani. 1Children’s Hospital of Los Angeles, Los Angeles, CA; 2UC Irvine School of Medicine, Orange, CA; 3Children’s Hospital of Orange County, Orange, CA

10.1136/jim-2022-WRMC.87

Purpose of Study The COVID-19 pandemic has disproportionately impacted children from low socioeconomic and minority groups. Parents encounter new decisions regarding vaccinating their child against COVID-19 and return to school in fall of 2021. Prior studies show COVID-19 vaccine hesitancy is associated with income, race, and marital status. However, few studies examine the demographics behind COVID-19 vaccine hesitancy in relation to return to school in vulnerable communities. Understanding both are crucial to addressing challenges for children with healthcare inequities.

Methods Used A cross-sectional survey was conducted at inpatient and outpatient settings at an academic center and its affiliated site between September 2020 - September 2021. Parents were recruited to complete an anonymous mobile-based survey using REDCap regarding perspectives on COVID-19 vaccines and factors affecting children’s return to school during the pandemic. Statistical analyses were performed to examine the association between demographic factors (gender, marital status, education, ethnicity, and household income), COVID-19 vaccine hesitancy, and healthcare inequities affecting return to school.

Summary of Results Of 189 respondents, 65.5% were married, 41.9% had less than college education, and 37.0% had household of > 2 people. 64.6% were minorities and 53.9% were from low income families. COVID-19 vaccine acceptance was positively associated with marital status and number of household members: 60.9% of married individuals reported they would vaccinate their child compared to 30.4% of unmarried individuals (p = 0.001). 62.1% of households of > 2 people would vaccinate compared to 43.1% with households 2 or less (p = 0.015, table 1). Those who accepted or rejected COVID-19 vaccines were more likely to prefer onsite school compared to those who were unsure (p = 0.020). Education, ethnicity, and income were not associated with COVID-19 vaccine acceptance (table 1) or parental decisions in having their child return to school. Those with less than college
Abstract #88 Table 1 COVID-19 vaccine hesitancy and demographic factors

<table>
<thead>
<tr>
<th>Demographic Factor (N)</th>
<th>Yes (%)</th>
<th>Unsure (%)</th>
<th>No (%)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender (171)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mother (143)</td>
<td>68</td>
<td>51</td>
<td>24</td>
<td>0.119</td>
</tr>
<tr>
<td>(47.6%)</td>
<td>(35.7%)</td>
<td>(16.8%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Father (28)</td>
<td>19</td>
<td>5</td>
<td>14</td>
<td></td>
</tr>
<tr>
<td>(67.9%)</td>
<td>(17.9%)</td>
<td>(14.3%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Marital Status (171)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Unmarried (56)</td>
<td>17</td>
<td>25</td>
<td>14</td>
<td>0.001</td>
</tr>
<tr>
<td>(30.4%)</td>
<td>(44.6%)</td>
<td>(25.0%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Married (115)</td>
<td>70</td>
<td>31</td>
<td>14</td>
<td></td>
</tr>
<tr>
<td>(60.9%)</td>
<td>(27.0%)</td>
<td>(12.2%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Educational Level (171)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>College and Up (100)</td>
<td>52</td>
<td>30</td>
<td>18</td>
<td>0.605</td>
</tr>
<tr>
<td>(52.0%)</td>
<td>(30.0%)</td>
<td>(18.0%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Grade and High School (71)</td>
<td>35</td>
<td>26</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>(49.3%)</td>
<td>(36.6%)</td>
<td>(14.1%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Number of Household Members (168)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1–2 people (102)</td>
<td>44</td>
<td>35</td>
<td>23</td>
<td>0.015</td>
</tr>
<tr>
<td>(43.1%)</td>
<td>(34.3%)</td>
<td>(22.5%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3–7 people (66)</td>
<td>51</td>
<td>20</td>
<td>5</td>
<td>0.767</td>
</tr>
<tr>
<td>(82.1%)</td>
<td>(30.3%)</td>
<td>(8.0%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ethnicity (171)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Non-Minority (White, Asian) (81)</td>
<td>32</td>
<td>19</td>
<td>10</td>
<td>0.941</td>
</tr>
<tr>
<td>(52.5%)</td>
<td>(31.1%)</td>
<td>(16.4%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Minority (Hispanic, Black, Pacific Islander) (110)</td>
<td>55</td>
<td>37</td>
<td>18</td>
<td></td>
</tr>
<tr>
<td>(50.0%)</td>
<td>(33.6%)</td>
<td>(16.4%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Income (171)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>High (&gt;50K) (78)</td>
<td>46</td>
<td>21</td>
<td>11</td>
<td>0.151</td>
</tr>
<tr>
<td>(59.0%)</td>
<td>(26.9%)</td>
<td>(14.1%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Low (&lt;50K) (93)</td>
<td>41</td>
<td>35</td>
<td>17</td>
<td></td>
</tr>
<tr>
<td>(44.1%)</td>
<td>(37.6%)</td>
<td>(18.3%)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Abstract

COVID-19 vaccine hesitancy and demographic factors

Purpose of Study

Black individuals in the United States have historically faced barriers to accessing healthcare, only exacerbated by the COVID-19 pandemic. The purpose of this study was to examine the self-reported likelihood of utilizing care within this population in the midst of the COVID-19 pandemic.

Methods

Housing authority residents in Broward County, Florida were asked about their likelihood of visiting their doctor during the COVID-19 pandemic as part of a COVID-19 testing and education initiative conducted by the YMCA of South Florida, in partnership with the Housing Authority of the City of Fort Lauderdale and the Broward County Housing Authority. Secondary data analysis of program data was conducted, including descriptive statistics for describing respondents, chi-square and t-tests for detecting significant differences around likelihood of seeking care between groups, and logistic regression to determine the odds of particular groups’ likelihood of seeking care.

Summary of Results

Significant differences were found between respondents (n=147) reporting they were more likely to visit their doctors in terms of race/ethnicity ($X^2 [1,n=147] = 8.15, p < .01$). Black respondents had three times the odds of claiming to be more likely to visit their doctor (aOR 2.76, 95% CI 1.36–5.60) than other groups. However, Black respondents reported being significantly more afraid of contracting the virus that causes COVID-19 on the way to the doctor’s office than non-Black respondents ($X^2 [1,n=147] = 4.23, p < .05$). Black respondents also reported being more concerned of contracting the virus that causes COVID-19 at the doctor’s office than non-Black respondents ($X^2 [1,n=147] = 5.29 p < .05$).

Conclusions

Black Housing Authority residents seemed to have a high likelihood of utilizing care if needed. However, this high utilization is coupled with a fear of contracting the virus that causes COVID-19 in the process of utilizing care. Areas for further research include determine the rationale behind this positive attitude toward healthcare utilization and figuring out specific areas of concern for contracting COVID-19 in the process of utilizing care (ex. fears of contracting the virus through the use of public transportation on the way to the clinic, etc.).

Abstracts

#90 THE EFFICACY OF EDUCATIONAL WORKSHOPS TO IMPROVE HUMAN PAPILLOMAVIRUS LITERACY IN HISPANIC POPULATIONS

Purpose of Study

Hispanic populations experience disparities with regards to human papillomavirus (HPV) vaccine uptake despite ranking highest among racial groups for rates of cervical cancer. It is well-established that HPV vaccination confers a high degree of protection against HPV-related cancers. Yet barriers to HPV vaccination contribute to low rates of vaccine initiation and series completion in Hispanic populations with only 35–46% of adolescents fully vaccinated against HPV. Notably, literature suggests low health literacy as a common deterrent to vaccine uptake. The purpose of this study is to assess the utility of educational workshops in the improvement
Abstracts

#91 UNDERREPRESENTED IN MEDICINE MINORITY PHYSICIANS IN EMERGENCY MEDICINE RESIDENCY LEADERSHIP

J Smittick, ET Reibling, B Jones, M Kienyeni*; Loma Linda University School of Medicine, Loma Linda, CA; Loma Linda University, Loma Linda, CA

Purpose of Study It’s evident in academic literature that representation of underrepresented in medicine (URiM) minorities in Emergency Medicine (EM) is sparse. This disparity is more drastic amongst EM leadership. Faculty and residents are directly involved in recruiting, interviewing, and ranking potential incoming residents. The lack of URiM participation in these processes impacts the potential for future URiM physicians to be appointed to EM residency leadership positions. Our study sought to identify potential areas for increased representation in the future and factors that may increase URiM involvement.

Methods Used We administered a survey to the U.S. Emergency Medicine Residency Program Directors (PDs) listed on FREIDA, the American Medical Association (AMA) residency and fellowship database. We drafted and piloted the online survey instrument before sending it to participants via Qualtrics. Survey items focused on ethnic identity in program leadership, career preparation such as mentors and previously held roles, and strategies used to encourage URiM recruitment. Participants received one announcement email and three reminder emails following the survey distribution. We used Microsoft Excel for primary data analysis.

Summary of Results We received 57 completed surveys. 22% of respondents identified as URiM, of which 9% identified as Black and 7% identified as Latinx. The median percentage of Residents identifying as URiM was 13% (IQR: 1%-32%). Eight programs (14%) reported having at least one Chief Resident identifying as URiM. 72% of respondents reported that a mentor was instrumental in their ascension to PD. 11% reported that their mentor identified as URiM. We asked PDs to confirm which strategies they’ve implemented to encourage URiM participation (Boatright 2008). The most commonly implemented strategies were, ‘Know the institution’s local and community demographics, and address those needs’ (51%), followed by ‘Broaden selection criteria beyond USMLE scores to include intangibles such as leadership, community service, and other life experiences’ (49%), and ‘Develop curricula to address topics on diversity, cultural competence, and implicit bias’ (47%).

Conclusions The disparity of URiM PDs in EM may be a result of a lack of URiM mentorship. 29% of respondents were URiM but only 11% reported having a URiM mentor. This lack of mentor-mentee concordance may be an area of further study and improvement. More intentional utilization of URiM recruitment strategies could also drastically improve representation. Increased URiM participation in EM leadership has great potential to improve diversity, equity, and inclusion in EM overall.


#92 NOVEL HEALTH EQUITY ADVOCACY CURRICULUM TO INCITE ENGAGEMENT IN MEDICAL TRAINEES

C Corkin*, N Rodman, D Smith, W Joshi; University of California San Diego School of Medicine, La Jolla, CA; Rady Children’s Hospital San Diego, San Diego, CA

Purpose of Study Medical education health equity curriculums rarely emphasize advocacy and community engagement, further exploiting the minority tax in pursuing health equity work. Health equity curriculums must include three components: history, outcomes and interventions. The Journal Club and Advocacy Lab (JC-AL) schema was added to the Health Equity Thread (HET) preclinical curriculum at UC San Diego (UCSD) School of Medicine to teach and support interventions to health disparities.

Methods Used Preclinical students receive HET credit by attending JC-ALs. JC-AL workflow is depicted in figure 1; the JC and AL are held 1–2 weeks apart. Participants took a survey, approved by the UCSD Institutional Review Board, before the JC and after the AL. Survey responses from November 2020–June 2021 were gathered and summarized for each time-point using R.

Summary of Results Of participants surveyed, 141 (28.5%) identified as underrepresented in medicine. About a quarter of participants saw an increase in mood (25%), resilience (27%).
Abstract #92 Figure 1  Workflow for JC-AL

sense of community (24%) and/or motivation (29%) regarding health equity work after the intervention. 158 participants (67.2%) reported being somewhat or very likely to stay involved in the advocacy project, and 93 participants (39.6%) reported being likely to lead a session in the future. Almost all of the JC-ALs have manifested long term projects including:

- Educational material for healthcare providers and preclinical students regarding removal of race from eGFR and adoption of cystatin C and addition of cystatin C in UCSD Health Laboratory Medicine Formula
- Learning modules for preclinical students on gender affirming and trauma informed care
- Elective on obtaining a health equity history in the emergency department

Conclusions The JC-AL schema is a feasible approach to engage trainees in the community and institution to enact change. It is a well-received component of the HET ranging from 30–100 participants at each event.

Abstract #93 USING VALIDATED INSTRUMENTS TO ASSESS LONELINESS, SOCIAL SUPPORT, AND BURNOUT AMONGST RACIAL AND LGBTQ+ RESIDENTS

DA Mesa*, University of Colorado, Denver, CO

10.1136/jim-2022-WRMC.92

Purpose of Study There is very little data published exploring the impact that racial or sexual minority identity has on a resident’s training experience. Given that a high percentage of internal medicine training programs are predominately white it’s important to understand the emotional and supportive barriers minority residents face. We began the important work looking into these barriers with a survey-based needs study.

Methods Used 174 residents enrolled in the University of Colorado Internal Medicine Residency Program were asked to participate in an online survey. This survey consisted of several validated instruments including: the PHQ-4, the MOS Social Support Survey, and the UCLA Loneliness Scale. The survey included a demographics section and each respondent utilized a unique PID to maintain anonymity. 65 out of 174 residents responded to the survey. The answers to the survey were coded and scored per the original publications. Analysis of the data was done using two tailed T-Tests in the SAS software.

Summary of Results The average MOS total support score was significantly lower in LGBTQ+ residents compared to Non-LGBTQ+ residents (Mean 65.80 vs. 79.16; P = 0.035). LGBTQ+ residents also trended towards having higher amounts of burnout, though this wasn’t statistically significant (Mean 0.38 vs 0.17; P = 0.19). Notably 3 out of 9 LGBTQ + residents reported feeling burnt out compared to 9 out of 56 Non-LGBTQ+ residents (33% vs 16%). Notably most of the significant findings were amongst single vs non-single residents with significance in: UCLA loneliness scale (P = 0.03), MOS total support score (P < 0.0001), MOS emotional support score (P = 0.008), MOS affectionate support score (P < 0.0001), MOS tangible support score (P < 0.0001), MOS positive interactions score (P = 0.001) and PHQ-Depression sub-domain (P = 0.025). Racial minority residents had lower average levels of burnout compared to non-minority residents (Mean 0.12 vs 0.23; P = 0.32). However, racial minority residents had lower average levels of overall social support compared to non-minority residents (Mean 73.16 vs 78.74; P = 0.270) with the tangible support subdomain score being the closest for significance (Mean 13.55 vs 15.93; P = 0.120).

Conclusions The sample size for the survey-based study was smaller than anticipated. However, it was large enough to find significance for LGBTQ+ residents having less social support, and also revealed higher levels of burnout. It’s also important to note that while it didn’t reach significance minority residents experienced lower average levels of overall social support. Surprisingly minorities had fewer burnout numbers which may be an indicator of increased resilience or utilization of protective mechanisms. Further research needs to be conducted to better understand the needs of LGBTQ+ and racial minority residents. Future directions include expansion nationwide to gather a larger sample size and assess for geographic differences.

Abstract #94 IDIOPATHIC SUBGLOTTIC STENOSIS IN NON-CAUCASIAN WOMEN

1A Suk*, 1L Reder, 1K O’Dell, 5S Verna, 5M Harmon, 5P Weissbrod, 5P Krishna, 5B Crawley, 1Loma Linda University School of Medicine, Loma Linda, CA; 2Kaiser Permanente Baldwin Hills, Los Angeles, CA; 3University of Southern California, Los Angeles, CA; 4University of California Irvine, Irvine, CA; 5University of California San Diego, La Jolla, CA; 6Loma Linda University School of Medicine, Loma Linda, CA

10.1136/jim-2022-WRMC.93

Purpose of Study To analyze the presentation, disease course, and treatment of idiopathic subglottic stenosis of non-Caucasian women.

Methods Used In this multi-institutional retrospective study, information extracted included date of birth, age at symptom onset, age and date of diagnosis, race, Cotton Meyer grade, stenosis length and distance from glottis, BMI, comorbidities, medication to manage iSGS, age at first surgery, additional treatment with serial intraläsional steroid injections, the history of each surgery, occupation, autoimmune labs, and family history of autoimmune diseases.

Summary of Results 35 non-Caucasian women with idiopathic subglottic stenosis were identified. Of the 35 women, 31 were Hispanic while one was African-American, two were Asian, and one was non-Hispanic mixed race. Their average BMI was 31.8 ± 2.19 kg/m² and 51.4% of the patients were obese (BMI>30). 31.4% had hypertension. Their average age of onset was 45.8 years old (95% CI, 42.2–49.3) with a range of 26–69 years old. The average age at diagnosis was 47.8 years (95% CI, 44.3–51.3) with a Charlson comorbidity index of 0.85 (95% CI, 0.42–1.28). At diagnosis, 13.4% were CM I, 43.3% were CM II and 43.3% were CM III (n=30). The average age at their first surgery was 46.8 (95% CI, 43.2–50.4) years and 17 received SILLI. While treatment type

J Investig Med 2022;70:110–345

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Abstracts

#95 SCOPING REVIEW OF SOCIOECONOMIC FACTORS AND HIDRADENITIS SUPPURATIVA

1OS Cherepakhin*, 2K Deliro. 1University of Washington School of Medicine, Seattle, WA; 2University of Washington, Seattle, WA

Purpose of Study Hidradenitis suppurativa (HS) is an autoimmune inflammatory disease characterized by painful boils beneath the skin. While socioeconomic factors have been linked to HS individually, there has been no scoping review that synthesizes these correlations. Our objective was to assess the published data on the associations between HS and the factors of income, education, and work.

Methods Used A search limited to English publications was conducted in PubMed, Embase, Web of Science and Cochrane from database origin to 07/26/21. The terms used were ‘hidradenitis’ combined with ‘socioeconomic,’ ‘insurance,’ ‘class,’ ‘disparities,’ ‘disparity,’ ‘education,’ ‘income,’ ‘work,’ ‘employment,’ ‘job,’ ‘insurer,’ ‘medicaid,’ or ‘professional activity.’ Eligible publications were peer-reviewed and examined the association between HS and income level, educational attainment, occupation class, employment status, work impairment, or insurance status. Records were evaluated by O.C. and K.D. In the event of a disagreement, another reviewer was available to resolve the discrepancy.

Summary of Results After duplicate removal, 413 records were screened by title/abstract. 79 full-text records were then assessed for eligibility and 33 articles met inclusion criteria. By manually searching article references, an additional 3 papers were included. 29 research articles, 6 reviews, and 1 case report from 13 different countries were qualitatively synthesized according to the defined categories of associations.

3 articles found that HS patients had lower income levels but one of those studies, after adjusting for age/sex, found that this was not significant. 6 articles elucidated an association between HS and lower educational attainment. An association between HS and lower class of occupation was found by 1 study, and 7 publications (6 articles, 1 review) demonstrated a higher probability of being unemployed as an HS patient. 16 articles, 1 case report, and 5 reviews discussed the association between HS and work impairment. A higher likelihood for HS patients to have government-funded insurance was found by 3 studies. 4 articles utilized a combination of the factors as measures of SES. 3 of them found associations between low SES and HS, while one Israeli study found the opposite.

Conclusions Our qualitative synthesis demonstrates that HS globally is linked with lower income levels, reduced educational attainment, unemployment, work impairment, and government-funded insurance coverage. Though one study found that higher SES is associated with HS, this can be explained by their usage of dermatologist-diagnosed HS patients and the fact that in Israel, dermatology encounters require co-payments unlike primary care visits. Though the directionality between HS and lower SES cannot be determined from the current research, our work shows the importance of considering SES when treating HS patients.

#96 AMERICAN OSTEOPATHIC ASSOCIATION OTOLARYNGOLOGY AND OPHTHALMOLOGY PROGRAM CLOSURES AS A MODEL TO HIGHLIGHT CHALLENGES OF MAINTAINING GRADUATE MEDICAL EDUCATION IN HIGH NEED AREAS

1K Vo*, 2H Ahmed, 3V Robbins. 1Western University of Health Sciences, Pomona, CA; 2Loma Linda University, Loma Linda, CA; 3OhioHealth, Columbus, OH

Purpose of Study While 90% of former American Osteopathic Association (AOA) residency programs transitioned to Accreditation Council for Graduate Medical Education (ACGME) accreditation, surgical subspecialty programs such as otolaryngology (ENT) (62%) and ophthalmology (47%) struggled to gain accreditation. DOs have actively participated in serving underserved communities, and losing AOA surgical specialty programs may decrease access to surgical care in rural and non-metropolitan areas.

Methods Used A directory of former AOA ENT and ophthalmology programs was obtained from the American Osteopathic Colleges of Otolaryngology and Ophthalmology-Head and Neck Surgery (AOCOO-HNS). A secured survey was sent to 16 eligible ENT and ophthalmology program directors. The survey contained both quantitative and qualitative aspects to help assess why these programs did not pursue or failed to receive ACGME accreditation.

Abstract #96 Table 1 Percentage of remaining osteopathic ENT and ophthalmology programs from 2014–2015 to 2020–2021 academic year under the SAS

<table>
<thead>
<tr>
<th>Specialty</th>
<th>2014-2015</th>
<th>2020-2021</th>
<th>% Total Programs Remaining</th>
<th>% Initial Accreditation After Merger</th>
<th>% Continuing Accreditation</th>
<th>% Continuing Accreditation without Outcomes</th>
<th>Net Decline</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ophthalmology</td>
<td>15</td>
<td>7</td>
<td>46.7%</td>
<td>26.7%</td>
<td>25.0%</td>
<td>6.7%</td>
<td>-53.3%</td>
</tr>
<tr>
<td>Otolaryngology</td>
<td>21</td>
<td>13</td>
<td>61.9%</td>
<td>47.6%</td>
<td>0.0%</td>
<td>14.3%</td>
<td>-38.1%</td>
</tr>
</tbody>
</table>

Table 1 depicts a net decline of 53.3% and 38.1% in osteopathic ophthalmology and ENT programs from 2014-2015 to 2020-2021 academic year under the SAS.
Abstract #96 Figure 1

Summary of Results 12 of 16 eligible programs responded: 6 ophthalmology and 6 ENT program directors. 83% of respondents did not pursue accreditation (6 ophthalmology and 4 ENT programs), and 17% (2) were unsuccessful in achieving accreditation despite pursuing accreditation. Across 12 respondents, 58% (7) cited lack of hospital/administrative support and 42% (5) cited excessive costs and lack of faculty support as reasons for not pursuing or obtaining ACGME accreditation.

Conclusions The survey results reflect financial issues associated with rural hospitals. Lack of hospital/administrative support and excessive costs to transition to the ACGME were key drivers in the closures of AOA surgical specialty programs. Considering these results, we have 4 recommendations for various stakeholders, including program directors, designated institutional officials, hospital chief medical officers, and health policy experts. These recommendations include expanding Teaching Health Center Graduate Medical Education to surgical subspecialties, identifying and learning from surgical fields such as urology that fared well during the transition to ACGME, addressing the lack of institutional commitment and prohibitive costs of maintaining ACGME accredited subspecialty programs in under-resourced settings, and reconsidering Centers for Medicare & Medicaid Services (CMS) pool approach to physician reimbursement.

Hematology and oncology I
Concurrent session
3:15 PM
Thursday, January 20, 2022

#97 NOVEL PERINEPHRIC NEOVASCULARITY SCORING SYSTEM IN RENAL CELL CARCINOMA TUMOR STAGING

C Fateri*, A Peta, N Kar, T Bui, B Roth, J Glavis-Bloom, L Limfueco, J Landman, R Houshyar, University of California Irvine, Irvine, CA

10.1136/jim-2022-WRMC.96

Purpose of Study Renal cell carcinoma (RCC) is the most common type of kidney cancer worldwide. Angiogenesis plays a major role in providing adequate blood flow and nutrients to promote tumor growth and RCC progression. While radiologists assess enhancement patterns of renal tumors to predict tumor pathology, to our knowledge, no formal scoring system has been created and validated to assess the level of neovascularity in RCC, despite its critical role in cancer metastases. In this study, we characterized and analyzed the level of angiogenesis in tumor-burdened kidneys and their benign counterparts. We then created and validated a scoring scale for neovascularity that can help predict tumor staging for RCC.

Methods Used After Institutional Review Board approval, the charts of patients who had undergone surgery for RCC between January 13, 2014 and February 4, 2020 were retrospectively reviewed for inclusion in this study. Inclusion criteria were a diagnosis of RCC, simple/radical nephrectomy, pre-operative contrast enhanced computed tomography (CT) scans, and complete pathology reports. Neovascularity was scored on a scale of 0 to 4 where 0= no neovascularity detected, 1= a single vessel <3 mm wide, 2= a single vessel ≥3 mm wide, 3= multiple vessels <3 mm wide, and 4= multiple vessels ≥3 mm wide. Each patient was scored by a senior medical student and then validated by a board-certified abdominal radiologist. Statistical analysis was performed using RStudio® Version 3.5.1. Demographics and tumor characteristics were compared using a Kruskall-Wallis ANOVA or Chi-squared test; neovascular score was compared using a Wilcoxon Rank-Sum test. Statistical significance was considered as p < 0.05.

Summary of Results A total of 217 patients were included in this study. There was no significant difference in patient demographics between tumor stages. Additionally, the majority of tumor pathology was clear cell carcinoma, regardless of tumor staging. The average neovascularity score was 1.07 for pT1x tumors, 2.83 for pT2x tumors, and 3.04 for pT3x tumors. The average neovascularity score for the benign counterparts was 0.124, 0.385, and 0.458, respectively. There was a significant difference in neovascularity score between pT1x and pT2x tumors (p = 0.0046), pT1x and pT3x tumors (p <
0.0001), and benign kidneys and kidneys with RCC (p = 0.0001).

Conclusions Our novel vascular scoring system for renal cell carcinoma demonstrates a significant correlation with RCC pathological tumor staging. This scoring system may be utilized as part of a comprehensive radiological assessment of renal tumors, potentially improving tumor characterization and clinical decision making.

Abstracts

#98

STRESS-INDUCED DIFFERENTIAL MIR-4633–5P EXPRESSION IN THYROID CANCER HEALTH DISPARITIES

1) Mancao*, 1) S. Khan, 1) J. Lee, 1) R. Rood, 1) R. Davis, 1) M. Perez, 1) A. Simental, 1) S. Roy, 1) K. Loma Linda University School of Medicine, Loma Linda, CA; 1) Loma Linda University Department of Basic Sciences, Loma Linda, CA; 1) Loma Linda University Medical Center, Loma Linda, CA;

1) The University of Texas at El Paso, El Paso, TX

10.1136/jim-2022-WRMC.97

Purpose of Study Filipino Americans (FA) are known to have higher rates of thyroid cancer incidence and disease recurrence compared to European Americans (EA). FA are also known to be two times more likely to die of thyroid cancer compared to EA. Epidemiological studies in California have shown that thyroid cancer is the second most common cancer among FA women. Currently, there are no studies that demonstrate the mechanism behind these discrepancies. Evidence shows a strong correlation between obesity and more aggressive forms of thyroid cancer; obesity has an increased frequency in FA populations. The exact connection between the mechanisms of obesity and cancer is poorly understood. This epigenetic phenomenon may be due to microRNAs (miRNAs), which post-transcriptionally regulate gene expression. Dysregulated miRNA profiles have been associated with various diseases including obesity and cancer. MiRNAs are linked to different types of cancer; tumor suppressor genes and oncogenes are subject to modulation by dysregulated miRNAs. No study elucidates the association of miRNAs to tumor staging or prognosis in thyroid cancer health disparities.

Methods Used In this study, we determined miRNA expression profiles and found significant differences in the miRNA profiles between FA and EA thyroid cancer patients. Our pilot study showed several dysregulated miRNAs, from which we chose to assay dysregulated miR-4633–5p segments that are known to be associated with thyroid cancer signaling. We used QIAGEN’s miRNA extraction kit to obtain high-quality miRNA from paraffin-embedded thyroid tissues. We performed next-generation miRNA sequencing using equal number of FA and EA samples and identified the top ten significantly up- and down-regulated miRNAs from the pool of differentially expressed miRNAs by qPCR assays.

Summary of Results Our investigation demonstrated a 1.5–2-fold higher expression of an upregulated miR-4633–5p in FA versus EA miRNA samples (n = 70) after normalization to controls. In contrast, miR-323b–3p showed no difference between FA and EA after normalized to controls.

Conclusions For our future work, we plan to analyze multiple up- and down-regulated miRNAs by qPCR, determine whether the miRNA signatures are consistent between samples from FA versus EA, and explore the use of these miRNA signature differentials for affordable and rapid thyroid cancer screening and prognosis.

#99

MELANOMA OF UNKNOWN PRIMARY: A SINGLE-INSTITUTION EVALUATION OF CLINICAL OUTCOMES

1) S. Dwabe*, 1) G. In, 1) Los Angeles County University of Southern California Medical Center, Los Angeles, CA; 1) Keck Hospital of USC, Los Angeles, CA

10.1136/jim-2022-WRMC.98

Purpose of Study Melanoma of unknown primary (MUP) is clinically uncommon and is understudied as a disease. There have been studies evaluating the utility of local resection with radiation therapy for treatment of MUP. However, it has been only within the last few years that MUP has been routinely treated with targeted or immunotherapy.

Methods Used We conducted a retrospective review of patients with MUP treated at LAC-USC Medical Center and Norris

Abstract #99 Table 1 Clinical outcomes

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Median overall survival, all patients (months, range) Total 1 year survival</th>
<th>Median overall survival, patients treated with radiation (months, range) Total 1 year survival</th>
<th>Median overall survival, patients treated with immunotherapy (months, range) Total 1 year survival</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 organ (n = 16)</td>
<td>14.3 (0.2 – 72.6)</td>
<td>13 (100%)</td>
<td>8.6 (85.7%)</td>
</tr>
<tr>
<td>2+ organs (n = 14)</td>
<td>15 (83.3%)</td>
<td>10 (71.4%)</td>
<td>66.2 (99.3%)</td>
</tr>
<tr>
<td>3 (n = 18)</td>
<td>24.8 (7.1 – 35.9)</td>
<td>6 (87.5%)</td>
<td>11 (100%)</td>
</tr>
<tr>
<td>4+ (n = 12)</td>
<td>23.7 (0.6 – 66.2)</td>
<td>16 (88.9%)</td>
<td>10 (100%)</td>
</tr>
<tr>
<td>5 cm (n = 12)</td>
<td>35.0 (12.1 – 66.2)</td>
<td>35.0 (12.1 – 66.2)</td>
<td>35.0 (12.1 – 66.2)</td>
</tr>
<tr>
<td>Largest met</td>
<td>37.0 (12.1 – 72.6)</td>
<td>37.0 (12.1 – 72.6)</td>
<td>37.0 (12.1 – 72.6)</td>
</tr>
<tr>
<td># of metastases, months, range</td>
<td>8 (66.7%)</td>
<td>8 (66.7%)</td>
<td>8 (66.7%)</td>
</tr>
<tr>
<td>Site</td>
<td></td>
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</tr>
</tbody>
</table>
Cancer Center from 2008 to December 1st, 2020. We recorded the presentation, treatment course, and outcomes of each patient within our database. Data points collected include demographic information, clinical staging, size of largest metastases, location and number of metastatic sites. Treatment modalities, including metastatectomy, and systemic therapy were reviewed. The primary outcomes studied were median overall survival and 1-year overall survival.

**Summary of Results** Data was collected from 32 patients identified as having MUP. Sites of melanoma metastases included lymph node, soft tissue/muscle, lung, liver, brain/leptomeningeal, and bone. Thirteen patients (40.6%) were found to have one metastatic tumor, 6 (18.8%) were found to have 2–3 metastases, and 13 (40.6%) were found to have 4+ metastatic tumors on presentation. Two patients were lost to follow-up shortly after their diagnosis. The 30 remaining patients had a 14.3-month median survival with 17 (56.7%) surviving past one year.

In total, 15 patients underwent surgical metastatectomy, and 18 patients were treated with immunotherapy. With all patients surviving after one year, the 13 patients who had a complete resection of their tumor burden were noted to have a median survival time of 37.0 months following their diagnosis compared to a 2.2-month median survival among the 17 patients who did not have surgery or who had incomplete resection of tumor. The 18 patients treated with immunotherapy (PD-1 + CTLA-4 inhibition) were found to have a median survival time of 23.7 months with 16 (88.9%) surviving past one year. Eleven patients who were treated both with complete surgical resection and adjuvant immunotherapy, were found to have a median survival time of 35.0 months with 11 (100%) surviving past one year. When analyzing outcomes of patients with MUP based on the number of metastases, number of organs involved, and largest size of metastases, survival was correlated with less than 4 metastases and less than 2 organs involved. Size of largest metastases had no effect on survival outcomes.

**Conclusions** Outcomes among patients with MUP may vary, depending on treatment modality, and tumor burden. Based on our data, patients who have MUP with low burden of disease may benefit from multi-modality therapy, including both surgical metastatectomy, and immune checkpoint blockade. Further validation using larger cohorts is warranted to help confirm these findings.

**#101** IMPACT OF SPONSOR ON ADVANCED NON-SMALL CELL LUNG CANCER CLINICAL TRIAL ENROLLMENT CRITERIA

1RA Cooper*, 2Y Chai, 1J Nieva. 1University of Southern California Keck School of Medicine, Los Angeles, CA; 2Children’s Hospital of Los Angeles Saban Research Institute, Los Angeles, CA

10.1136/jim-2022-WRMC.100

**Purpose of Study** Clinical trials use inclusion and exclusion criteria to control for confounding variables in patient populations. Largely inspired by the ASCO-Friends of Cancer Research recommendation documents (2017 and 2021), there has been a recent drive to loosen clinical trial enrollment criteria to improve generalizability in trial outcomes. We sought to determine if the sponsor of a clinical trial impacted the transparency and selection of inclusion and exclusion criteria.

**Abstract #101 Figure 1** Percentage of studies with strict, loose, and no restrictions on performance status. Actual number of studies within each group are included as data labels
Methods Used Using clinicaltrials.gov, phase 2 and 3 non-small cell lung cancer (NSCLC) drug trials were sorted into one of three sponsor categories: Industry, government/cooperative group, and academic. Fisher Exact tests were used to assess variability in strictness of specific criteria and level of transparency in listing organ function requirements. Independent sample t tests were used to analyze differences in total number of criteria. Summary of Results Industry sponsored NSCLC drug trials more often omit from clinicaltrials.gov complete organ function requirements compared to government/cooperative group (p = 2.3 x 10^-10, α = 0.01) and academic (p = 1.8 x 10^-4, α = 0.01) sponsored trials. Industry sponsored trials are also more likely to have stricter performance status requirements compared to government/cooperative group sponsored studies (p = 5.7 x 10^-5, α = 0.01). Conclusions Industry funded NSCLC clinical trials are more rigorous in excluding patients with worse performance status and are less transparent in listing all study requirements on clinicaltrials.gov.

#102 ASSESSING ADULT PATIENTS’ UNDERSTANDING OF TERMS IN CONSENT RELATED TO SECONDARY MALIGNANCY RISK IN RADIATION THERAPY

N Vartanian*, M Wilson, R Eroonian. University of Washington School of Medicine, Seattle, WA

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Purpose of Study Informed consent entails that healthcare providers effectively describe adverse effects associated with medical treatments to patients. In radiation oncology, the terms ‘second tumors’, ‘secondary malignancies’, or ‘secondary tumors’ are used in patient consents to describe the appearance of new and different tumors caused by radiation treatment. Furthermore, these incidences are sometimes described in consents as ‘rare’, although the incidence varies greatly from nearly negligible in patients treated with palliative intent, to 20% in patients undergoing myeloablative total body irradiation for stem cell transplant. We evaluated whether non-cancer patients without prior knowledge of or exposure to radiation therapy interpret the terms ‘secondary malignancy’, ‘rare’, and ‘small chance’ in a way consistent with physician intent.

Methods Used We screened 164 adult subjects who did not require medical interpreters at a university affiliated family medicine clinic, excluding cancer patients and those with prior knowledge of or experience with radiation treatment. One hundred subjects were eligible for and completed our 12-question multiple choice questionnaire, which assessed their understanding of the term ‘secondary tumor’, and how they would interpret the terms ‘small chance’ or ‘rare’ in the context of a ‘bad side effect’ arising from medical treatment. Summary of Results Twenty-nine percent of subjects correctly identified that ‘secondary tumors’ referred to new and different tumors caused by treatment. Forty-nine percent thought the term referred to their original tumor coming back, and twenty-two percent thought the term referred to new and different tumors not caused by radiation therapy. In the context of a ‘bad side effect’ occurring ‘rarely’, 2% of subjects attributed ‘rare’ to a 1/10 chance; 16% to a 1/100 chance; 33% to a 1/1000 chance; and 49% to a 1/100,000 chance.

In the context of a ‘bad side effect’ having a ‘small chance’ of occurrence, 8% of subjects attributed ‘small chance’ to odds of 1/10; 33% to 1/100; 41% to 1/1000; and 18% to 1/100,000.

Conclusions Patients without prior radiation therapy exposure have a demonstrably different understanding than radiation oncologists of the terms ‘secondary malignancy’, ‘second tumor’, or ‘secondary tumor’. Additionally, there is great variability in patient understanding of the terms ‘rare’ or ‘small chance’. Radiation oncologists must use different and more descriptive terms for secondary malignancies and their incidence, to ensure patients are truly informed when undergoing treatment. The results of this study may have implications for all medical fields in which patients are consented for procedures associated with a risk for consequential side effects.

#103 REGULATION OF CHEMORESISTANCE BY CHRM1 IN NON-SMALL CELL LUNG CANCER

S Simpson*, T Bland. WWAMI, Seattle, WA

10.1136/jim-2022-WRMC.102

Purpose of Study Chemotherapy is a mainstay treatment for late-stage non-small cell lung cancer (NSCLC), yet most tumors develop resistance to these agents. Studies in our lab have shown that chemoresistant NSCLC cells overexpress the muscarinic acetylcholine receptor 1 (CHRM1). We hypothesize that CHRM1 regulates chemoresistance in NSCLC cells, and that the combination of a repurposed CHRM1 antagonist dicyclomine, clinically used to treat IBS, and a chemotherapeutic agent has the potential to sensitize and kill chemoresistant NSCLC cells.

Methods Used Chemosensitive (A549) and chemoresistant (A549R) NSCLC cells were utilized in this study. Cell survival and colony formation assays were utilized to measure DTX sensitivity by pretreating with designated drug (24 hr) before addition of DTX (48 hr). Western blot and phosphokinase array were utilized to measure protein expression and intracellular pathway activation. The designed receptors exclusively activated by designer drugs (DREADD) system was utilized to isolate CHRM1 signaling. All data are expressed as the mean ± SEM. Multiple comparisons were analyzed using one-way ANOVA with post-hoc Tukey’s analysis and single comparisons were analyzed using a two-tailed, unpaired Student’s t test.

Summary of Results CHRM1 expression is enhanced in A549R cells, suggesting that CHRM1 may play a role in chemoresistance. This was supported by the ability of a CHRM1 agonist, dicyclomine (Dic), to sensitize A549R cells to the chemotherapeutic agent docetaxel (DTX) measured by cell survival (IC50: DTX, not reached > 1M; DTX + Dic (10µM), 49.91µM; DTX + Dic (25µM), 12.11µM). Furthermore, these results were duplicated by colony formation assay. However, activation of a CHRM1 in A549 cells by the acetylcholine mimetic carbachol did not protect cells from DTX-induced cell death, suggesting that CHRM1 expression is necessary for chemoresistance in the A549R cells, but not sufficient. A phosphokinase array was used to determine the intracellular signaling pathway activated by CHRM1, which showed increased phosphorylation of multiple kinases including CREB, EGFR, STAT3, and ERK1/2. Increased CREB phosphorylation was
validated by western blot with carbachol stimulation in A549R and M1D samples suggesting these as possible targeting pathways downstream of CHRM1.

**Conclusions** Chemoresistant NSCLC shows increased CHRM1 expression, which when antagonized, resensitizes these cells to DTX-induced cell death. While CHRM1 expression is not sufficient to instill resistance, it is necessary in the A549R cells, and may play a role in enhancing EGFR signaling. This provides a potential promising new therapy for lethal chemoresistant NSCLC which utilizes the repurposed IBS drug dicyclomine.

### #104 TISSUE IS THE ISSUE: CHEMOTHERAPY RESPONSE SCORE (CRS) IS MOST PREDICTIVE OF RESPONSE TO NEOADJUVANT CHEMOTHERAPY IN ADVANCED, HIGH GRADE SEROUS OVARIAN CANCER

1. Knickerbocker*, 2K Kuchta, 3C Donaldson, 2E Diaz Moore, 2M Lippitt, 2G Rodriguez, 2W Watkin, 2T Jenkins Vogel. 1Washington State University, Spokane, WA; 2NorthShore University HealthSystem, Evanston, IL

10.1136/jim-2022-WRMC.104

**Purpose of Study** Favorable chemotherapy response score (CRS) has prognostic value and correlates with progression free and overall survival in advanced ovarian cancer. CRS has not been compared to other clinical measures used to gauge response to neoadjuvant chemotherapy (NACT). We sought to examine whether CRS is a better predictor of outcome compared to traditional clinical and radiographic response measures.

**Methods Used** Clinical data from 2003–2020 was obtained through retrospective chart review. Radiographic review pre- and post-NACT was performed via RECIST 1.1 with responses characterized as complete/near-complete (CR/NCR), partial with >50% reduction in tumor (PR>50), partial with <50% reduction in tumor (PR<50), stable disease (SD) and progressive disease (PD). Histologic response in surgical specimens was characterized using CRS 1–3. Survival was assessed using the Kaplan-Meier method with log-rank tests, and Cox regression with hazard ratios (HR).

**Summary of Results** 128 patients who underwent NACT for high grade serous ovarian cancer (HGSOC) were included. Increasing CRS was associated with improved recurrence free (RFS) and overall survival (OS). OS at 5 years for CRS 1, 2 and 3 was 24.7%, 57% and 73.7% (p<0.0001). More favorable radiographic response was predictive of decreased recurrence risk with RFS at 3 years for PR<50 and CR/NCR (16.3% and 54.8%, p=0.0005) but not predictive of OS. Patients with CR/NCR more commonly had CRS 3 vs CRS 1 (47.1 vs 17.7%, p=0.022). Among radiographic response groups, increasing CRS was associated with decreased risk of recurrence and death. For example, in patients with CR/NCR risk of recurrence HR 5.38 (p=0.0243) and risk of death 8.24 (p=0.006) with CRS 1 vs 3. Number of NACT cycles prior to surgery did not differ among patients regardless of CRS. Rate of R0 resection was similar among all three CRS subgroups, and for the entire cohort was 85.2%. Recurrence rates were significantly higher with CRS 1 (89.1%) and 2 (73.9%) compared to CRS 3 (38.9%) (p<0.0001). Median CA 125 prior to surgery was lower with CRS 3 compared to CRS 1 (28 vs 81, p=0.0017). Of the 12 germline BRCA2+ patients in the study, 7 (58.3%) had a pathologic CRS 3.

**Conclusions** Our data confirms that favorable CRS is associated with improved overall and recurrence free survival in HGSOC. While radiographic response appears predictive of recurrence, it was not associated with overall survival in our study. Among patients with similar radiographic response, CRS remained predictive of outcome and is associated with other clinical factors traditionally felt to confer favorable prognosis. Pathologic CRS is an important predictive factor in determining response to neoadjuvant chemotherapy in HGSOC and may provide the best means to characterize prognosis.

### #105 THE IMPACT OF AN EDUCATIONAL TRAINING PROGRAM ON COVID-19 AND CANCER: ENHANCING COMMUNITY UNDERSTANDING, FOR DISEASE PREVENTION, BETTER TREATMENT, AND OPTIMAL OUTCOMES

J Miller*, E McGhee. Charles Drew University of Medicine and Science, Los Angeles, CA

10.1136/jim-2022-WRMC.104

**Purpose of Study** The CDC reported cancer patients as at-risk for severe illness from severe acute respiratory syndrome-related coronavirus 2 (COVID-19). Cancer patients were 2 times more likely than non-cancer patients to exhibit cellular sequelae due to COVID-19. Those with hematological malignancies exhibited a case-fatality rate 2 times more than those with solid tumors. This research aims to educate and enhance community understanding of factors that lead to increased mortality rates in COVID-19 cancer patients by using a community training program in SPA 6 of Los Angeles, California.

**Methods Used** Data were obtained from post-training surveys of SPA 6 community members which included cancer diagnosis, demographics, and knowledge of COVID-19 with cancer. Impact assessments utilizing Likert-scale response options to analyze and measure the data. Fisher’s exact test was utilized to measure and evaluate participant understanding of the community training program in regards to increased mortality rates of COVID-19 cancer patients. Data analyses were performed using statistical tests SPSS. P-values of <0.05 were considered significant.

**Summary of Results** There is a significant need for COVID-19 educational training programs for cancer patients in African American and Latino underserved communities. Impact assessments distributed to 100 participants demonstrated positive change in social behavior and willingness to be vaccinated. Post lecture reviews, quizzes, and feedback surveys were distributed to 100 participants. The information received showed a notable change in participants' overall knowledge of COVID-19 regarding the increased risk in cancer patients.

**Conclusions** The data exhibit educational training programs in underserved communities that were hardest hit by COVID-19 increase the understanding of COVID-19 cancer knowledge. The educational training program indicated an association with a greater increase in willingness to participate in COVID-19 prevention practices and willingness to be vaccinated. This research demonstrated the positive
change from educational training programs to be utilized to make a significant impact on health outcomes and cancer mortality rates.

**#106 PREDICTIVE VALUE OF THE MINUTE WALK TEST IN ONCOLOGY AND PRE-REHABILITATION: A LITERATURE REVIEW**

1RC Ellis*, 2A Blough, 1M Clark. 1Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA; 2University of Kansas Medical Center, Kansas City, KS

Purpose of Study To review the predictive value of the minute walk physical function test in hematologic malignancy.

Methods Used A literature review of PubMed using the terms and synonyms of ‘hematologic cancers’ and ‘functional evaluation’ on June 3, 2021 elicited 1,256 manuscripts. After reviewing each abstract for clinical outcomes in relation to minute walk physical function tests in hematologic malignancy, and with the exclusion criteria of confounding intervention or lack of original research, we included 3 published studies.

Summary of Results Increased frailty before and during cancer treatments has been demonstrated to predict mortality, disability, and hospitalization for cancer patients. The minute walk test is an objective measurement of frailty that measures the distance walked in a set amount of time, with decreased distance walked indicative of increased frailty. While there are multiple published manuscripts documenting the association between the minute walk test and clinical outcomes in cancer patients, few studies validate this test in hematologic cancer patients. Our review found 3 studies using the minute walk test as a functional correlate for rates of mortality. Only 1 of the reviewed manuscripts reported significant increase in mortality with decreased physical function measured by the minute walk test, while the other 2 studies showed no significant change. The study that showed a significant change used a follow up period of 2 months, while the studies with nonsignificant results used a 1 to 2 year follow up.

Conclusions The frequency of nonsignificant results and the shorter follow up period of the significant results suggest that the minute walk test may be an unreliable predictor of mortality in hematologic cancers. This affects oncology and physiatry alike. The minute walk is one of multiple frailty assessments that oncologists use to determine the intensity and type of treatment a patient should receive. This would also impact physiatry, as there is a growing practice of ‘pre-rehabilitation’, of improving physical function before and during cancer treatment to improve clinical outcomes. If the minute walk test is an inaccurate predictor of mortality, then pre-rehabilitation may focus less on walking mobility. It is possible that the minute walk test may be an accurate predictor of other outcomes in this patient population, such as patient satisfaction or unplanned hospitalizations. Further research, including a meta-analysis, is necessary to determine the predictive value of the minute walking test in hematologic malignancies. As more rehabilitation and oncology practices embrace pre-rehabilitation, the need for validated and standardized methods of objectively assessing physical mobility increases.

**#107 PREVENTING GROWTH FAILURE AND CHARACTERIZING MICROBIOME IN NEONATES WITH GASTROSCHISIS**

1K Strobel*, 2K Kramer, 3E Fernandez, 4C Rottkamp, 2C Ug, 2L Moyer, 2SL Liebel, 2M Adam, 4P Poulan, 1KL Calkins. 1University of California Los Angeles, Los Angeles, CA; 2University of California San Francisco, San Francisco, CA; 3University of California San Diego, La Jolla, CA; 4University of California Davis, Davis, CA; 5University of California Irvine, Irvine, CA

Purpose of Study 55% percent of infants with gastroschisis in the University of California Fetal Consortium (UCFC) have growth failure (GF). The etiology of GF is multifactorial and associated with caloric/nutrient deficiencies. Intestinal dysbiosis may play a role. In this prospective study of infants with gastroschisis, we aimed to investigate 1) if a nutritional pathway would decrease GF, and 2) the relationship between the microbiome and GF.

Methods Used The UCFC implemented a pathway to decrease GF by standardizing parenteral nutrition dosing, human milk feedings, and GF detection and treatment. Adherence was monitored, and a contemporary cohort (n=45) was compared to a historical cohort (2015–2019, n=125). GF was defined as a decline in weight or length z-score ≥0.8. Shotgun next generation sequencing of the fecal microbiome was performed in a subset of gastroschisis (n=7) and late preterm infants (n=7).

Summary of Results Good adherence to the pathway was noted. Demographics were similar for the cohorts except...
ARACHIDONIC AND DOCSAHEXAENOIC ACID AND RETINOPATHY OF PREMATURITY
1T Gillespie, 2E Kim, 3H Kim, 4T Tsui, 5A Chu, 6KL Calkins. 1University of California Los Angeles David Geffen School of Medicine, Los Angeles, CA; 2University of California Los Angeles, Los Angeles, CA

Purpose of Study Worldwide, 20,000 infants each year are legally blind from retinopathy of prematurity (ROP). We have demonstrated that premature infants develop docosahexaenoic (DHA) and arachidonic acid (ARA) deficits after birth. These polyunsaturated fatty acids play an important role in regulating inflammation and angiogenesis. The purpose of this research is to investigate DHA and ARA status in infants at risk for ROP.

Methods Used Inclusion criteria for this single site retrospective study: ≤30 weeks gestational age (GA) or ≤birthweight (BW) <1.5 kg, and ROP screenings until ROP development, complete vascularization, or 42 weeks postnatal age. DHA and ARA in the red blood cell membrane were quantified with gas chromatography-mass spectrometry. DHA, ARA, and ARA:DHA were compared throughout the first month of life, stratified by either severity of (Type 1 ROP, low grade ROP, no ROP) or treatment for ROP.

Summary of Results Table 1 depicts subject demographics. At week 1, ARA was lower in the Type 1 ROP group vs. the no ROP group (17.9±2.2% vs. 20.5±1.7%, p<0.01). At week 2, significant differences were noted in DHA and ARA (figure 1) but not ARA:DHA. No significant differences in DHA, ARA, and ARA:DHA were observed in weeks 3–4.

Conclusions This study demonstrates that premature infants with more severe ROP, either Type 1 or ROP requiring treatment, have lower ARA and DHA levels than infants without ROP. It remains unclear if DHA and ARA supplementation shortly after birth will improve ROP outcomes.

#109 TYPE OF INTRAVENOUS LIPID EMULSION AND CLINICAL OUTCOMES IN INFANTS WITH GASTROINTESTINAL DISEASE
1LJ Lee*, E Kim, T Romero, KL Calkins. University of California Los Angeles David Geffen School of Medicine, Los Angeles, CA

Purpose of Study Intravenous lipid emulsions (ILEs) are an important component of parenteral nutrition (PN) for neonates with gastrointestinal disorders (GD). Neutones with GD are at high risk for parenteral nutrition associated cholestasis (PNAC) and associated complications, including liver failure. 100% soybean oil (SO) contains a high concentration of hepatotoxic phytosterols and omega-6 fatty acids, which contribute to PNAC. A composite oil (CO) containing 15% fish oil has high amounts of a-tocopherol and omega-3 fatty acids, and less phytosterols. This study aims to compare PNAC and clinical outcomes in infants with GD who received SO or CO.

Methods Used Inclusion criteria for this observational study included: 1) born between 2014 and 2019, 2) GD
Abstract #109 Figure 1

(gastroschisis, omphalocele, intestinal atresia, motility disorder, volvulus, necrotizing enterocolitis, or intestinal perforation), 3) exposure to SO or CO >7 days, and 4) survival to discharge. The primary outcome was cholestasis (conjugated bilirubin (CB) >1 mg/dL). Gas chromatography/mass spectrometry was used to measure fatty acids in the red blood cell membrane in a subset of infants.

Summary of Results The mean ±SD gestational age was 37 ±3 and 36±3 weeks for the SO (n=29) and CO (n=21) groups, respectively (p=0.47). The two groups were well matched for GD diagnosis (p=0.5) and number of GI surgeries (1.8±0.8 for both groups, p=0.90). Nutrition delivery was similar for the SO and CO groups, including days to full enteral feeds (33±12 vs. 30±25 days, p=0.85) and ILE days (25±21 vs. 30±27 days, p=0.77). Weight %SD from birth to discharge (-1.0±0.9 vs. -0.8±1.0, p=0.01 for both), but there was no difference between groups (p=0.52). There was no difference in PNAC incidence (48% vs 48%, p=0.99) and maximum CB (2.0±1.8 vs. 1.9±1.6 mg/dL, p=0.79) when the SO group was compared to the CO group (figure 1). Fatty acid profiles were similar between the two groups.

Conclusions In this study of infants with GD, when compared to infants who received SO, infants who received CO had similar fatty acid trajectories, growth, and clinical outcomes, including PNAC. Further investigation is needed to determine the optimal ILE to decrease PNAC incidence in this population.

#110 NEONATAL ENCEPHALOPATHY FOLLOWING SELECTIVE SEROTONIN REUPTAKE INHIBITOR EXPOSURE IN THE THIRD TRIMESTER OF PREGNANCY: A POPULATION-BASED STUDY

1M Koren*, 2H Forquer, 1A Scheffler, 1A Yeaton-Massey, 1T Newman, 2M Kuznieczuk, 2Y Wu. 1University of California San Francisco, San Francisco, CA; 2Kaiser Permanente, Oakland, CA

Purpose of Study About 4–8% of pregnant women are treated with selective serotonin reuptake inhibitors (SSRI). SSRI exposure in the third trimester may cause poor neonatal adaptation and abnormal movement in neonates, both potential signs of encephalopathy. We assessed whether exposure to SSRI during the third trimester of pregnancy, and dose of SSRI, are associated with neonatal encephalopathy (NE).

Methods Used In a cohort study comprising all Kaiser Permanente Northern California births ≥35 weeks from 2011 to 2019, we defined NE as 5-minute APGAR score <7 and abnormal level of consciousness, activity, tone, or reflexes. We used logistic regression to adjust for potential confounders.

Summary of Results Of 305,426 infants, 8,024 (2.6%) were exposed to SSRI in the third trimester, and 510 (0.17%) had NE. After adjusting for maternal depression or anxiety, maternal age, race, and hospital, exposed neonates had 2.7 times higher odds of NE (95% CI 1.9–3.8). The average risk difference between SSRI-exposed and unexposed mothers was 2.7/1000 (95%CI 1.8–4.1/1000). This relationship was dose-dependent. Each 25mg/d increase in the sertraline equivalent dose was associated with a 31% (95% CI: 23–39%) increase in the odds of developing NE.

Conclusions Exposure to SSRI in the third trimester is associated with increased risk of neonatal encephalopathy. Given that NE is rare, if this association is causal the number needed to cause is high (N~370) and should be balanced with the potential maternal and neonatal benefits of treatment. Future directions include EEG and MRI analyses to correlate SSRI exposure with severity of NE and brain abnormalities.

Abstract #110 Table 1 Neurologic outcomes of neonates exposed and unexposed to SSRI

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<th>No SSRI exposure in 3rd trimester</th>
<th>SSRI exposure in 3rd trimester</th>
<th>Risk ratio (95% CI)</th>
<th>Risk difference (95%CI)</th>
<th>NNH (assuming causality)</th>
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<td></td>
<td>(N=297,403)</td>
<td>(N=8,024)</td>
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<tr>
<td>Encephalopathy</td>
<td>1.6/1,000</td>
<td>4.4/1,000</td>
<td>2.7</td>
<td>(1.9–3.8)</td>
<td>2/71,000</td>
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<tr>
<td>Metabolic acidosis (BE&lt;-16)</td>
<td>16/1,000</td>
<td>21/1,000</td>
<td>1.32</td>
<td>NS</td>
<td>NS</td>
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Abstract #111 BREASTFEEDING RATE REMAINS LOW DURING COVID-19 PANDEMIC IN INFANTS WITH NEONATAL OPIOID WITHDRAWAL SYNDROME (NOWS)

A Fisher*, A McDougal, M McLaren, C Fung. University of Utah Health, Salt Lake City, UT

Purpose of Study Breastfeeding is a well-established non-pharmacological way to improve severity of NOWS in infants with in-utero drug exposure. Given the barriers imposed on mothers during their inpatient stay in the COVID-19 pandemic, the purpose of this study was to assess breastfeeding rate among infants with NOWS during that time. This study also
Identifying barriers to mother’s milk feeding in early preterm black infants in an urban neonatal intensive care unit

Purpose of Study Greater use of mother’s milk (MM) is associated with improved outcomes for preterm infants admitted to neonatal intensive care units (NICUs). Healthcare disparities exist in MM provision to preterm infants and further research is needed to identify barriers to providing MM in high-risk populations. Our urban Level IV NICU serves a patient population who are 60% non-Hispanic Black, allowing us to better study this important demographic. The study objective was to evaluate incidence and predictors of provision of MM to early preterm non-Hispanic Black infants in Baltimore, Maryland.

Methods Used We performed a retrospective medical record review of non-Hispanic Black infants (as identified by their mother) born <34 weeks gestational age (GA), between 9/2014 – 12/2020 in an urban Level IV NICU. We performed bivariate analyses comparing: 1) maternal and neonatal characteristics of infants who received MM at any point during hospitalization, 2) feeding status at discharge, and 3) the proportion of infants who received MM compared to maternal milk (MM) during hospitalization and at discharge.

Results The study included 422 preterm non-Hispanic Black infants. Of these, 332 (79%) received some MM during their NICU admission. Maternal factors associated with receiving any MM during admission included higher maternal gravidity (p=0.0011), increased parity of term deliveries (p<0.0001) and mothers with increased number of living children (p<0.0001). Maternal age and medical comorbidities such as pre-eclampsia, chronic hypertension, and diabetes did not have a significant impact on provision of MM. Infants of mothers with bipolar disorder were less likely to receive MM (p=0.0068) while those of mothers with anxiety were more likely to receive MM (p=0.0243). There was no difference in MM provision for those whose mothers had pre-existing depression or who screened positive for postpartum depression. Mothers of infants who did receive MM were significantly more likely to have had documented lactation consultation during admission (74% vs. 20%, p<0.001). Infants who received no MM had higher birth weights (p<0.0001), were born less prematurely (p=0.0002), and were more likely to have a documented breastfed status at discharge (p<0.0001), while those who received MM were less likely to have a documented breastfed status at discharge (p<0.0001).

Conclusions Despite a higher rate of maternal MAT with no change in the substance exposure rates, infants with NOWS during COVID-19 suffered from the loss of benefits of breastfeeding/breastmilk feeding. The provision of maternal milk when medically safe in infants with NOWS is vital to optimizing short- and long-term outcomes. However, in this population of vulnerable mother-infant dyads, establishing and sustaining breastfeeding remains a complex challenge particularly during the COVID-19 pandemic when additional psychosocial factors and unanticipated barriers may dominate.

#112

Identifying barriers to mother’s milk feeding in early preterm black infants in an urban neonatal intensive care unit

1University of Southern California, Los Angeles, CA
2University of Maryland School of Medicine, Baltimore, MD

10.1136/jim-2022-WRMC.111

Abstract #111 Table 1

<table>
<thead>
<tr>
<th>Infant Demographics</th>
<th>n (%) or median (Q1, Q3)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gestational age at delivery (weeks)</td>
<td>38 (37, 39)</td>
</tr>
<tr>
<td>Female sex</td>
<td>41 (40.2)</td>
</tr>
<tr>
<td>Male sex</td>
<td>61 (59.8)</td>
</tr>
<tr>
<td>Birth weight (grams)</td>
<td>2995 (2724, 3240)</td>
</tr>
<tr>
<td>SGA (BW ≤ 10%)</td>
<td>10 (9.8)</td>
</tr>
<tr>
<td>Inborn</td>
<td>99 (97)</td>
</tr>
<tr>
<td>Infants discharged to non-maternal guardianship</td>
<td>18 (17.6)</td>
</tr>
<tr>
<td>Maternal Demographics</td>
<td>n (%) or median (Q1, Q3)</td>
</tr>
<tr>
<td>Maternal age (years)</td>
<td>31 (28, 34)</td>
</tr>
<tr>
<td>COVID-19 tested</td>
<td>54 (52.9)</td>
</tr>
<tr>
<td>COVID-19 positive</td>
<td>1 (1)</td>
</tr>
<tr>
<td>Caucasian/White</td>
<td>69 (67.6)</td>
</tr>
<tr>
<td>African American/Black</td>
<td>3 (2.9)</td>
</tr>
<tr>
<td>American Indian/Alaskan Native</td>
<td>2 (2)</td>
</tr>
<tr>
<td>Other/Unknown</td>
<td>26 (25.5)</td>
</tr>
<tr>
<td>Primigravida</td>
<td>10 (9.8)</td>
</tr>
<tr>
<td>Mothers receiving medication-assisted treatment</td>
<td>79 (77.5)</td>
</tr>
<tr>
<td>Mode of Delivery</td>
<td>n (%)</td>
</tr>
<tr>
<td>Vaginal</td>
<td>65 (63.7)</td>
</tr>
<tr>
<td>Cesarean section</td>
<td>37 (36.3)</td>
</tr>
<tr>
<td>Breastfeeding</td>
<td>n (%)</td>
</tr>
<tr>
<td>Maternal milk eligible</td>
<td>65 (63.7)</td>
</tr>
<tr>
<td>Any maternal milk provided</td>
<td>57 (55.9)</td>
</tr>
<tr>
<td>Breastfeeding attempted</td>
<td>46 (45.1)</td>
</tr>
<tr>
<td>Breastfeeding at discharge</td>
<td>37 (36.3)</td>
</tr>
<tr>
<td>Prenatal Exposures</td>
<td>n (%)</td>
</tr>
<tr>
<td>Opioids alone</td>
<td>21 (20.6)</td>
</tr>
<tr>
<td>Polysubstance (opioids + ≥ 1 nonopioid)</td>
<td>81 (79.4)</td>
</tr>
</tbody>
</table>
were more likely to have been on a ventilator (p=0.0219) during their admission, though there was no difference in rates of intraventricular hemorrhages.

**Conclusions** Identifying barriers to MM provision for non-Hispanic Black infants will allow clinicians to focus supportive and educational interventions. Interestingly, although medical comorbidities such as diabetes, hypertension, and depression did not lower likelihood of providing MM, having more living children did decrease incidence of MM provision. Inpatient lactation consultation had one of the strongest associations, so enhancing access to lactation consultation may significantly increase MM provision in early preterm neonates.

### #113 IMPROVING ENTERAL FEEDING PRACTICES AND HUMAN MILK CONSUMPTION IN MULTIPLE NICUS: A QUALITY IMPROVEMENT PROJECT

1. Parker *, 2 Ulm, 2 Kaproet, 2 Wieweck, 2 Wetherbee, 2 W Ruben, 2 O Kudin, 2 Weber, 2 M Ellsworth, 2 K Allred, 2 K Marshall, 2 N Whittenberg, 2 M Eklund, 2 G Martin, 2 S Bhopal. 1 The University of Arizona College of Medicine Phoenix, Phoenix, AZ, 2 Division of Neonatology, Phoenix Children’s Hospital, Phoenix, AZ

10.1136/jim-2022-WRMC.112

**Purpose of Study** Optimal nutrition is essential to overcome common disease processes in preterm and high-risk term newborns; however, introduction of enteral feedings creates a possible risk of developing necrotizing enterocolitis (NEC). NEC is a potentially devastating inflammatory disease of the gastrointestinal tract, which can result in intestinal perforation and possibly death. Simple interventions such as prioritizing human milk over formula feeds and following a standardized feeding protocol for initiating and advancing feeds are well established practices for improving outcomes and reducing NEC. Phoenix Children’s Hospital Division of Neonatology was established in 2020, providing medical services to a level 4 NICU and to two level 2 NICUS. We introduced a standardized feeding protocol and used quality improvement methodology to measure compliance with our non-surgical infants <37 weeks, with a goal to increase compliance by >10% and measured human milk use during hospitalization and at discharge in infants of all gestational ages with a goal to increase human milk consumption by >10%.

**Methods Used** We collected data on our feeding practices at all 3 NICUS to measure compliance with the feeding protocol from December 2020 to July 2021. Outcomes were compared in 2 distinct epochs: Epoch 1 from December 2020 to March 2021 and Epoch 2 from April to July 2021. To increase compliance with our feeding protocol, we educated providers, and nurses about the protocol upon its roll-out. Awareness was increased by posting copies at medical provider work stations (February 2021), reviewing interim compliance data with the medical team (April 2021) and placing copies in bedside charts (May 2021).

**Summary of Results** Feeding data was tracked on 265 infants. The mean gestational age and birth weight were 36 weeks (± 3 weeks) and 2700g (± 100g). In both epochs, breast milk was used for the initial feed in 58% of all babies admitted to the NICU. The mean time to full feeds was 4 days ± 2 days in preterm infants <34 weeks. Compliance with protocol improved with time from 72% in epoch 1 to 77% in epoch 2 in babies <37 weeks, and from 66% to 75% in babies <34 weeks. Babies discharging home exclusively on breast milk increased from 16.9% in epoch 1 to 43.5% in epoch 2. There was 1 case of medical NEC in both epochs and no cases of surgical NEC.

**Conclusions** In this quality improvement project, we improved compliance with a feeding protocol and increased exclusive human milk usage through hospital discharge. While there was 1 case of medical NEC in both epochs, there were no cases of surgical NEC in our data set.

<table>
<thead>
<tr>
<th>#114 CLINICAL AND ECONOMIC IMPACT OF USING EXCLUSIVE HUMAN MILK IN VERY LOW BIRTH WEIGHT INFANTS</th>
</tr>
</thead>
</table>

M Tetarbe *, M Chang, L Barton, E Toushin, R Ramanathan, R Cayabyab. University of Southern California, Los Angeles, CA

10.1136/jim-2022-WRMC.113

**Purpose of Study** The AAP recommends use of expressed breast milk (EBM) or donor human milk (DHM) in preterm infants fortified with proteins, minerals, and vitamins to ensure optimum nutrient intake. Unfortunately, the implementation of EBM/DHM fortified with human milk-based fortifiers (EHM) can place an economic burden on individual institutions raising concerns on the economic feasibility of such products. The objective of this study is to assess the clinical impact of using EHM in very low birth weight infants (VLBW) infants, and to perform a cost-benefit analysis of its use.

**Methods Used** Retrospective study of all VLBW infants admitted to neonatal intensive unit before and after the implementation on the use of EHM. Neonatal demographics and clinical outcomes such as necrotizing enterocolitis (NEC), severe retinopathy of prematurity (ROP), bronchopulmonary dysplasia (BPD), late-onset sepsis (LOS), and average length-of-stay (ALOS) were collected from January – December 2016 (before implementation) and January – December 2020 (after implementation) using the Pediatric Data Collection System (PDACS) database. We used EHM for all infants in NICUs and EBM and DHM for infants less than 1500g.

**Abstract #114 Table 1**

<table>
<thead>
<tr>
<th>Clinical Outcome</th>
<th>Pre-EHM 2016 N=45</th>
<th>Post-EHM 2020 N=27</th>
<th>Change Post vs Pre</th>
<th>Estimated cost per case</th>
<th>Cost Avoidance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Medical necrotizing enterocolitis, n</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>$74,004</td>
<td>$74,004</td>
</tr>
<tr>
<td>Late-onset sepsis, n</td>
<td>4</td>
<td>5</td>
<td>-1</td>
<td>$10,055</td>
<td>$10,055</td>
</tr>
<tr>
<td>Bronchopulmonary dysplasia, n</td>
<td>9</td>
<td>5</td>
<td>4</td>
<td>$31,565</td>
<td>$126,260</td>
</tr>
<tr>
<td>Severe retinopathy of prematurity, n</td>
<td>2</td>
<td>3</td>
<td>-1</td>
<td>$35,749</td>
<td>$35,749</td>
</tr>
<tr>
<td>Total parenteral nutrition, days*</td>
<td>34</td>
<td>27</td>
<td>7</td>
<td>$1,436</td>
<td>$271,404</td>
</tr>
<tr>
<td>Length-of-stay, days*</td>
<td>87.2</td>
<td>74.3</td>
<td>12.9</td>
<td>$3,500</td>
<td>$1,219,050</td>
</tr>
<tr>
<td>Cost Avoidance</td>
<td>$1,644,914</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Product Acquisition</td>
<td>$313,784</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2020</td>
<td>$1,331,130</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*mean
Summary of Results After excluding deceased infants in both time periods, 45 infants were included in the pre-EHM analysis period, (mean birth weight (BW): 1034 g, mean gestational age (GA): 27.9 weeks), and 27 infants were included in the post-EHM analysis period (mean BW: 1070 g, mean GA: 28.8 weeks). Our institution’s product acquisition cost in 2020 was estimated to be $313,784. The implementation of the EHM protocol saw a reduction in the ALOS by 12.9 days and average total parenteral nutrition (TPN) use by 7 days per infant in the post-EHM group equating to a net savings of $1,176,670. While there was a small difference in the number of morbidities between the two time periods, when combining the cost avoidance to include medical NEC, and BPD, the estimated financial impact excluding insurance reimbursement rose to $1,331,130 (table 1).

Conclusions Our preliminary findings suggest that implementation of exclusive human-milk feeding in VLBW infants is a cost-effective option for NICUs that can result in decrease in NEC, BPD, TPN use and length of stay for these infants at nominal cost.

### Neonatology pulmonary II

#### Concurrent session

3:15 PM Thursday, January 20, 2022

**#115** **PERINATAL ELECTRONIC CIGARETTE-EXPOSURE INDUCES ASTHMA IN RAT OFFSPRING**

A Harb*, C Yu, J Liu, R Sakurai, Y Wang, V Rehan. The Lundquist Institute, Torrance, CA

**Purpose of Study** An exponential increase in the use of electronic cigarettes (e-cig), including by pregnant women, exposes an increasing number of fetuses to potentially harmful e-cig chemicals with little knowledge of its repercussions. Perinatal nicotine exposure-induced asthma is associated with downregulated PPARγ signaling and upregulated Wnt signaling in the developing lung. However, the impact of maternal nicotine vaping on the developing lung is unknown. Here, we use an established rat model to determine the effect of perinatal maternal e-cig vaping on offspring pulmonary function and markers of airway contractility.

**Methods Used** Pair-fed pregnant rat dams received saline, vehicle (e-cig without nicotine), or e-cig with nicotine daily from embryonic day 6 until postnatal day (PND) 21. Using an established e-cig delivery system and mimicking real-life puffing topography, dams were exposed to four-sec puffs, one puff (puff volume 35 ml) every 30s, 3h/day, and 7 days/week. Average maternal plasma nicotine level (7 ± 4 ng/ml) using this vaping regimen is well within the range observed in moderate cig smokers. Pups delivered spontaneously at term and breastfed ad-lib, but not directly exposed to e-cig aerosols at any time. At PND21, lung resistance and compliance were determined following the methacholine challenge. At sacrifice, the lungs were collected to determine the expression of airway contractility markers, i.e., α-SMA, Calponin, Fibronectin, Collagen I/III, and key Wnt/PPARγ signaling intermediates by qRT-PCR, immunoblotting, and immunostaining.

**Summary of Results** Compared to controls, perinatal e-cig exposure resulted in a significant increase in airway resistance and decreased airway compliance following the methacholine challenge. mRNA levels of Collagen III and LIF-1 increased, and those of PPARγ and ADRP decreased in the e-cig group. Immunoblotting showed that in the e-cig group, airway contractility markers (α-SMA, Calponin, Fibronectin, Collagen I and III), Wnt signaling intermediates (β-catenin and LIF-1), and nicotinic acetylcholine receptors α3 and α7 levels increased. In contrast, compared to controls, PPARγ, which interacts directly with Wnt signaling intermediates, levels decreased. Immunostaining of whole lung sections confirmed immunoblotting data.

**Conclusions** For the first time, we unequivocally demonstrate offspring asthma following perinatal maternal e-cig vaping and explain likely molecular mechanisms involved. Our data add to the accumulating evidence contradicting the idea that e-cigs are safe.

**Grant Support** NIH (HL151769, HD127237, HD071731, and HL152915)) and TRDRP (23RT-0018, 27IP-0050, and T29IR0737).

**#116** **COMBINED PRE- AND POSTNATAL GROWTH RESTRICTION INCREASE EXPRESSION OF A NOVEL DOMINANT NEGATIVE PPARG SPICE VARIANT IN THE RAT LUNG**


10.1136/jim-2022-WRMC.115

**Purpose of Study** Preterm infants frequently suffer growth restriction, increasing the risk and severity of neonatal lung disease, characterized by impaired alveolar development and worse outcomes in male infants. We showed that growth restriction in the prenatal (IUGR) or postnatal (PGR) period reduces rat lung PPARγ gene expression, which results in impaired alveolar development. PPARγ variants, including the novel delta 5 splice variant (PPARγδ5), can impact the downstream effects of PPARγ activation. As PPARγδ5 is a dominant negative variant, the effect of increasing PPARγδ5 is a reduction in PPARγ signaling. Whether PPARγδ5 is expressed in the rat lung, and the effect by growth restriction on expression is unknown. We hypothesize that PPARγδ5 will be expressed in rat lung, and that the combination of IUGR and PGR will increase expression of PPARγδ5.

**Methods Used** IUGR and PGR were generated in Sprague Dawley rat pups by bilateral uterine artery ligation and variation in litter size respectively. Lungs were collected at postnatal day 12 from Control, IUGR only, PGR only, and PGR +IUGR rat pups. Male and female rats were treated as separate groups. PCR, gel electrophoresis, and sequencing were used to confirm the presence of PPARγδ5 in the rat lung. Full length and PPARγδ5 mRNA and protein were assessed using real-time RT PCR and western blotting. Differences were assessed by one-way ANOVA and fishers post hoc test.

**Summary of Results** Results are IUGR as% of control±SD, *P<.05. Sequence confirmed PPARγδ5 mRNA is expressed in...
rattung at postnatal day 12. PGR model resulted in significantly lower weights on D12 (66.9±3% for PGR only, 64.8±3.2% in IUGR+PGR). In male rat lung, PPARγΔ5 mRNA was increased (325±79%*) by IUGR+PGR. Similarly, in the male rat lung, PPARγΔ5 protein was increased by IUGR+PGR (163±14%). In female rat lung, neither PPARγ transcript was affected. However, lung protein levels of PPARγΔ5 were increased in female IUGR+PGR (146±24%*).

**Conclusions** We conclude that the PPARγΔ5 is expressed in rat lung, and that IUGR+PGR increase expression. We speculate that increased PPARγΔ5 expression in male IUGR+PGR rat lungs may further impair PPARγΔ5 signaling, leading to impaired alveolar development.

### EVIDENCE FOR THE INTERGENERATIONAL PULMONARY EXPRESSION OF GENES DIFFERENTIALLY METHYLATED IN SPERM CELLS OF PERINATALLY NICOTINE EXPOSED RATS

L Afrose*, Y Wang, J Liu, C Yu, T Rho, T Dao, D Hatai, V Rehan. The Lundquist Institute, Torrance, CA

10.1136/jim-2022-WRMC.116

**Purpose of Study** Nicotine exposure to the developing fetus results in asthma that can be transmitted across generations. However, the underlying mechanism remains unknown. We recently demonstrated differential DNA methylation in the proximity of nicotine-response genes in sperms of the perinatally nicotine exposed F1 animals. Gene ontology and pathway enrichment analysis suggested a possible link between the spermatozoidal differential DNA methylation and the offspring asthma phenotype. We hypothesize that nicotine-induced spermatozoidal epigenetic changes drive the intergenerational transmission of nicotine-induced asthma. The expression of genes in F2 lungs differentially methylated in the spermatoozoa of nicotine exposed F1 males was determined to test this hypothesis.

**Methods** Used Sprague Dawley rat dams (F0) received nicotine (1 mg/kg, sc) or saline from embryonic day 6 (E6) until postnatal day 21 (PND21). Pups (F1) were weaned at PND21 and used as breeders to generate F2 without any subsequent exposure to nicotine in the F1 progeny. F2 pups were weaned at PND21. At PND60, F2 males (n=20; 10 control, 10 nicotine) were sacrificed, and their lungs were collected and flash frozen for performing qRT-PCR for the top 11 differentially methylated genes (AABR07051515.1, Dio1, Gabra4, Htr6, Map4k2, Men1, Mntu, Orai2, Rars, Sec1415, and Scl7a11) in sperm cells of the nicotine exposed F1 males.

**Summary of Results** In line with data from F1 lungs, the expression of the top 2 differentially hypermethylated genes AABR07051515.1, a lincRNA, known to modulate lung function, and Dio1 (iodothyronine deiodinase 1) in the nicotine exposed F1 sperm cells was upregulated or downregulated, respectively (p<0.05), in F2 lungs. In addition, similar to the F1 progeny, Mntu and Sec1415 genes' expression was downregulated (p<0.05) in F2 lungs of the nicotine-exposed group. In contrast, the expression of the other 7 differentially methylated genes in F1 spermatoozoa did not change significantly.

**Conclusions** Our data further support the concept that perinatal nicotine exposure-induced spermatoozoal epigenetic reprogramming, specifically DNA methylation alterations in nicotine response- and lung development-related genes, likely drive the intergenerational transmission of perinatal nicotine-induced asthma.

**Grant Support** NIH (HL151769, HD127237, HD071731, and HL152915) and TRDRP (23RT-0018; 27IP-0050; and T29IR0737).

### THE EFFECT OF SEROTONIN DEPLETION ON HYPOXIA INDUCED NEONATAL MURINE BRONCHOPULMONARY DYSPLASIA AND PULMONARY HYPERTENSION

D Roberts*, JN Posey, J Archambault, E Nazik, C Delaney. University of Colorado – Anschutz Medical Campus, Aurora, CO

10.1136/jim-2022-WRMC.118

**Purpose of Study** Pulmonary hypertension (PH) associated with bronchopulmonary dysplasia (BPD) leads to worse outcomes in former preterm neonates. Serotonin (5-hydroxytryptamine, 5-HT) is a potent pulmonary vasoconstrictor, smooth muscle mitogen, and is increased in the lungs of infants who died with severe BDP. Tryptophan hydroxylase 1 (TPH1), the rate limiting enzyme in 5-HT synthesis, is increased in adult patients and animals with experimental PH. Serotonin signaling blockade decreases pulmonary vascular resistance and prevents pulmonary vascular remodeling in preclinical models. We hypothesized that TPH1 knock-out (KO) neonatal mice would be protected from hypoxia induced BPD associated with PH.

**Methods** Used Neonatal wild-type (WT) and TPH1 KO offspring were placed in hypoxia or remained in normoxia at Denver altitude for 2 weeks. To assess alveolar development, inflation fixed lungs were analyzed for surface area (SA) and mean linear intercept (MLI). To identify total number of small vessels (<30 μm), lung sections were immunostained with Factor VIII. PH was assessed by Fulton’s index and right ventricular systolic pressures (RVSP). Platelet poor plasma (PPP), platelet, and lung homogenate 5-HT levels were measured by ELISA. Data were analyzed by Prism with unpaired t-test or 2-way ANOVA with Bonferroni post-hoc analysis. Significance level p<0.05.

**Summary of Results** At baseline, WT mice have more platelet poor plasma, platelet, and lung 5-HT than KO mice (53±6 — 91±1, p<0.0001; 275±11 — 56±12, p<0.0001; 18±2 — 10±2, p<0.004; respectively, ng/mL). TPH1 KO mice were not protected from hypoxia-induced alveolar simplification, shown by no difference compared to WT mice MLI and SA, nor were they protected against hypoxia-induced pulmonary vascular simplification, shown by no difference compared to WT mice vessel density. TPH1 KO mice were attenuated to hypoxia-induced pulmonary vasoconstriction, shown by reduction in RVSP (32±0.66 — 29±0.55, p<0.006, mmHg). There was less PPP and platelet 5-HT in hypoxia-exposed WT mice compared to WT mice at baseline (20±2 — 53±6, p<0.0001 and 117±27 — 275±11, p<0.0001, respectively, ng/mL). There was less lung 5-HT in hypoxia-exposed KO mice than in KO mice at baseline (2±1 — 10±1, p<0.001, ng/mL).

**Conclusions** Neonatal TPH1 KO mice are not protected against hypoxia-induced lung injury. Surprisingly, this study contradicts the current understanding of the role of 5-HT in adults with PH and in adult models of hypoxia-induced PH. We found decreased plasma and platelet 5-HT following hypoxia exposure. We speculate that decreased 5-HT observed in hypoxia may contribute to neonatal hypoxia-induced...
alveolar simplification and impaired vascular development. Further studies are needed to elucidate the role of 5-HT in the developing lung.

**THE EFFECT OF PLATELET ALPHA GRANULE DEFICIENCY ON HYPOXIA INDUCED NEONATAL PULMONARY HYPERTENSION**

D Roberts*, JN Posey, ENozi, C Delaney. University of Colorado – Anschutz Medical Campus, Aurora, CO

Purpose of Study Pulmonary hypertension (PH) associated with bronchopulmonary dysplasia (BPD) leads to worse outcomes in former preterm neonates. Elevated platelets at birth are an independent predictor of BPD, increased platelet derived protein after birth is associated with higher rates of neonatal pulmonary vascular disease, and perinatal platelet transfusions are associated with higher rates of mortality and BPD. Circulating platelets from neonatal mice with experimental PH are increased and express a higher percentage of active αIIbβ3, a marker of platelet activation. NBEAL2 knock-out (KO) mice lack platelet alpha granules, have low platelet counts, and have decreased platelet function in vitro and in vivo. We hypothesized that NBEAL2 KO neonatal mice would be protected from hypoxia-induced PH.

Methods Used Neonatal wild-type (WT) and NBEAL2 KO offspring were placed in hypobaric hypoxia (18,000 feet) or normoxia at Denver altitude for 2 weeks. Hypoxia was assessed by Fulton’s index as a marker of right ventricular hypertrophy (RVH) and right ventricular systolic pressures (RVSP). Data were analyzed by Prism with unpaired t-test or 2-way ANOVA with Bonferroni post-hoc analysis. Significance level p<0.05.

Summary of Results Right ventricular systolic pressure is higher in NBEAL2 KO mice than in WT mice at baseline (24.2±0.5 vs 21.5±0.3, p<0.001, mmHg). There is no difference between baseline right ventricular hypertrophy between NBEAL2 KO mice and WT mice (0.27±0.01 vs 0.31±0.01, ns). NBEAL2 KO mice display comparable hypoxia-induced increase in RVSP compared to WT mice (29±1 vs 30±1, ns, mmHg). NBEAL2 KO mice display comparable hypoxia-induced increase in RVH compared to WT mice (0.3±0.03 vs 0.4±0.03, ns).

Conclusions Platelet alpha granule deficiency is a risk factor for neonatal pulmonary vasoconstriction at baseline. Further studies are needed to elucidate the role of platelets in neonatal PH associated with BPD.

**METABOLITES IMPORTANT IN THE DIFFERENTIATION OF HUMAN PLURIPOTENT STEM CELLS TO LUNG PROGENITOR CELLS**

SL Leibel*, 2I Tseu, 3A Zhou, 4A Hodges, 5Y Yin, 2C Bilodeau, 7O Goltsis, 1M Post. 1University of California San Diego, La Jolla, CA; 2SickKids Research Institute, Toronto, ON, Canada; 3Johns Hopkins University, Baltimore, MD; 4Sanford Burnham Prebys Medical Discovery Institute, La Jolla, CA

Purpose of Study Metabolism is vital to cellular function and tissue homeostasis during human lung development. In utero, embryonic stem cells undergo endodermal differentiation towards a lung progenitor cell (LPC) fate that can be modeled in vitro using pluripotent stem cells (hPSCs). We previously showed differences in lung cell composition and gene expression between wild type and surfactant protein B (SP-B) deficient lung organoids. These differences may be impacted by changes in metabolites during early lung development. We hypothesize that SP-B deficient cells will express a different metabolomic profile compared to wt cells during the differentiation to lung progenitor cells.

Methods Used To examine metabolites that differ during endodermal differentiation, we used an untargeted metabolomics approach to evaluate the changes in metabolites at the stem cell (hPSC), definitive endoderm (DE), anterior foregut endoderm (AFE) and lung progenitor (LPC) stage between wt and SP-B deficient cell lines. At each differentiation step, the cells were sorted for surface markers specific to their differentiation stage in quadruplicate. The homogeneous cell lysates were analyzed using a Biocrates p180 metabolite kit including hexoses, amino acids, phosphatidylcholines, lysophosphatidylcholines, sphingolipids, acylcarnitines, and biogenic amines. The metabolomic multivariate data analysis was performed using XLSTAT2016 software (Addinsoft) and MetaboAnalyst.

Summary of Results We found that the largest metabolic changes during endodermal differentiation occurred from hPSC to DE with a change from glycolytic respiration to oxidative phosphorylation. The metabolites most enriched during the differentiation from hPSC to LPC, independent of cell line, were sphingomyelin and lecithin. In the wt cell lines, metabolites for oxidation of fatty acids and tryptophan metabolism were up-regulated, while metabolites for ammonia recycling and aspartate metabolism were down-regulated. In the SP-B deficient cells, metabolites in fatty acid oxidation and carnitine synthesis were up-regulated and metabolites for amino acid metabolism, the urea cycle, and multiple energy-based pathways were down-regulated.

Conclusions Differentiation to lung progenitor cells from pluripotent stem cells resulted in increased fatty acid metabolism and decreased urea cycle and aspartate metabolism in both wt and SP-B deficient cell lines. Therefore, metabolite composition in early lung development is not influenced by the loss of SP-B expression.

**EFFECTS OF VITAMIN A AND VITAMIN D TREATMENT ON LUNG GROWTH AND FUNCTION IN OFFSPRING FROM MATERNAL VITAMIN D DEFICIENT RATS**

N Galambos*, 1E Bye, 1T Gonzalez*, 1G Seedorf, 1G Smith, 1JW Alman, 1E Mandell. 1University of Colorado – Anschutz Medical Campus, Aurora, CO; 2The University of Texas at Austin College of Natural Sciences, Austin, TX

Purpose of Study Maternal vitamin D deficiency (M-VDD) is associated with perinatal pulmonary morbidities. We have demonstrated that offspring of rodent maternal VDD dams have sustained abnormalities of distal lung structure, increased airway hyperreactivity and abnormal lung mechanics. In pulmonary endothelial cells, vitamin A (VA) and vitamin D (VD) co-dimerize on retinoid x receptor. VA therapy has been shown to improve lung development in pre-clinical and clinical studies, but whether combined postnatal (PN) treatment with VA and VD further enhances lung development in offspring of M-VDD dams is unknown. Therefore,
we seek to determine if PN VA and VD supplementation improves lung development and function in offspring of M-VDD dams.

Methods Used Newborn rats from control (CTL) and M-VDD dams received daily treatment of retinoic acid (VA) alone, VA and 1,25-OHD (VD) (VA-VD) or saline (SAL) for 14 days. On DOL 14 lung structure was assessed by mean linear intercept (MLI), radial alveolar count (RAC) and pulmonary vessel density (PVD). Lung mechanics were measured using flexiVent.

Summary of Results Lungs from VDD-SAL rats had increased MLI (p<0.001) and decreased pulmonary vessel density (p<0.05) as compared to CTL-SAL. VDD rats that received VA had increased RAC compared to VDD-SAL (p<0.05). VDD-SAL rats had increased resistance (p<0.01) and decreased compliance (p<0.01) as compared to CTL-SAL. VDD-VA rats had decreased elastance as compared to VDD-SAL pups (p<0.05).

Conclusions M-VDD decreases distal lung and vascular development and impairs lung function in infant rats. PN VA therapy improved RAC and decreased elastance in VDD pups. These findings suggest that abnormal lung development after PN VA therapy may improve alveologenesis and lung mechanics of M-VDD pups. We speculate that M-VDD leads to persistent abnormalities in infant lung growth that may be responsive to PN VA.

#122 MATERNAL VITAMIN D DEFICIENCY ALTERS PULMONARY ENDOTHELIAL CELL GROWTH AND MRNA EXPRESSION IN NEWBORN RATS

1. Gonzalez, 2. E Bye, 1. N Galambos, 3. G Seedorf, 2. C Fleet, 3. SH Abman, 2. E Mandell. 1. University of Colorado – Anschutz Medical Campus, Aurora, CO; 2. The University of Texas at Austin College of Natural Sciences, Austin, TX

Purpose of Study Vitamin D deficiency (VDD) during pregnancy is associated with chronic lung disease in preterm infants, and the underlying mechanisms are not understood. We have shown that vitamin D (VD) preserves lung structure and prevents pulmonary hypertension (PH) in an experimental model of bronchopulmonary dysplasia, and that VD treatment increases pulmonary artery endothelial cell growth and function. However, the direct effects of maternal VD on pulmonary endothelial cell (PEC) growth and function are unknown. Thus, we seek to determine whether PEC from newborn rats exhibit altered growth and mRNA expression at birth after exposure to maternal VD and whether these changes persist during infancy.

Methods Used Female rats were fed VDD chow and shielded from UV-B light to achieve 25-OH levels less than 10 ng/ml before mating. PEC were isolated from offspring of maternal VDD (VDD) or control (CTL) dams at postnatal day 0 and 14. PECs were used for proliferation assays and response to exogenous VEGF and 1,25-OH. PEC lysates were also collected for RT-qPCR analysis.

Summary of Results PEC isolated from VDD pups at both D0 and D14 demonstrate decreased growth compared to CTL D0 and D14 (p<0.01). VEGF or 1,25-OH treatment increased CTL PEC growth from both D0 and D14 when compared to untreated CTL D0 and D14 PEC (p<0.01). In contrast, neither VEGF nor 1,25-OH treatment increased D0 VDD PEC growth. D14 VDD PEC showed an increased growth with VEGF treatment compared to untreated D14 VDD PEC (p<0.01). RNA isolated from D0 VDD PEC demonstrate decreased expression of KDR and eNOS and increased VEGF expression compared to D0 CTL PEC (p<0.01), no expression changes seen at D14.

Conclusions We found that D0 PEC from newborn offspring of maternal VDD dams demonstrate decreased baseline PEC growth and no responsiveness to angiogenic stimuli. At D14 VDD PEC grew poorly at baseline, and were responsive to VEGF but not 1,25-OH treatment. We speculate that maternal VDD disrupts normal PEC function, which persists into postnatal life and may contribute to high risk for late cardiopulmonary disease.

#123 THE EFFECT OF PERINATAL NICOTINE EXPOSURE ON THE LUNG CIRCADIAN MOLECULAR CLOCK

D Hata*, Y Wang, R Sakurai, J Liu, I Afrose, T Dao, T Rho, C Yu, V Rehan. The Lundquist Institute, Torrance, CA

Purpose of Study Dysregulated peripheral circadian rhythm is associated with enhanced inflammatory response and cellular senescence. Recent studies have demonstrated an association of exposure to cigarette smoke and dysregulated peripheral molecular clock in Chronic Obstructive Pulmonary Disease (COPD) and asthma patients. This has also been confirmed in rodent models. Although developmental smoke/nicotine exposure predisposes to asthma and COPD, its impact on circadian clock genes is unknown. Here, we test the hypothesis that developmental nicotine exposure alters the molecular clock, which lasts well into adulthood.

Methods Used Pair-fed pregnant Sprague-Dawley rat dams received once-daily 1mg/kg nicotine or saline dilluent from embryonic day 6 (E6) to postnatal day 21 (PND21). Lungs from pups were collected on E21, PND21, or PND60 and flash-frozen for later mRNA and protein analysis. The expression of core clock genes (Bmal1, Clock, Cry1, Cry2, Per2, Per2, Rev-erba, Rev-erbb, Rora, and Sirt1) was determined by qRT-PCR on mRNA isolated from lungs. Protein levels of key clock genes Bmal1, Clock, and Rev-erba were determined using western analysis on proteins extracted from PND 21 lungs.

Summary of Results Overall, the mRNA expression of Bmal1, Clock, Cry1, Cry2, Per1, Per2, Rev-erba, Rev-erbb, Rora, and Sirt1 was significantly decreased (p<0.05) in the nicotine treated group vs. the control group at E21 and PND21. Perinatal nicotine exposure-induced downregulation of key clock genes Bmal1 and Rev-erba was also confirmed by their down-regulated protein levels by western analysis at PND 21. Interestingly, the expression of several down-regulated clock genes at E21 and PND21 in the nicotine-treated group was either not different or upregulated versus the control group at PND60, suggesting a dynamic response of perinatal nicotine exposure on the peripheral molecular clock.

Conclusions Perinatal nicotine exposure leads to peripheral clock dysregulation in the lung that lasts at least through adolescence. These results suggest a new mechanism that underlies the effects of perinatal nicotine-induced lung injury. Further studies are needed to determine the impact of perinatal nicotine exposure-induced dysregulated peripheral clock on lung health, gender specificity, and how long these effects last.
Purpose of Study Pulmonary hypertension (PH) is a life-threatening condition that affects infants, children, and adults. However, treatment strategies are limited, and morbidity and mortality remain significant. We have previously demonstrated in robust animal models that serotonin (5-HT) contributes to the pathogenesis of experimental neonatal PH and know that infants who died due to severe lung disease have a 34-fold increase in lung 5-HT. We designed an exploratory pilot study to test the hypothesis that systemic 5-HT is increased in infants with persistent pulmonary hypertension of the newborn (PPHN).

Methods Used Near term and term infants (≥36 weeks) were recruited from the NICUs at Children’s Hospital Colorado and University of Colorado Hospital beginning in March 2021. Infants with culture proven sepsis, metabolic/genetic abnormality, major cardiac defect, renal failure, or anentanatal exposure to SSRIs were excluded. PH was defined on echocardiogram by an estimated systolic pulmonary artery pressure ≥40 mm Hg, end-systolic eccentricity index ≥1.16, or presence of a right-to-left shunt. 5-HT is an unstable neurotransmitter that degrades quickly; thus, we measured its more stable metabolite 5-hydroxyindolacetic acid (5-HIAA). Urine samples were collected on DOL 1 and DOL 3, and 5-HIAA was analyzed via mass spectrometry. Monthly follow up samples were collected if PH persisted. Demographics, clinical characteristics, and interventions were obtained through chart review and summarized for the patient cohort. 5-HIAA levels were summarized using medians and ranges.

Summary of Results To date, 6 infants with PPHN and 7 age-matched controls have been enrolled. 54% were male and 46% female. The mean gestational age was 38.2 weeks. All infants with PPHN were classified as having severe PH on initial echo. 83% were born with congenital diaphragmatic hernia (CDH) and 100% had a patent ductus arteriosus (PDA). 83% required vasopressors, with 60% initiated in the delivery room. 100% required steroids for blood pressure and/or respiratory support. 83% required pulmonary vasodilators with inhaled nitric oxide and sedenafil being the most common. 50% of infants with PPHN were followed for refractory PH for a mean of 2.67 months. At DOL 1, the median 5-HIAA level was 14.89 (min, max: 14.17, 19.1) in the PPHN group and 14.17 (11.56, 16.63) in the control group. At DOL 3, the median was 18 (8.89, 40.5) in the PPHN group and 15 (10.95, 17.14) in the control group.

Conclusions This study investigated the association of PPHN with systemic alterations in 5-HIAA. Our current results offer a preliminary description; however, enrollment is ongoing. With additional data we will test our hypothesis that 5-HIAA is significantly associated with PH severity. Ultimately, we aim to establish it as a noninvasive biomarker to follow treatment response and predict the later development of PH in high-risk infants.
Purpose of Study In humans, as well as other vertebrates, color vision requires the differential expression of specific cone opsinps in photoreceptor cone cells. One model for the regulation of the human long and medium wavelength sensitive (LWS/MWS) opsin tandem array suggests an upstream regulatory region interacts with replicated opsin genes at random, resulting in mutually exclusive expression of a specific opsin. A similar ontologous long wavelength sensitive (lus1/lus2) array in zebrafish provides a good model for study of this regulation. However, our prior investigations into this array suggest that thyroid hormone (TH) and retinoic acid serve as trans regulators in larval/juveniles (Mitchell et al., 2015, PLOS Genetics; Mackin et al., 2019, PNAS). This study investigates whether cone opsin expression remains plastic to TH treatment in adult zebrafish, where cone distribution is considered stable.

Methods Used Adult zebrafish (6–18 months old) were treated with NaOH (0.01%, control) or TH (386 nM) for 1 or 5 days. qRT-PCR was performed on homogenized eyes. Whole retinas were treated by hybridization chain reaction in situ and then analyzed by confocal imaging for mRNA expression.

Summary of Results In adult zebrafish, exogenous TH drastically increased lus1 expression in both 1 and 5 day-treated groups (p<1e-7, 0.01, respectively) while decreasing lus2 expression (p<0.001, 0.001). Other phototransduction-related transcripts (gopr2b, rh2–1) also demonstrated expression changes following TH treatment. Exogenous TH induced a drastic shift from lus2 to lus1 in adult zebrafish, consistent with previous studies of larvae and juveniles.

Conclusions This shift from lws2 expression to lws1 expression occurs as rapidly as 1 day when exposed to TH, which shows that cones remain highly plastic even into adulthood. Plasticity in spectral sensitivity (to be sensitive to higher wavelengths) in response to TH suggest a role in visual system function well into adulthood. These results oppose earlier models suggesting that regulation between tandemly replicated opsin genes is stochastic and fixed.

Purpose of Study The head-twitch response (HTR) is evoked following stimulation of postsynaptic serotonin 2A (5-HT2A) receptors in the prefrontal cortex (PFC). D-Fenfluramine (FF) is a selective 5-HT releaser, it produces the HTR via release of serotonin from nerve terminals through the 5-HT uptake carrier working in reverse. Methamphetamine (MA) is a non-selective releaser of monoamines 5-HT, noradrenaline (NE) and dopamine (DA). We investigate whether pretreatment with either MA (1–5 mg/kg, i.p.) or the 5-HT2A receptor selective antagonist EMD 281014 (0.001, 0.005, 0.01, 0.05 mg/kg, i.p.) can alter: 1) the mean frequency of FF-induced HTR at different ages (20-, 30- and 60-day old), and 2) the expression of c-fos evoked by FF in different regions of the PFC. We also explored whether blockade of serotonergic 5-HT1A or adrenergic α2-receptors can alter the effect of MA on FF-induced HTR across the above ages.

Methods Used The HTR was observed for 30 min following the injection of FF in each mouse. We use immunohistochemistry study to evaluate the changes of c-fos expression in the PFC.

Summary of Results Pretreatment with MA (1–5 mg/kg, i.p.) dose-dependently suppressed the FF-induced HTR across different ages. MA at 1 mg/kg in 20- and 30-day old mice, and at 5 mg/kg in 60-day old mice significantly suppressed the FF-induced HTR. Pretreatment with EMD 281014 (0.001, 0.005, 0.01, 0.05 mg/kg, i.p.) also blocked the FF-induced HTR in an age- and dose-dependent manner. The selective 5-HT1A receptor antagonist WAY 106635 (0.25 mg/kg, i.p.) and the adrenergic α2-receptor antagonist RS 759948 (0.1 mg/kg, i.p.) significantly reversed the inhibitory effect of MA on the mean frequency of HTR in 20-day old mice, but not in 30- and 60-day old mice. Moreover, FF significantly increased c-fos expression in several PFC regions in 30-day old mice. Despite the inhibitory effect of MA or EMD 281014 on FF-induced HTR, pretreatment with either MA (1 mg/kg, i.p.) or EMD 281014 (0.05 mg/kg, i.p.) significantly increased c-fos expression in different regions of the PFC in 30-day old mice.

Conclusions The inhibitory effect of MA on the FF-evoked HTR appears to be mainly due to functional interactions between the stimulatory 5-HT2A - and the inhibitory 5-HT1A- and/or adrenergic α2-receptors. The MA-induced increase in c-fos expression in different PFC regions is probably due MA-evoked increases in synaptic concentrations of 5-HT, NE and/or DA. EMD 281014 failed to prevent the increase in c-fos expression induced by FF, which may be due to the increased 5-HT synaptic concentration that activates other serotonergic receptors, such as, 5-HT1A.

Purpose of Study Geographic atrophy (GA) is a severe and poorly understood progression of dry age-related macular degeneration (AMD). Patients with GA are also more likely to develop choroidal neovascularization. Carbipoda-levodopa treatment has demonstrated successful reduction in neovascular AMD. In this study, we investigate the effects of carbipoda-levodopa treatment on progression of GA.

Methods Used A retrospective analysis of patients with already existing GA who participated in our proof-of-concept study was performed. Fundus autofluorescence (FAF) and optical coherence tomography (OCT) were utilized to confirm the presence of geographic atrophy. This study followed the 2018 retina consensus meeting requirements to measure geographic atrophy markers in patients. The primary outcomes measures
were complete retinal pigment epithelium and outer retinal atrophy (cRORA), hypertransmission through Bruch’s membrane, and mm/year change from initiation of study drug.

**Summary of Results** We included 5 patients with already existing geographic atrophy in 6 eyes. This cohort consented to carbidopa-levodopa treatment and was predominantly male (3 patients) with a median (IQR) age of 82.5. The mean GA change in cRORA 1 year before and after treatment initiation was -0.004433 mm/year and 0.0061 mm/year. The mean GA change in hypertransmission 1 year before and after treatment initiation was 0.0085 mm/year and 0.135 mm/year.

**Conclusions** Geographic atrophy progressed in all eyes except one. According to a 2021 Ophthalmic Research meta analysis, the average GA growth rate seen across 23 studies was 0.33 mm/year. Thus, our results indicate that the carbidopa-levodopa treatment provides benefit in slowing progression rates of GA. Further studies are indicated into the pathogenesis of GA and the role that carbidopa-levodopa might play in its treatment.

**#129** HOW DO MEDICAL AND PHYSICAL THERAPY STUDENTS LEARN ABOUT CONCUSSIONS?

1-3H Wu, 1-3Y Tangori, 1-3JL McKay, 1-3AD Kay, 1-3D Baron. 1Western University of Health Sciences, Pomona, CA; 2Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA; 3The University of Arizona College of Medicine Tucson, Tucson, AZ

10.1136/jim-2022-WRMC.128

**Purpose of Study** This study assesses medical and physical therapy students’ knowledge level in concussion symptoms, diagnosis, and treatment. Parameters we evaluated included how education level, sports background, and concussion history influenced students’ concussion knowledge. The study assessed how these students learn about concussions and whether gaps in knowledge exist. The ultimate goal is to use the survey results to help educators better prepare medical and physical therapy students for patient care.

**Methods Used** The first phase of our study involved sending a 14-question electronic survey to osteopathic medical schools across the United States, which assessed demographics, concussion knowledge level, source of concussion education, and interest in curriculum-based learning. The second phase consists of sending a similar electronic survey that expanded to allopathic and physical therapy schools. This survey consisted of 16 questions, with 2 additional demographic questions inquiring about gender and type of pursued degree.

**Summary of Results** Preliminary collection of over 800 responses and analysis of the data show that 60.2% of MD, DO, and physical therapy students played sports in either high school, college, or professionally. In addition, 42.7% of participants reported sustaining at least one concussion throughout their lives. 26.9% of our participants reported learning about concussions through non-academic means, while 70% reported learning via academic means such as through lectures, literature reviews, or clinical rotations. Our results showed that 80% of our participants agreed they would like more formal training in non-academic methods. While data is forthcoming, this may indicate that an alternative means to learning about concussions is through a sports background and/or concussion history. Participants agree that in order to solidify or supplement concussion knowledge, more education is needed to best prepare rising health care professionals in clinical settings.

**#130** NEW THROMBUS FORMATION IMMEDIATELY AFTER ANDEXANET ALFA INFUSION: A CASE REPORT AND LITERATURE REVIEW

1-2T Torbati, 1OA Elshaigi*, 1M Kayali, 1OM Dumitrascu. 1Western University of Health Sciences, Pomona, CA; 2Cedars-Sinai Medical Center, Los Angeles, CA; 3Mayo Clinic Arizona, Scottsdale, AZ

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**Case Report** Andexanet alfa was FDA approved in May 2018 to reverse anticoagulant effects of Factor Xa inhibitors like Apixaban and Rivaroxaban, thereby generating pro-thrombotic mechanisms. Thromboembolic complications within 30 days of Andexanet alfa administration have been reported. Here, we present for the first time a thrombotic cerebral event that appeared immediately after Andexanet alfa infusion in a patient with acute intraventricular hemorrhage (IVH).

A 73-year-old man presented to our emergency department with sudden onset of a severe headache. Head CT demonstrated 2.4 mL of IVH. CT angiogram showed 60% stenosis of the left supracholinoid internal carotid artery (ICA). The patient had been taking 5 mg Apixaban twice daily for atrial fibrillation, with his last dose 3.5 hours prior to presentation. IVH indicated the patient may benefit from anticoagulation reversal via Andexanet alfa.

A 400 mg bolus of Andexanet alfa was administered followed 30 minutes later by a 2-hour infusion of an additional 480 mg, immediately upon which the patient exhibited global aphasia, temporarily alleviated by head-of-the-bed flattening. A left ICA territory mismatch (342 mL) and 76 mL core infarct were observed on CT perfusion. Shortly afterwards, the patient developed a persistent and severe left middle cerebral artery (MCA) stroke syndrome with NIH stroke scale (NIHSS) score of 23. Emergent cerebral angiogram was then performed, revealing a new sizeable thrombus in the left cervical ICA. Successful thrombectomy yielded a resulting TICI score of 2B. However, neurologic status remained poor due to development of a large left MCA territory infarct, and the patient’s family chose to withdraw supportive care.

Our observation of a thrombotic event induced immediately after Andexanet alfa challenges current administration guidelines. The ANNEXA-4 phase 3 trial reported thrombotic events within 7 days of Andexanet alfa administration in 4% of subjects and within 30 days in 10%. The earliest event in literature was noted 1 day following treatment. Also, Andexanet alfa appears to have a higher thrombotic risk than other reversal agents like four-factor prothrombin complex concentrate by as much as 7%. Further elucidation of its effects on the coagulation cascade are warranted to improve safe clinical practices. Interestingly, no thrombotic events were reported in ANNEXA-4 patients who restarted anticoagulation protocols. Thus, it may be advisable to monitor patients closely and broadly for thrombotic events until future studies update protocols for resuming anticoagulation therapy after reversal treatment.
Pulmonary and critical care
Concurrent session
3:15 PM
Thursday, January 20, 2022

#132 PLASMA NEUTROPHIL EXTRACELLULAR TRAP LEVELS CORRELATE WITH ACUTE RESPIRATORY DISTRESS SYNDROME SEVERITY

J Aoki*, F Denorme, J Rustad, D Perry, M Cody, E Harris, EA Middleton, CC Yost. University of Utah Health, Salt Lake City, UT

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Purpose of Study Acute Respiratory Distress Syndrome (ARDS) is characterized by hypoxic respiratory failure, multi-organ dysfunction, and mortality. ARDS results from inflammatory alveolar injury precipitated by direct and indirect lung injury. Neutrophils play a central role in the pathology of ARDS and release neutrophil extracellular traps (NETs) to trap and kill pathogens. Dysregulated NET formation, however, can cause inflammatory tissue damage and exacerbate acute lung injury as in COVID-19 associated ARDS. Whether NETs participate pathogenically in non-COVID-19 associated ARDS remains unknown. We hypothesized that plasma NET levels correlate directly with disease severity and mortality in non-COVID-19 ARDS patients.

Methods Used We obtained previously collected plasma samples from patients (n=200) with moderate to severe ARDS enrolled in the Re-evaluation of Systemic Early Neuromuscular Blockade (ROSE) trial at three different time points (admission, 24 hours, and 48 hours after admission) complete with clinical outcome data through 28 days after admission. We also examined age- and gender-matched healthy donor plasma (n=20). We assayed cell-free DNA levels via fluorescence as a surrogate for NETs in each plasma sample. Clinical outcomes from ROSE trial participants were correlated with the quantification of NETs. We also assessed NET formation by neutrophils isolated from healthy adults following incubation with ARDS patient and healthy donor plasma samples using live cell imaging and confocal microscopy.

Summary of Results We demonstrated elevated cell-free DNA in ARDS plasma compared to healthy donor plasma. Deceased study participants demonstrated higher plasma cell-free DNA levels on admission and at 48 hours as compared to ARDS survivors (admission: p = 0.0045 and 48 hours: p = 0.005). Increased cell-free DNA on admission, at 24 hours, and 48 hours also correlated with illness severity. Furthermore, ARDS plasma samples induced NET formation in vitro in neutrophils isolated from healthy donors while control plasma did not.

Conclusions NET formation is increased in plasma from patients with ARDS compared to healthy donor plasma, consistent with the inflammatory alveolar injury seen in ARDS. Additionally, plasma from ARDS patients induces NET formation in vitro in PMNs isolated from healthy adult donors. We speculate that exaggerated NET formation may serve as a novel biomarker for inflammatory lung injury in ARDS resulting from multiple etiologies and strategies targeting NET formation may improve outcomes in ARDS.
#134 EVALUATION OF THE ACCURACY OF MINIMALLY-INVASIVE CARDIAC OUTPUT MONITORS BEFORE AND AFTER CARDIOPULMONARY BYPASS

K Cheung*, N Fleming. University of California Davis, Sacramento, CA

10.1136/jim-2022-WRMC.134

Purpose of Study Cardiac output (CO) monitoring is an important tool for hemodynamic optimization. Bolus thermodilution (iCO) with a pulmonary artery catheter (PAC) remains the gold standard for CO measurement, but is invasive and has been associated with complications. This study evaluates the level of agreement of CO values measured from multiple minimally-invasive CO monitor systems before and after cardiopulmonary bypass (CPB). CCO uses a modified thermodilution technology. Cheetah is based on thoracic bioreactance. ClearSight reconstructs the brachial arterial pressure waveform from the finger arterial pressure. CNAP CO is based on continuous non-invasive arterial pressure from the finger. LiDCO is based on the radial arterial blood pressure waveform. FloTrac calculates CO from the radial arterial pulse contour.

Methods Used The IRB reviewed and approved this quality improvement study. Sixty patients were enrolled. 8 patients were excluded due to missing iCO measurements. CO measurements from 52 patients were evaluated using Bland-Altman analysis. CO values were measured simultaneously by bolus thermodilution with a PAC and the CO monitors listed above.

Summary of Results All values were not available at all time points. The Bland-Altman plots are presented in figure 1 and the corresponding values are summarized in table 1.

Conclusions Based upon percentage errors, the relative accuracy of the minimally-invasive CO monitors when compared to iCO were: CCO>Cheetah>ClearSight>FloTrac>CNAP>LiDCO. Measurements after CPB have slightly smaller percentage of

Abstract #134 Figure 1 Bland-altman analysis of minimally-invasive CO monitors pre-bypass and post-bypass
Abstract #134 Table 1  Bland-altman analysis of minimally-invasive CO monitors

<table>
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<th>PRE-BYPASS:</th>
<th>CCO</th>
<th>Cheetah</th>
<th>ClearSight</th>
<th>FloTrac</th>
<th>CNAP</th>
<th>LiDCO</th>
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<td>178</td>
<td>183</td>
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<td>-3.5</td>
<td>-5.0</td>
<td>-6.7</td>
<td>-6.6</td>
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<td>To</td>
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<td>1.7</td>
<td>2.1</td>
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EVALUATION OF THE ACCURACY OF MINIMALLY-INVASIVE CARDIAC OUTPUT MONITORS BEFORE AND AFTER CARDIOPULMONARY BYPASS INvasive CARDIAC OUTPUT monitors

K Cheung*, N Fleming.

University of California Davis, Sacramento, CA

Cardiac output (CO) monitoring is an important tool for hemodynamic optimization. Bolus thermodilution iCO with a pulmonary artery catheter (PAC) remains the gold standard for CO measurement, but is invasive and has been associated with complications. This study evaluates the accuracy of minimally-invasive CO monitors that reconstruct the brachial arterial pressure waveform from technology. Cheetah is based on thoracic bioreactance. CCO uses a modified thermodilution method and has been associated with complications. The IRB reviewed and approved this quality improvement study. The corresponding values are summarized in table 1. Summary of Results

POST-BYPASS: CCO>Cheetah>ClearSight>FloTrac>CNAP>LiDCO. CCO was the most accurate monitor, followed by Cheetah, ClearSight, FloTrac, CNAP, and LiDCO. The percentage errors for CCO were: 54.5%, 53.3%, 68.1%, 90.8%, 110.2%, and 110.6% for Cheetah, ClearSight, FloTrac, CNAP, and LiDCO, respectively. The bias and SD of the Bland-Altman analysis are also provided in table 1. Significant differences were observed between the monitors.

Purpose of Study

Signaling in lung epithelial cells plays a role in respiratory disease pathogenesis. ERK, NfkB, and AMPK are key kinases regulating cell growth and proliferation that are implicated in airway inflammation and disease. Importantly, ERK and AMPK display heterogenous and temporally dynamic signaling activity that can be linked to cell behavior but has yet to be investigated in the context of airway disease. We hypothesize that unique signatures of short-term oscillatory signaling activity (minutes) differentially regulate long-term (>24 hour) inflammatory responses in part via regulation of the transcription factor STAT3 at the cellular level.

Methods Used

Using fluorescent biosensors and live-cell imaging, we track single-cell kinase signaling activity in our Human Bronchial Epithelial (HBE1) cell line, and primary human bronchial epithelial cells (pHBE), continuously at 6-minute intervals, both in submerged and Air-Liquid Interface (ALI) culture conditions. Computational image analysis extracts kinase signaling activity profiles in response to growth factors, and inflammatory cytokines. After 24 hours of ligand exposure, cells are fixed and immunofluorescent stained for nuclear pSTAT3 is performed to measure cellular inflammatory response.

Summary of Results

Comparison of HBE1’s ERK signaling activity from control and ligand stimulated cells, reveals heterogenous and dynamic single-cell responses to inflammatory ligands relevant to both subtypes of asthma and COPD (IL-6, IL-1β, & TNFα), that are both ligand- and concentration-dependent. Intriguingly, pHBE cells in submerged culture display dynamic and heterogeneous ERK signaling activity, that is similarly dependent on ligand and concentration, but are unique from those seen in HBE1 cells. Notably, both HBE1 and pHBE cells displayed unique ERK responses to inflammatory ligands than those of EGF treated cells. We note a similar heterogeneity in STAT3 activation dependent on inflammatory ligand, that is attenuated in the presence of metabolic perturbation and AMPK activation.

Conclusions

These results support our central hypothesis and demonstrate the importance of this novel and unique approach using both airway epithelial cell lines and primary bronchial epithelial cells in ALI. Ongoing work: 1) Finishing data collection for ERK and NfkB in pHBE cells cultured in ALI, and AMPK activity in HBE1 cells, 2) Using statistical modeling to determine ERK, AMPK, and NfkB’s contribution to the patient had been vaping with THC products including dabs. Dabs, a wax-like THC product, are likely the mechanism by which lipids were introduced into his lungs resulting in pneumonia.

Discussion

This case represents a presentation of lipid pneumonia secondary to ‘dabbing,’ a relatively novel form of ingesting cannabis. There have been few reported cases of respiratory failure secondary to ‘dabbing,’ and this case identifies lipid pneumonia as the cause of lung injury. This case highlights the need for physicians to be aware of specific forms of recreational drugs and routes of delivery.
modulating the STAT3 response to inflammatory ligands, and
3) Assessing how pharmacological agents alter activity profiles to modulate long-term inflammatory responses in these model systems. Our technique will reveal deeper knowledge about airway epithelial kinase signaling mechanisms relevant to asthma and COPD, and potentially, other lung diseases.

#137 THE EFFECTS OF ACUTE AND CHRONIC KIDNEY INJURY ON MORTALITY IN OLDER PATIENTS CRITICALLY ILL WITH COVID-19

MD Rockstrom*, S Windham, KM Erlanson, J Hippensteel. University of Colorado, Denver, CO

10.1136/jim-2022-WRM.C135

Purpose of Study Older age is a well-recognized risk factor for increased mortality due to COVID-19. Similarly, older age is associated with high rates of critical illness and admission to intensive care units. Limited data suggests that frailty, as measured by the presence of comorbidities prior to admission, may predispose to severe illness. There is currently an inadequate description of what factors can be attributed to the increase in mortality in aged, critically ill patients with COVID-19.

Methods Used In this retrospective, cohort study, we examined 200 patients admitted to a quaternary referral center intensive care unit (ICU) with COVID-19 from March to June of 2020. Data collected included demographics, number of comorbidities, residence in assisted living or skilled nursing facilities prior to admission, vital signs, and laboratory findings at time of admission to the hospital and the ICU.

Bivariate analysis was performed with age, comorbidities, and place of residence prior to admission as independent variables and mortality, length of stay, rates, and severity of acute respiratory distress syndrome (ARDS), rates of acute kidney injury (AKI) and hemodialysis, and rates and severity of shock as dependent variables. Multivariate analysis evaluated the relationship of age, comorbidities, end organ dysfunction, and disease severity.

Summary of Results ICU mortality was positively correlated with age (p < 0.001) and number of comorbidities (p < 0.001). Furthermore, age and number of comorbidities were directly correlated with development of AKI (p = 0.012); age was inversely correlated with severity of lung disease as measured by P:F ratio (p = 0.0092). Pre-existing chronic kidney disease (CKD) was highly predictive of development of AKI (OR 6.93, p = 0.003). Multivariable analysis demonstrated that the odds of mortality were higher with increasing age (OR = 1.06 [95% CI 1.02, 1.11] per year, p = 0.005), decreased with increasing P:F ratio (OR 0.336 [95% CI 0.16, 0.61] for every increase in P:F of 50, p < 0.001), and increased with AKI requiring hemodialysis (OR = 4.61 [95% CI 1.185, 19.7], p = 0.031).

Conclusions Recognition of CKD in older adults critically ill with COVID-19 is critical in identifying those who are high risk of severe kidney dysfunction and death. CKD has been shown previously to be associated with mortality in COVID-19 and this study offers a mechanistic understanding of this relationship. This is especially important given that older patients do not seem to have the same propensity for severe lung disease seen in younger patients. The role of kidney disease in the mortality of older patients critically ill with COVID-19 needs to be better defined prospectively.

#138 EFFECT OF POSITIVE END EXPIRATORY PRESSURE IN PREVENTION OF POSTOPERATIVE PULMONARY COMPLICATIONS IN OBSE Patients UNDER GENERAL ENDOTRACHEAL ANESTHESIA: SYSTEMATIC REVIEW AND META-ANALYSIS OF RANDOMIZED TRIALS

JY Choi*, MA Al-Saedy, BJ Carlson. Washington State University, Spokane, WA

10.1136/jim-2022-WRM.C136

Purpose of Study In non-obese patients, low positive end-expiratory pressure (PEEP) of < 5cmH2O is a widely accepted practice. Although the safety of low tidal volume plus low PEEP has been established in non-obese patients, this strategy may induce atelectasis and post-operative pulmonary complications in obese patients. In obese patients, use of PEEP > 10cmH2O has shown to prevent intraoperative atelectasis but there is no conclusive data for the postoperative complications from high PEEP. Therefore, this meta-analysis aims to compare the rate of postoperative complications (PPCs) associated with high PEEP and low PEEP in obese patients who received surgery under endotracheal general anesthesia.

Methods Used The protocol for this meta-analysis was published on PROSPERO (CRD42021224041). We searched MEDLINE, EMBASE, CENTRAL, and CINAHL databases using pre-specified search strategies. We included only RCTs that compared the effect of low PEEP and high PEEP in obese patients (BMI>30 kg/m2) who received open or laparoscopic surgeries under endotracheal general anesthesia. All patients received the tidal volume of 6–8 ml per kg of predicted body weight. Risk of bias was assessed using the Covidence review manager. Statistical analysis was conducted using Cochrane RevMan software (Mantel-Haenszel, Fixed Effects, Risk Ratio). Primary outcome was the number of PPCs. Secondary outcomes included extrapulmonary complications, intraoperative complications, and mortality during hospital stay.

Summary of Results There was no significant difference in the rate of postoperative complications between low PEEP and high PEEP groups (RR=0.93; 95% CI 0.79 to 1.10; P=0.41). We identified 4 randomized controlled trials involving 2,116 participants. Two trials had a low risk of bias, and the other two trials had an immediate and substantial risk of bias. We observed significant heterogeneity within included trials (I²=53%). Secondary outcomes were not reported by all trials. Incidence of intraoperative hypotension was reported in 3 trials and was significantly lower in the low PEEP group (RR=1.84; P<0.01).

Conclusions Overall, the evidence is not robust enough to determine the protective effect of high PEEP and low PEEP in obese patients. Out of 2,116 total participants, 1,976 participants were from 1 trial that included both laparoscopic and open abdominal surgery. This may explain the heterogeneity observed since the other three trials only included patients who received laparoscopic abdominal surgeries. Although exclusion of the large study significantly lowered the incidence of PPCs in the low PEEP group with relative risk of (RR=8.00 P=0.05), the other trials were smaller studies and had higher risk of publication bias. More multi-center RCTs are warranted to compare the rate of PPCs between low PEEP and high PEEP in obese patients under endotracheal general anesthesia.
Case Report Pulmonary Vasculitis is a manifestation of a specific set of disorders pathologically defined by inflammatory destruction of blood vessels within the lungs. Secondary immune mediated vasculitis, as in autoimmune processes like rheumatoid arthritis (RA), is one subset of these disorders. We present a rare case of alveolar hemorrhage caused by pulmonary vasculitis in a patient positive for anti-CCP antibody but no laboratory or imaging to meet diagnostic criteria for RA.

CASE A 50-year-old female with history of pulmonary embolism (PE) presented to the ED with three episodes of hemoptysis. She also complained of chest pain, arthralgias and fatigue. She was not on anticoagulation therapy. Family history was positive for lupus and RA in two sisters. Vital signs were stable. Labs showed ESR 36, CRP 1.53, and D-dimer 749. CT chest angiogram revealed scattered ground glass and patchy alveolar densities throughout lung fields, no evidence of PE. Flexible bronchoscopy with bronchoalveolar lavage (BAL) was diagnostic for alveolar hemorrhage. Cytology from BAL yielded numerous red blood cells and macrophages with hemosiderin granules. Autoimmune panel was significant for elevated ANA 1:80 and CCP Ab IgG >250. Patient was started on Prednisone 60 mg/day for diagnosis of alveolar hemorrhage secondary to immune mediated capillaritis and discharged home. As an outpatient, X-rays of wrists and hands revealed no inflammatory or crystal line arthropathy. Labs showed normal rheumatoid factor (RF), but persisting elevation in CRP and ESR. Methotrexate was started to prevent recurrent alveolar hemorrhage and Prednisone was continued pending rheumatology referral.

Discussion Although there have been some rare cases of diffuse alveolar hemorrhage (DAH) preceding the diagnosis of RA, DAH commonly arrives as a manifestation of long standing RA. Anti-CCP antibody is known to not only be as sensitive as RF but significantly more specific when diagnosing RA. Studies have shown a subgroup of patients testing positive for anti-CCP antibodies with no evidence of RA, who subsequently developed RA within short followup.

Conclusion The unique disease course of DAH in patients with underlying features of autoimmune disease is not well documented. Implications of positive anti-CCP antibody but no definitive diagnosis of RA in patients with lung disease still requires further investigation. Anti-CCP antibody has been shown to be highly predictive of future development of RA. Patients that present with pulmonary symptoms like hemoptysis due to autoimmune vasculitis and capillaritis should be closely monitored. Whether to initiate early treatment for RA should also be strongly considered. As such, clinicians need to remain vigilant when suspecting immune-mediated DAH in patients with unclear autoimmune disease.

Purpose of Study For decades, hard backboards have been the standard of care for emergency medical personnel and ski patrollers treating known or suspected spinal cord injuries in the field. Recent studies have shown that prolonged immobilization on hard backboards can cause injuries to patients and are no longer recommended unless specific criteria are met. In 2018, the National Ski Patrol implemented guidelines consistent with the Position Statement of the National Association of EMS Physicians and the American College of Surgeons Committee on Trauma. The effectiveness of those guidelines for ski patrollers in the field are quantified in this survey study.

Methods Used This study was approved by the Western University of Health Sciences Institutional Review Board. An anonymous online survey was distributed electronically to 168 ski patrollers across the United States using contact information obtained from the National Ski Patrol website. The survey was optional, and participants voluntarily completed the survey between April 2 and April 27, 2021. A total of 19 ski patrollers responded to the survey and 19 are represented in the data. They were asked questions regarding how many backboards they used per season before and after the new guidelines were implemented, frequency of backboard-specific training, whether the mountain also has a paid patrol division, as well as confidence of their patrol utilizing the new spinal protection guidelines.

Summary of Results The majority of respondents (14 out of 19) reported a decrease of at least 5 backboards used in the season following implementation of the new guidelines. Patrols that conducted more than two backboard-specific trainings per year reported a larger decrease in backboards (6–10 per season) in the season following the new guidelines than patrols that conducted 1–2 yearly trainings (up to 5 per season). The average reduction in backboards in both patrols that have a paid division and those that do not was approximately 5 backboards per year. 10 out of 19 patrols reported that they were ‘very confident’ their patrols could implement the new guidelines and 9 out of 19 reported they were ‘somewhat confident’. No respondents reported that they were ‘not very confident’.

Conclusions The new guidelines and training from the National Ski Patrol resulted in a decrease in hard backboard use by the majority of surveyed patrols while still allowing ski patrollers to utilize potentially life-saving devices when necessary.

Surgery II
Concurrent session
3:15 PM
Thursday, January 20, 2022

Purpose of Study The Garden classification has been used to grade femoral neck fractures since 1961. Nondisplaced
Alterations in Shoulder Tendon Structural Proteins in Atherosclerosis

WH Fang*, S Sekhon, FG Thairkam, DK Agrawal. Western University of Health Sciences, Pomona, CA

Purpose of Study Hyperlipidemia is a hallmark of the atherosclerotic process and can impact every system of the body, including the musculoskeletal system as evident from the increased comorbidity of tendinopathies in atherosclerotic patients. Indeed, lipid deposits within the extracellular matrix (ECM) have been found in tendon tissues with changes in the biomechanical properties of the tendon. However, there is limited information on the development and progression of tendon pathology in atherosclerotic patients. Here, we examined the expression status and molecular crosstalk of the ECM proteins in atherosclerosis using hyperlipidemic microswine model.

Methods Used Shoulder tendons tissues (N=12) from hyperlipidemic Yucatan microswine were harvested, fixed, embedded in paraffin and longitudinal thin sections were used for tissue morphology with hematoxylin and eosin (H&E), Masson trichrome, and Pentachrome staining.

Immunofluorescence staining (IF) was performed for collagen types I, III, IV, V, VI, and XVII, MMP2, and MMP9 following standard protocols. The fluorescence intensity (MI) of each protein was quantified using ImageJ software. The variation with respect to control (tendon tissue harvested from normal swine) was calculated from the average MFI/nuclei and the results are presented as log2 fold-change (FC).

Summary of Results H&E staining showed disorganized ECM in atherosclerotic tissue with greater deposition of adipocytes. Trichrome staining revealed disorganization in collagen fibers with poorly defined vasculature and increased infiltration of adipocytes in atherosclerotic tendons compared to the control. The pentachrome staining highlights decreased collagen expression and increased mucin deposition in the atherosclerotic tissues. Also, the atherosclerotic shoulder tendons demonstrated decreased expression of COL III (FC=-0.38 ± 1.49), COL IV (FC= -0.61 ± 1.75), and a pronounced decrease in COL XVII (FC=-1.6 ± 1.85) and COL I (FC=-2.24 ± 0.41). However, there was an increased expression of COL V (FC=1.02 ± 2.13), MMP-9 (FC=0.9 ± 0.60), and a marked increase in MMP-2 (FC=2.05 ± 1.53).

Conclusions The findings demonstrated that there was considerable structural alteration in ECM composition and components in atherosclerotic tendon when compared with normal shoulder tendons. There was a decrease in collagen proteins and an upregulation of the MMP class of proteolytic enzymes. The decreased collagen and increased MMP expression are correlated with increased tendon injury and rupture. Such pathological alterations support the existence of increased comorbidity of tendinopathies in hyperlipidemic patients.

#143 AN INVESTIGATION IN THE EVIDENCE BASE FOR ADAPTOGENS IN WOUND HEALING AND SKIN REJUVENATION

D Sokolov*, S Gupta. Loma Linda University, Loma Linda, CA

10.1136/jim-2022-WRMC.141

Purpose of Study Despite having a long history of use in traditional and herbal medicine, adaptogens have recently reentered the spotlight due to their potential to augment the body’s response to stress. Originally mentioned in 1974 in a Soviet literature review as ‘New substances of plant origin that Increase non-specific resistance’, the term ‘adaptogen’ has grown to include most any compound that increases the body’s resilience against a variety of mechanisms of cellular stress. Considering that the skin is highly affected by many of these pathways, this implies a potential application for adaptogens in the context of specialties concerned with aesthetic outcomes. This study identified adaptogens with an evidence-based clinical application in the areas of skin health and wound healing.

Methods Used A literature search was conducted using the term ‘adaptogen’ in the article databases of Google Scholar,
Inclusion criteria consisted of articles pertaining to skin improvement and anti-inflammatory properties. Additionally, special attention was given to studies that used human keratinocyte cell lines. Exclusion criteria included general information articles, and articles with described benefits outside the realm of skin and regenerative capacity.

**Summary of Results** The number of relevant articles found spoke to a large need for further research. While there are numerous articles and studies done on the systemic benefits of various adaptogens, research specific to skin and wound healing is very limited. However, the research completed consistently shows agreement in the nature of the benefits of various adaptogens. A common theme of inflammation reduction quickly emerged through the course of the search, with the most prevalent pathway affected being NFkB. While Ashwagandha, Schisandra and Triphala held the most promise in terms of an evidence base, the number of applicable articles often corresponded with the total amount of available literature on a certain compound.

**Conclusions** This serves to highlight how much potential there is in this area for further research. If the benefits proposed in the current literature stood the test of clinical studies, adaptogens could become part of standard perioperative and post traumatic care.

#144 **HOW DOES DIMETHYL SULFOXIDE AFFECT IN-VITRO FAT GRAFT VIABILITY?**

NM Safi*, S Gupta. Loma Linda University Adventist Health Sciences Center, Loma Linda, CA

Purpose of Study Since the popularization of fat grafting in the 1980s, surgeons have sought to improve the viability of the grafts. Research demonstrates that the majority of harvested cells die within hours to days of transplantation. One of the most promising interventions which may improve survival during this time are pharmacologic agents added to the graft or recipient site. Several additives have been tried. One such compound is dimethyl sulfoxide (DMSO). In one study, rats treated with DMSO showed improved fat graft survival. Due to the challenge of studying new treatments in humans we sought to observe the impact of DMSO on viability of adipocytes in vitro.

**Methods Used** DMSO was added to samples of human lipoaspirate to produce three concentrations: 500:1, 2000:1, and 8000:1 and a control sans DMSO. The samples were centrifuged to isolate the fat from the aqueous fraction. Each sample was then divided into several 900µL portions and incubated with 100µL of AlamalBlue at 37°C for 4 hours. Each portion was then diluted with 3mL of saline and centrifuged to suspend the dye. Absorbance was measured at 570nm and 600nm. The experiment was repeated thrice. The patients were females aged 41, 61, and 48. Normalized absorbances were calculated by subtracting the 600nm absorbance from the 570nm absorbance.

**Abstract #144 Figure 1**

**DMSO Dosing**

<table>
<thead>
<tr>
<th>DMSO Dilation</th>
<th>Normalized Absorbance</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.000</td>
<td>0.450</td>
</tr>
<tr>
<td>0.300</td>
<td>0.400</td>
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<tr>
<td>0.450</td>
<td>0.350</td>
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<td>0.400</td>
<td>0.300</td>
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<tr>
<td>0.000</td>
<td>0.050</td>
</tr>
</tbody>
</table>

**Abstract #143 Table 1**

<table>
<thead>
<tr>
<th>Name</th>
<th>Number of Included Articles</th>
<th>Pathway of action</th>
<th>Methods</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ashwagandha</td>
<td>8</td>
<td>FOKO3, cortisol, NFkB, 8 hyaluronic acid, testosterone</td>
<td>Literature Review (2), Case Study (1), Research Study (6)</td>
<td>Decreased inflammation, decreased aging protein markers,</td>
</tr>
<tr>
<td>Schisandra</td>
<td>10</td>
<td>ROS scavenging, antioxidant, UVA/B damage reduction, elastase, MPP, glutathione,</td>
<td>Literature Review (2), Research Study (8)</td>
<td>Increased collagen synthesis, decreased damage, increased antioxidant pathways</td>
</tr>
<tr>
<td>Triphala</td>
<td>8</td>
<td>NFkB, Collagen I, antioxidant, MPP, skin oxidative</td>
<td>Literature Study (3), Research Study (5), Case Study (2)</td>
<td>Inhibited MPP, reduced inflammation, reduced RBC hemolysis</td>
</tr>
<tr>
<td>Meca</td>
<td>1</td>
<td>UVA/B damage reduction</td>
<td>Research Study</td>
<td>Topically applied Meca created a reduction in UVA/B induced damage</td>
</tr>
<tr>
<td>Rhodiola Rosea</td>
<td>2</td>
<td>HGF, MPP, skin oxidative</td>
<td>Research Study (2)</td>
<td>Reduced tissue edema, reduced cell damage</td>
</tr>
<tr>
<td>Arctium Lappa</td>
<td>1</td>
<td>NFkB, INOS, TNF-alphaph</td>
<td>Literature Review</td>
<td>Reduced inflammation</td>
</tr>
<tr>
<td>Ganoderma Lucidum</td>
<td>1</td>
<td>MMP, CK2/6, p53</td>
<td>Research Study</td>
<td>Increased cell proliferation and migration</td>
</tr>
</tbody>
</table>
Summary of Results Trial #1, (figure 1), showed a decrease in absorbance across all doses of DMSO tested [Control (0.386), 1:8000 (0.251 p = 0.004), 1:2000 (0.317), and 1:500 (0.381)]. Trial #2 (figure 2), demonstrated a dose-dependent decrease in absorbance across the three concentrations [Control (0.201) 1:8000 (0.186, p = .03), 1:2000 (0.183, p = 0.2), 1:500 (0.145, p = 0.02)]. Finally, trial #3, (figure 3) yielded an increase in absorbance at a low (1:8000) concentration of DMSO (0.182, p = 0.02) vs Control (0.140). 1:2000 and 1:500 were found to have a negative effect on absorbance (0.106 p = 0.1 and 0.102 p = 0.07, respectively).

Conclusions Our results are, thus far, inconclusive. However, we believe that there are confounds which can be eliminated. The researcher noted an inconsistent mixing of assay with fat and also noted that small variations in pipetting technique introduced bubbles in the cuvette that could affect readings. If minor technique changes rectify these errors, more consistent results are possible.

If future in vitro trials find DMSO to offer a viability benefit to graft cells, further research could be done on topical applications in humans as DMSO has been safely used topically for years.

#145 COMPARISON OF COMPLICATIONS IN PATIENTS WITH NORMAL AND COMPROMISED RENAL FUNCTION UNDERGOING ADJUVANT RADIATION THERAPY FOLLOWING ROBOTIC-ASSISTED LAPAROSCOPIC RADICAL PROSTATECTOMY

1R Chen*, 2S Song, 2A Amasyali, 2C Ritchie, 1C Baas, 2D Baldwin. 1Loma Linda University, Loma Linda, CA; 2Loma Linda University Adventist Health Sciences Center, Loma Linda, CA

Purpose of Study Chronic kidney disease (CKD) and end-stage renal disease (ESRD) are associated with high morbidity and mortality. Kidney transplantation may be a life-saving therapy for these patients. However, ESRD patients with untreated prostate cancer may be excluded from transplantation due to risk for cancer progression, while patients treated with robotic-assisted laparoscopic radical prostatectomy (RALP) who have undetectable PSA’s may be cleared for immediate transplantation. Patients with detectable PSA’s may require adjuvant radiation, which may result in radiation cystitis, a clinical diagnosis ranging from mild dysuria to severe hematuria. In patients with prior RALP, adjuvant radiation therapy (XRT) may result in a higher mean radiation dose to the bladder due to its low volume and new post-surgical location, which may increase the risk for hemorrhagic cystitis. The purpose of this study was to compare the complications between patients with normal and compromised renal function undergoing adjuvant radiation therapy following RALP.

Methods Used A retrospective review was conducted of all patients in a single academic institution undergoing RALP followed by XRT between December 2006 and July 2020. Demographic variables, surgical parameters, cancer stage, outcomes and complications were compared between patients with CKD 0–2 and CKD 3–5. The primary outcome was radiation related bladder complications including hematuria, catheter insertion, blood transfusions, continuous bladder irrigation, surgical clot evacuation, and bladder fulguration. Statistical analysis was performed by Mann Whitney U and Independent T test with p<0.05 considered significant.

Summary of Results When comparing the 101 patients with CKD 0–2 who underwent RALP followed by XRT, to the 9 patients with CKD 3–5, there was no difference in age (67.7...
#146 FACTORS ASSOCIATED WITH SPONTANEOUS LATERAL SKULL BASE CEREBROSPINAL FLUID
JS De Armas*, E Miles, A Miller, Y Liu, N Wyckoff. Loma Linda University Medical Center, Loma Linda, CA
10.1136/jim-2022-WRMC.144

Purpose of Study Most lateral skull base CSF leaks have an identifiable cause, however, spontaneous cerebrospinal fluid (CSF) leaks do not. We aimed to determine risk factors for spontaneous lateral skull base CSF leaks.

Methods Used Retrospective chart review of all patients seen at a tertiary referral center over the last 10 years for spontaneous lateral skull base CSF leak who underwent temporal bone CT scan. Patients with spontaneous CSF otorrhea were included. Patients with lateral skull base CSF leaks secondary to known causes such as trauma, cholesteatoma, or iatrogenic were excluded. Sex and age-matched controls were randomly selected from all patients seen at the same center with temporal bone CT scan and without major otologic pathology. Demographics collected and main outcomes analyzed included age, sex, ethnicity, body mass index (BMI), smoking status, diagnosis of obstructive sleep apnea, and other comorbid medical conditions. Statistical analysis included analysis of variance, student t-test, and chi-squared between CSF leak patients and controls.

Summary of Results Fourteen patients were identified with spontaneous lateral skull base CSF leak. 32 sex and age-matched controls with CT temporal bone imaging were randomly selected. 93% of patients in the CSF leak group were female with an average age of 66.9 years. Patients in the CSF leak group had higher BMI (33.4 ± 27.4) vs 22.4 (p < 0.01) with higher rates of cardiovascular disease (93% vs. 63%, p = 0.04) and sleep apnea or snoring (43% vs. 3%, p < 0.01).

Conclusions Risk factors associated with spontaneous lateral skull base CSF leaks include female gender, elevated BMI, OSA, and cardiovascular disease. Further research is warranted to identify potential associations between lateral skull base thickness and spontaneous lateral skull base CSF leaks.

#147 UTILITY OF PEDIATRIC VESICOSTOMY IN PROTECTING THE URINARY TRACT
J Byer*, C Ritchie, J Chamberlin. Loma Linda University School of Medicine, Loma Linda, CA; *Loma Linda University, Loma Linda, CA
10.1136/jim-2022-WRMC.145

Purpose of Study In pediatric patients, a vesicostomy is a surgical option for temporary or permanent urinary diversion as a protective measure for the integrity of the patient’s urinary tract and renal system. The objective of this study is to quantify the indications for the pediatric patient population undergoing vesicostomies at an academic children’s hospital.

Methods Used We performed a retrospective chart review of all pediatric patients (0–18 years) who underwent cutaneous vesicostomy from a single tertiary children’s hospital from 2002 to 2021. Demographic information, pathological indications for vesicostomy, medical management, urinary tract infections, urodynamics, renal function, complications, and reversal of vesicostomy were evaluated. The primary outcome was improvement in hydrenephrosis and vesicoureteral reflux, comparing before and after vesicostomy. Two-tailed, Student-t tests were calculated, with p values less than 0.05 considered significant.

Summary of Results At our institution, 33 pediatric patients (20 males and 13 females) underwent vesicostomy. The age at time of surgery ranged from 0 to 179 months (mean 70.5 months) with median follow-up 70.5 months (range 2–210). The indication for vesicostomy included neurogenic bladder (14), chromosomal anomalies (8), anatomical malformation (7), prune belly syndrome (4), secondary vesicoureteral reflux (3), posterior urethral valves (3), and solitary kidney (3). Two patients underwent vesicostomy prior to kidney transplant clearance and one had a vesicostomy concurrently with kidney transplant. Prior to surgery, 21 (63.6%) were started on clean intermittent catheterization and 20 (60.6%) were on an anticholinergic. Fifteen (45.5%) had ≥2 confirmed UTIs before surgery (range 0 to 11). Either unilateral or bilateral SFU Grade ≥2 Hydrenephrosis was present in 25 patients (75.8%) prior to surgery and present in 12 (36.4%) after surgery (p < 0.01). Vesicoureteral reflux (3) Grade ≥2 was present in 15 patients (40.5%) prior to surgery and persistent in 5 (15.2%) after surgery (p = 0.036). The median percentage of expected bladder capacity was 83.3%. Complications after vesicostomy included vesicostomy site fungal rash (23), sepsis due to a UTI (10), and renal or bladder calculi (4). Additionally, 4 developed stricture, 4 developed stomal stenosis, 8 had to catheterize the stoma to maintain patency, and 5 had bladder prolapse through the vesicostomy site with 2 requiring revisional surgery. Six underwent excision and closure of their vesicostomy. One had their cutaneous vesicostomy modified to an ileovesicostomy. Two underwent repeat vesicostomy.

Conclusions Cutaneous vesicostomy are useful and effective forms of urinary diversion in pediatric patients with unsafe urinary tracts that have otherwise been refractory to medical treatment.

#148 DOES A NEEDLE HOLDER REDUCE FLUOROSCOPIC RADIATION EXPOSURE TO THE SURGEON’S HAND COMPARED TO LEAD GLOVES AND CONVENTIONAL GLOVES?
E Joo*, C Baas, R Chen, JD Hartman, JD Belle, A Amasyali, D Baldwin. Loma Linda University School of Medicine, Loma Linda, CA; *Loma Linda University Adventist Health Sciences Center, Loma Linda, CA
10.1136/jim-2022-WRMC.146

Purpose of Study Fluoroscopy is commonly employed while gaining needle renal access prior to removal of kidney stones during percutaneous nephrolithotomy (PCNL). Excess radiation...
exposure has been linked to possible deleterious outcomes for both patients and surgeons. Some of the radiation the surgeon receives is due to scatter radiation, but the surgeon’s hand may be exposed to the highest radiation dose when placed directly in the fluoroscopy beam. To reduce this exposure, some surgeons wear lead gloves, and more recently, a specialized needle holder has been developed. The purpose of this study is to compare direct radiation dose to the surgeon’s hand while holding the needle with a conventional surgical glove, a lead glove, or a novel needle holder.

Methods Used A PCNL procedure was simulated using a cadaver to represent a patient and a separate cadaver arm to mimic a surgeon’s hand. Three different techniques for holding the needle were tested: holding the needle directly in the fluoroscopy beam while wearing a conventional surgical glove, holding the needle while wearing a lead glove, and using a novel needle holder designed to distance the surgeon’s hand from the direct beam. Thermoluminescent dosimeters were attached to the thumb, middle finger, hypothenar eminence, wrist of the cadaver hand, and skin of the cadaver patient, dorsal to the kidney. Five trials were performed for each treatment arm. In each trial, five minutes of fluoroscopy was delivered using automatic exposure control. Radiation doses between treatment arms were compared using ANOVA with p < 0.05 considered significant.

Summary of Results Using a lead glove resulted in a significant reduction of radiation to the surgeon’s hand compared to when the surgeon directly held the needle wearing a conventional glove (mean dose 248.4 vs. 391.9 mRad; p < 0.001). However, the greatest radiation reduction was seen when using the novel needle holder (137.3 mRad) compared to a lead glove (248.4 mRad) and a conventional glove (391.9 mRad; p < 0.001). An unexpected finding was a significant reduction in the cadaver patient’s radiation dose when the novel needle holder was used, compared to the use of a lead glove and a conventional glove (703 vs 842.90 and 816.65 mRad, respectively; p = 0.024).

Conclusions This study showed that using a needle holder during PCNL reduced radiation exposure to the surgeon’s hand compared to the use of a lead glove and a conventional glove. In addition, by removing dense objects from the radiation field, the dose to the patient was significantly less when a needle holder was used. These findings demonstrate that keeping the surgeon’s hand out of the beam will reduce radiation exposure, not only to the surgeon, but also to the patient, and may thereby decrease the risks of radiation exposure, including cancer.

Abstract #150

COMPARISON OF SMARTPHONE TO WEARABLE SENSORS FOR ASSESSMENT OF GAIT IN AN ORTHOPAEDIC CLINIC

Purpose of Study Recently, a host of smartphone-based applications have been developed that advertise the ability to effectively measure gait kinematics compared to traditional methods. These applications, if proven to have clinical utility, would provide a more accessible and affordable alternative for measuring gait on a long-term, continual basis. Thus, the aim of this study is to compare the utility of a smartphone-based gait analysis platform to that of an inertial motion capture (IMC) wearable sensor in a clinical setting.

Methods Used Gait data was collected from 7 patients presenting to the Orthopaedic clinic over a 1-month period. During each appointment, the patient performed a Self-Paced Walking Test (SPWT) with 5 LEGSys wearable sensors attached to their thighs, shanks, and waist. After this test, the most recent
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values for the patient’s Step Length and Walking Speed were obtained from the Health application found in their iPhone and recorded in RedCap, a secure web application for managing online databases. A paired t-test was performed to compare the means of the values of Stride Length (m) and Walking Speed (m/s) recorded from the patient’s iPhone compared to that of the LEGSys sensors. This quality study was granted an IRB exemption, and patient data was de-identified before analysis.

Summary of Results Overall, the means for the iPhone and LEGSys Stride Length were 1.13 (m) and 1.17 (m), respectively, with a p-value of 0.6. The means for the iPhone and LEGSys Walking Speed were 0.906 (m/s) and 0.963 (m/s), respectively, with a p-value of 0.3. (Table 1).

Conclusions In this study of 7 participants, it was shown that there is not a notable difference between the mean measurements of Stride Length (m) and Walking Speed (m/s) as measured by iPhone and LEGSys wearable sensors, thus providing evidence that smartphone sensors may be useful in measuring these gait parameters. Smartphone-based gait sensors have the potential to serve as a useful clinical alternative to traditional wearable sensors in measuring gait. They are cost-effective and easily accessible in a clinical setting, saving time for the clinician during a patient encounter. Future research should focus on collecting data on more patients comparing these two technologies to increase the sample size and power of the study.

Poster session

Adolescent medicine and general pediatrics

6:00 PM
Thursday, January 20, 2022

#151 ABSTRACT WITHDRAWN

#152 IMPLEMENTING TELEHEALTH INTO PEDIATRIC TRAUMA CONSULTATIONS

S Garrison*, JP Marcin, J Galante, JL Rosenthal, T Rinderknecht, K Grether-Jones, MY Hamline, M Zwienerberg, B Haus, K Matthews, R Orominger, A Sanders, R Dizon, N Kuppermann, UC Davis Health, Sacramento, CA; UC Davis Children’s Hospital, Sacramento, CA; University of California Davis Health System Department of Orthopaedic Surgery, Sacramento, CA

10.1136/jim-2022-WRMC.149

Purpose of Study We are evaluating the implementation of telehealth and image sharing services to connect pediatric trauma specialists to rural and community hospital emergency departments to create a Virtual Pediatric Trauma Center (VPTC). The goal of the study is to compare the current standard of care to the VPTC model of care with regards to the parent/family experience of care, distress, healthcare utilization, and out-of-pocket cost burden.

Methods Used We are comparing the current standard of care to the VPTC model of care with regards to: 1) the parent/family experience of care and distress; 2) the 30-day healthcare utilization following the injury event; and 3) the out-of-pocket costs and financial burdens experienced by parents/families 3-days and 30-days following the injury. We’ve implemented a stepped wedge trial among a stratified selection of 10 hospital EDs in Northern California, with a goal of enrolling 380 patients by November 2022. We’ve deployed pole mounted videoconferencing units with high-definition monitors, omnidirectional microphones, and remote-controlled pan-tilt-zoom cameras. Pediatric trauma providers were provided access to workstations as well as video capable mobile devices.

Summary of Results To date, 122 pediatric trauma patients have been enrolled. Protocol adherence for videoconferencing has occurred in 18 of 43 eligible patients; adherence for image sharing has occurred in 16 of 38 eligible patients. Collection rates of surveys of parent/family experience of care, distress, and financial burden surveys has been 65%. The ability of the specialty providers to connect using telemedicine has been limited by other clinical responsibilities, and protocol relies heavily on the NPs to administer telehealth communications and recommendations. Changes in implementation strategies and the workflow were made to increase the reliability and fidelity of the intervention, which will be shared in the presentation.

Conclusions Implementing telehealth into acute pediatric trauma care is challenging. Challenges include incorporating videoconferencing by busy providers and implementing project-specific platforms at partner sites. However, with this commitment, acutely injured children can receive regionalized pediatric trauma expertise at the bedside in a receiving hospital ED. Current findings illustrate the need for qualitative data to improve the family experience.

Poster Session

Behavior and Development

6:00 PM
Thursday, January 20, 2022

#153 A SURVEY OF STUDENT ATTITUDES ABOUT ONLINE LEARNING VIA THE ZOOM PLATFORM

C Bauer*, A Craft. Western University of Health Sciences, Pomona, CA

10.1136/jim-2022-WRMC.150

Purpose of Study Despite decreasing attendance at live in-person lectures, this teaching method has remained a mainstay in medical education. Prior surveys of medical students have found that M1 students attend live lectures more than M2 students, while both years utilize the lecture recordings to supplement their education. With the COVID-19 pandemic, more lecture content has moved online, with some delivered synchronously and some recorded for asynchronous review.

A study of dental students’ perception found that 44% of students felt learning ‘somewhat worsened’ and 26% thought learning ‘significantly worsened’ with the move to online platforms during COVID-1. The same study found that students preferred synchronous learning experiences despite often choosing not to attend live lectures prior to the pandemic. This study seeks to evaluate similar issues in osteopathic medical students to determine a preferred mode of
online learning, perception of their learning, and signs of burnout.

Methods Used An anonymous survey was created and distributed via email to the students in the Classes of 2022, 2023, and 2024 at the College of Osteopathic Medicine of the Pacific. The survey consisted of 8 questions, three were yes or no responses, two were categorical responses, and three questions were based on a 10-point Likert scale. A goal of a 20% return of surveys was pre-established.

Summary of Results Of the 995 students who received the survey, 292 (29.3%) completed the anonymous online survey. 54.1% of respondents were in the Class of 2024, 34.2% were in the Class of 2023, and 11.6% were in the Class of 2022.

On a 10-point scale, with 1 being highly dissatisfied, 55 (18.8%) students selected a score between 1 and 4 when evaluating satisfaction with a synchronous lecture via Zoom video conference technology compared to a live-in-person lecture. 231 (79.1%) students found the video conference lecture as or more helpful than an in-person lecture. 165 (56.5%) respondents noted they were MORE likely to ask a question via the video conference lecture platform, compared to an in-person lecture.

When asked how likely they were to attend live in person lectures in the future, the responses demonstrated that students were evenly distributed, with 41 students (14.1%) responding absolutely not and 45 students (15.4%) responding definitely. Overall, 171 (58.6%) students stated that synchronous video conference-based lectures could fully replace live in person lectures.

Conclusions Overall, students at this single medical school appear to have no significant preference for synchronous video conference-based lectures compared to live in-person lecture. Some students even suggested a greater willingness to participate and ask questions, whether verbally or via the chat function, during video, conference-based lectures. This preliminary data may allow for further investigation into how technology may impact the pedagogy for future generations of medical students.

#154 HOW DO MEDICAL STUDENTS RATE RESILIENTY IN A PANDEMIC?

S Dulani*, E Salimi, Al Nelson, M Hudson. Western University of Health Sciences College of Osteopathic Medicine of the Pacific-Northwest, Lebanon, OR

Purpose of Study To assess the perceptions of medical students about the importance of resiliency in general and their own resiliency in a time of pandemic and to assess what external factors contribute to resilience, especially in the midst of a pandemic.

Methods Used This IRB approved a survey for osteopathic medical students. The student survey questions were beta tested 3 times by students at other universities as well as examined by a professional in the field. All enrolled students at COMP/COMPNW were sent an invitation to participate (~ 1,300 students) ages 18 and up. The invitation letter asked them to self-exclude if they were pregnant or believed that participation in the study could cause them to feel uncomfortable. We based our questions on the academic resilience scale to see if there are any associations between self-perception of stress, relationship style, negative affect and resilience levels in students. The survey consisted of 9 Likert scale questions, 4 multiple choice questions, and 1 open ended question. Multiple questions are derived from previous resilience scales.

Summary of Results We obtained 130 responses (10% response rate). 46% were first year, 24% second year, 14% third year and 16% fourth year students. 51% were female. 40% said they had above average self-esteem, while 30% were neutral and 30% had below average self-esteem. 96% admitted to having anxiety in personal or professional relationships. 52% were likely to reflect on a stressful day, while 54% were likely to reflect on a successful day. This may show that students can be optimistic amid ongoing stress. 85% of students felt confident in their ability to persevere through challenging times. 65% of students were willing to ask for help in times of need while 16% were not willing to, and 19% were unsure. Students are most likely to turn to their spouse, friends, and themselves for support. 75% of respondents who had an opinion said that medical school had been the hardest time of their life. Only 40% of students were satisfied with their academic performance, and only 42% were satisfied with their general wellbeing. Financial hardship contributes to stress levels in 40% of students. The financial hardship does not seem to correlate with those that are unhappy in their academic performance and general well-being.

Conclusions The students surveyed are optimistic; more of them were likely to reflect on a successful day compared to a stressful one. The high rate of anxiety in personal as well as professional relationships may impede students’ ability to seek personal or professional support when they need it. Students who do not have access to friends or family during times of hardship may have a harder time coping with stress.

#155 IMPACT OF COVID-19 ON ROCKY VISTA UNIVERSITY MEDICAL STUDENTS’ MENTAL HEALTH: A CROSS-SECTIONAL SURVEY

D Paz*, V Kuo, M Bairs, V Bandi, M Zueger, M Payton, R Ryznar. Rocky Vista University, Parker, CO

Purpose of Study The high prevalence of anxiety and depression among medical students has been previously linked to the rigor of medical school curriculums. Since late 2019, the COVID-19 pandemic has forced schools to make unprecedented yet unavoidable changes to their educational curriculums. This study is a quantitative analysis of the educational and psychological impacts of COVID-19 on first and second-year osteopathic medical students at Rocky Vista University.

Methods Used A cross-sectional survey was administered to students at Rocky Vista University during the COVID-19 pandemic. Anxiety and depression levels were measured using validated questionnaires GAD-7 and PHQ-9. The remainder of the survey was a self-designed questionnaire that investigated potential associated factors, including sociodemographic characteristics, educational changes, and COVID-19 related concerns. The questionnaire also gauged the impact of COVID-19 on behavioral changes and strategies used to cope with the pandemic.

Summary of Results One hundred fifty-two Rocky Vista University OMS-I and OMS-II students across both campuses participated. Overall, depression and anxiety levels due to the COVID-19 pandemic were increased from previously reported
prevalence rates. Results showed that while the transition to an online based-curriculum was met with mixed feelings, most students found ways to effectively adapt their study habits amidst the pandemic. Notably, students reported feeling concerned that the transition in the curriculum will negatively impact their preparedness for clinical rotations. Overall, COVID-19 was found to have a negative impact on students’ mental health according to nearly all measures. However, the majority of students also reported finding new ways to cope with their stress and anxiety levels and nearly all students reported a willingness to assist with the pandemic.

Conclusions This study demonstrates the significant detrimental effects of the pandemic on medical student mental health, which could impact the quality of care these future physicians provide. Thus, it is imperative to establish effective interventions to mitigate the negative repercussions of the pandemic.

* This work was presented in part at Rocky Vista University Appreciation Day (virtual) on October 15, 2021.

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**SURVEYS OF THE PERCEPTIONS OF FACULTY ABOUT STRESS AND RESILIENCE**

E Salimi*, S Dulani, AL Nelson, M Hudson. Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA

10.1136/jim-2022-WRM153

Purpose of Study To find out what the faculty’s perceptions are about resilience and its relationship to academic success in osteopathic medical school. Can resilience be acquired or is it an innate trait? If it can be acquired, then can it be formally taught? Who is responsible for this formal training and when could it be provided? Should resilience be a part of the DO curriculum?

Methods Used This is an IRB-approved survey for all COMP faculty who interact within students in the pre-clinical years. The questionnaire was beta tested by 10 outside physicians and researchers. It consisted of 13 multiple choice questions and 2 open ended questions. 4 of the 13 questions used Likert scales. Letters of invitation that outlined the study purpose and its voluntary and anonymous nature were sent with the link to the survey, conducted via Qualtrics. Faculty were asked not to participate if they were pregnant or if they anticipated that the topic might cause them discomfort. We also sent out a reminder ~3 weeks after the initial letter that thanked those who had already responded and ask them not to do so again. At the same time, select faculty were approached to see if they would be willing to encourage their colleagues to participate in the study. We calculated the response rate and percentage distribution of the answers.

Summary of Results The response rate is 21%. 55% of the respondents were clinical faculty, 30% basic science faculty, 5% both. 60% thought that resilience is more learned than intrinsic. 34% believed that resilience can be learned through life experiences and another 34% believed that it can be learned through informal teaching in context of individual challenge. 35% believed that there should be a more formal screening for resilience in applicants during the admissions process while 30% disagreed. 100% believed that resilience is either very important or extremely important to a medical student’s success. 75% believed that financial burdens are either very important or extremely important to students’ stress levels to their growth as a physician. 90% believed that resilience improves student’s academic performance by a lot or a great deal. All respondents reported that they work with students in some way to increase their resilience. 65% believe that formal resilience training should be a part of the DO curriculum for students and 85% believe that faculty would benefit from formal training about how to teach resilience.

Conclusions Although osteopathic medicine places importance on holistic approaches to health, there are no requirements in the COCA curriculum to explicitly provide resilience training. The majority of the faculty thought that resilience is more learned than intrinsic. Therefore, it could be worthwhile to explore the possibilities of providing resilience training for both students and faculty.

**NOVEL CHOLESTEROL LOWERING AGENTS IN SETTING OF STATIN Intolerance**

I S Noh*, K Mai, I M Shaver, I S Yong, I M Mostaghimi, G Oh, M Mostaghimi, Western University of Health Sciences, Pomona, CA; University of the Pacific Thomas J Long School of Pharmacy, Stockton, CA

10.1136/jim-2022-WRM154

Purpose of Study Statins are the gold standard lipid-lowering therapy for atherosclerotic diseases. However, there is increasing evidence in support of combination therapies providing greater benefits to patients, particularly those intolerant to statins. Here, we discuss the efficacy, safety, and practical use of novel alternative and adjuvant agents that target low-density lipoprotein cholesterol (LDL-C) or lipoprotein(a) (Lp(a)).

Methods Used A literature review was conducted on PubMed and Google Scholar based on research and clinical trials of cholesterol modulators primarily within the past decade.

Summary of Results PCSK9 antibodies have been effective as an adjuvant therapies in further reducing LDL-C levels and coronary atherosclerosis progression when added to statins. Importantly, the FOURIER and ODYSSEY OUTCOMES trials demonstrated that the addition of evolocumab and alirocumab remarkably reduced the risk of major cardiovascular events, respectively. Yet, expensive cost and administrative burden are impediments to practical use. PCSK9 siRNA incisirs is an appealing alternative because of the marked reduction of LDL-C without significant adverse events as well as its lower cost and administrative burden. Bempedoic Acid (BmA) is another promising LDL-C lowering agent highlighted for its efficacy as both monotherapy and add-on to statins. BmA exhibits a favorable tolerability profile due to its exclusive activation in the liver, suggesting its potential as an alternative to statin-related myopathies. Future studies on its impact on cardiovascular outcomes may provide a greater appreciation of where BmA stands among LDL-C therapies. Antisense oligonucleotide therapies IONIS-APO(a)rx, IONIS-APO(a)Lrx, and AKCEA-APO(a)-Lrx have emerged as encouraging alternatives due to their efficacy and safety as Lp(a)-lowering agents.
Continued investigation as phase 3 trials may establish the potential of Lp(a) therapies as treatments for atherosclerotic disease.

Conclusions Further investigation for all these novel approaches is needed to better appreciate the long-term efficacy, safety, and effects on cardiovascular outcomes. Overall, these cholesterol targeting therapies provide a promising outlook as alternatives for atherosclerotic patients for whom statins have been ineffective or intolerable.

Poster session
Diversity, equity, inclusion research
6:00 PM
Thursday, January 20, 2022

#158 IMPACT OF IMPLEMENTATION OF SMART GOALS FOR RADIOLOGY TRAINEE RESEARCH AND MENTORSHIP

Purpose of Study Active research programs are particularly crucial to the field of radiology because the development of imaging technologies and subsequent translation of them into clinical practice drives the field forward. The SMART (Specific, Measurable, Attainable, Realistic, Time-bound) Goals framework is a validated goal-setting tool that has been successfully applied to medical education. In this prospective cohort study, we evaluated the implementation of the SMART Goals framework in radiology research. We hypothesized that engaging research trainees in formalized goal-setting would result in higher engagement in research and greater productivity.

Methods Used After Institutional Review Board approval, medical students and radiology residents from a radiology research lab at a major academic center between the academic year of August 1, 2020 to July 31, 2021 were invited to participate. Inclusion criteria were those who filled out a SMART Goals form and participated in a formal review meeting. The comparison group was comprised of lab members from previous academic years (August 1, 2016 to July 31, 2020). Productivity was assessed by the number of publications, abstracts, and grants. For purposes of streamlining the data, the acceptance year was defined as the same as the submission year to best match the timeframe during which most of the work was completed. Additionally, participants’ beliefs about their knowledge and abilities were assessed using five-point Likert scale pre- and post-surveys. Descriptive statistics and two-tailed t-tests were calculated. Statistical significance was considered to be \( p < 0.05 \).

Summary of Results There were 19 individuals in the post-SMART group and 28 individuals in the pre-SMART group. The mean number of publications was 0.79 and 1.54 per individual in the pre-SMART and post-SMART groups, respectively (\( p = 0.02 \)). The mean number of abstracts was 1.04 and 2.37 per individual in the pre-SMART and post-SMART groups, respectively (\( p = 0.02 \)). The mean number of successful grants was 0.05 and 0.347 per individual in the pre-SMART and post-SMART groups, respectively (\( p = 0.01 \)). Based on survey results, participants’ knowledge and comfort with research goals increased 1.54 points (\( p < 0.01 \)); with manuscript writing increased 1.21 (\( p < 0.01 \)); and with abstract submission increased 0.60 points (\( p = 0.01 \)).

Conclusions Dedicated learning goal creation through the SMART Goals framework has the potential to significantly increase the number of radiology trainees producing high-quality research, improve medical student and resident knowledge in research, and increase trainees’ confidence in their research skills.
Purpose of Study Interpreting medical literature is an essential skill set for physicians to acquire as physicians must be able to determine the validity of research that may influence their practice. A baseline competency in the basic components of research (i.e., identifying bias, data interpretation, etc.) is imperative for effective clinical practice and clinical research. The purpose of this study is to assess the impact of a journal club on trainee's confidence and knowledge base in fundamental components of research literature.

Methods Used After Institutional Review Board approval, members of the Computational Abdominal Radiology research lab, predominantly medical students and radiology residents, at a major academic center were invited to participate in this study. Journal club sessions were conducted from April to September 2021. Articles from leading peer-reviewed radiology journals were selected to highlight a specific component of research. Pre and post journal club session surveys were distributed among lab members via Microsoft Forms using Likert scales to assess trainee confidence and multiple-choice questions to assess knowledge base. Survey results were compared using a paired two-tail t-test with statistical significance set as p < 0.05.

Summary of Results On average, participants displayed an increase in confidence in topic knowledge (pre-test mean = 4.29 ± 0.31 versus post-test mean = 5.91 ± 0.33, p = 0.004). However, the difference in percent correct answers between the pre-test (mean = 0.77 ± 0.25) and post-test (mean = 0.85 ± 0.13) indicated that there was not a significant increase in correct answers after the journal club sessions (p = 0.42).

Conclusions The results of this pilot study indicate that a journal club can be effective in increasing participants' confidence in the basic fundamentals of research, although its limited application has not yet been shown to increase aptitude. Additional journal club sessions, repeated exposure to key topics, and longer follow up beyond this pilot study are needed to better assess the efficacy of the impact of journal clubs in the setting of research and clinical mentorship in radiology.

Purpose of Study The two most recent community health need assessments performed in Valdez, Alaska identified ‘lack of physical activity/overweight’ as a top health need. 66% of Valdez adult residents are reportedly overweight or obese and about 10% report they do not engage in any form of physical activity. Rural residents are generally less physically active than their urban counterparts and disproportionately affected by chronic diseases associated with insufficient activity. Getting the amount of recommended exercise in rural areas is a challenge where lack of infrastructure, weather dependency, and safety are common barriers.

Methods Used Information was gathered in the context of a clinical immersion in Valdez, Alaska. Patient and provider interviews were conducted at the Valdez Medical Clinic to discover community health concerns and local organizations that are working to counter concerns. An interview at the Department of Parks, Recreation, and Cultural Services (DPRCS) for the City of Valdez detailed available programs, resources, and future proposals. A literature review was then conducted to examine interventions that could build on Valdez’s existing assets.

Summary of Results A literature review identifies that increasing steps per day is an easy way to start and maintain a healthy lifestyle, however this is a challenge during the extended winter months. Step count drops precipitously as snow level increases, daylight decreases, and temperatures drop. Fear of falling is a major environmental barrier for older adults in rural areas. DPRCS has an ongoing feasibility study for a new recreation facility that could include an indoor track if community members show interest. This research highlighting the potential benefits of an indoor walking track has been sent to a manager at the DPRCS. This could potentially be used to advance the current feasibility study.

Conclusions An indoor track, as part of a new recreation facility, could provide a safe, affordable, and accessible place for older adults to exercise year-round. Annual property taxes paid to Valdez by the Trans-Alaska Pipeline could provide an avenue to pay for the infrastructure. This indoor track could combat the tough environmental conditions that make exercise particularly challenging during the Valdez winter.
NOVEL HETEROZYGOUS P3H2 VARIANTS IN A CHILD WITH EARLY-ONSET NON-SYNDROMIC DEGENERATIVE HIGH MYOPIA

1VQ Tang, 1A Egenes, 1M Estrada, 1;2SP Shankar, 1University of California Davis, Sacramento, CA; 2University of California Davis, Davis, CA

Purpose of Study To report on P3H2 variants causing non-syndromic high degenerative myopia guiding precision medical care.

Methods Used Chart review, Next-generation sequencing (NGS) panel, and exome trio genetic testing.

Summary of Results A 3-year-old male with bilateral degenerative high myopia (-12.75 diopter spherical (DS)) and refractive amblyopia diagnosed at 2 years of age, was referred to genomics medicine. He was born full-term with an unremarkable prenatal and birth history. His developmental milestones were normal other than mild articulation difficulty. Family history was significant for myopia of -3.00 DS. Physical exam was normal other than midfacial hypoplasia and depressed nasal bridge. Testing in the connective tissue panel, including Stickler syndrome genes and FBNN1 for Marfan’s syndrome, was negative. Several rare non-syndromic myopia genes have been reported; however, given the lack of NGS panel, an exome trio was ordered. This identified two pathogenic variants in the P3H2 gene, c.1372 C>T, p.Q458* and c.1328del, p.G443Vfs* in trans, confirmed by parental testing. P3H2, also known as LEPRLEL, encodes the Prolol 3-hydroxylase-2 enzyme, which performs the 3-hydroxylation of proline residues in collagen. Tissues from the eye, including the sclera, cornea (type 1), and lens capsule (type 4), are composed of collagen. The abnormal collagen molecule processing may cause weakened collagen in eye tissues. Mutations in the P3H2 gene have been associated with non-syndromic early-onset severe myopia with cataracts, lens instability, vitreoretinal degeneration, and risk of retinal detachment mostly in consanguineous families as homozygous variants. There are no reported skeletal issues or syndromic features. The variants in our proband have not been published in the past, but they have been reported once in ClinVar with different variants considered pathogenic.

Conclusions The degenerative high myopia in our proband is due to pathogenic heterozygous variants in the P3H2 gene, consistent with non-syndromic severe myopia and refractive amblyopia. This molecular diagnosis guides management of potential ocular risks and prevents unnecessary systemic evaluation, such as echocardiograms. It will influence the patient’s future lifestyle choices, such as avoiding contact sports and choosing suitable careers. It also eased parental anxiety and offered more accurate inheritance and recurrence risk counseling for parents and extended family.

#164 IMPROVED ACCESS TO CLINICAL GENETICS SERVICES VIA GENETIC COUNSELOR LED HEARING LOSS CLINIC

M Dutra-Clarke*, S Glover, B Russell. University of California Los Angeles, Los Angeles, CA

Purpose of Study There are over 200 genes associated with non-syndromic and syndromic hearing loss. Knowing the underlying genetic etiology helps to determine prognosis including whether it is expected to be stable vs progressive, isolated, or associated with other organ systems requiring surveillance (e.g. eyes, heart, kidneys), and when planning for cochlear implant. It guides medical management such as early access to developmental services, inform recurrence risks for the family, eligibility for future gene therapy, and avoidance of invasive imaging (e.g. CT scan) and aminoglycosides. Early genetic diagnosis is critical. Due to long wait times to see a clinical geneticist, we developed a genetic counseling (GC) clinic where patients with non-syndromic HL receive genetic counseling and testing by a certified genetic counselor. The purpose of the study was to compare wait times for a consult with a GC vs clinical geneticist and determine the diagnostic yield on genetic testing for apparently non-syndromic HL. The intent was to track productivity, measure the effort allocated for this clinic, and plan adjustments and future needs.

Methods Used We conducted a retrospective chart review of patients with apparently non-syndromic HL seen in the Genetics Division at UCLA between January 2020–September 2021 (ages 2 mos–58 yrs). We separated patients into two groups: (1) seen by a clinical geneticist with a genetic counselor (MD+GC) or (2) seen only by a genetic counselor (GC-only). We compared time from referral to initial visit to return of results. Panel-based genetic testing of 150–224 genes was done in a clinical setting.

Summary of Results There were 25 patients seen by MD+GC and 17 patients seen by GC-only. The average time from referral to initial visit was 80 days for MD+GC (n=22) and 16 days for GC-only (n=17). The average time for testing was 48 days in the MD+GC group (n=25) and 79 days in the GC-only group (n=6), which varied between labs. The average time from referral to return of results was 207 days.
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#165 UNCOMMON NEUROIMAGING FINDINGS IN INBORN ERRORS OF METABOLISM

1JA Morales*, 2FP Velez-Bartolomei, 1M Ruzhnikov, 1GM Enns. 1Stanford University, Stanford, CA; 2San Jorge Children and Women’s Hospital, San Juan

10.1136/jim-2022-WRMC.162

Purpose of Study Inborn errors of metabolism (IEM) are often associated with neurodevelopmental/neurological features such as intellectual disability, autism, epilepsy, hearing loss, hypotonia and movement disorders.

Underlying structural brain abnormalities, like disorders of neuronal migration and/or primarily symmetric signal abnormalities of basal ganglia or white matter, are also common features for certain IEMs.

In this study, we describe two individuals with distinct IEMs: Glutaric acidemia type 1 (GA1) and Congenital disorder of glycosylation type IIs, (CDGII), who presented with intracranial abnormalities not previously or rarely described.

Methods Used Literature & retrospective chart review, clinical evaluations.

Summary of Results 1. An 18-year-old male previously diagnosed with GA1, presented with a first-time generalized tonic-clonic seizure. Comorbidities include spastic quadriaparetic cerebro palsy and lack of verbal speech, likely secondary to past metabolic crises.

Brain MRI showed diffuse cerebral white matter, basal ganglia and brainstem abnormal diffusion. Multiple subependymal nodules along the ventricular margins were also detected.

A comprehensive brain malformations gene panel was not clinically relevant.

2. A 21-year-old male with an underlying diagnosis of CDGIIIs, presented with an episode of left face and arm numbness, and left facial droop at age 15 years. His comorbidities include complex partial epilepsy, hypogammaglobulinemia, beta-thalassemia intermedia, tethered cord, kyphosis, cirrhosis, and portal hypertension. Brain MRI demonstrated severe narrowing of the suprarnoid internal carotid arteries concerning for Moyamoya disease. This was confirmed through cerebral angiogram, and corrected by bilateral cerebral artery bypass grafts.

Conclusions These specific brain imaging findings have rarely been reported in IEMs. Subependymal nodules have been found in four other patients affected by GA1, three of whom were adults. Moyamoya disease is thought to arise from either tunica intima proliferation or abnormal angiogenesis and can be associated with genetic disorders such as Down syndrome or RNF213. While the number of known patients with CDGIIIs is small, none have been identified with Moyamoya disease and we are unaware of other CDG subtypes associated with this.

Given the low incidence of these rare conditions, it is not possible to determine a pathophysiologic link or direct correlation. However, in the absence of a secondary condition that could explain these findings, it is worth considering them as part of the broad phenotypic spectrum for these particular IEMs. Neuroimaging should be part of the diagnostic work-up upon presence of suggestive neurologic symptoms.

#166 PEDIATRICS RESIDENTS AND FUNDAMENTALS OF GENETIC TESTING: SELF-PERCEIVED KNOWLEDGE AND CONFIDENCE

R Gates*, L Hudgins, LC Huffman. Stanford University, Stanford, CA

10.1136/jim-2022-WRMC.163

Purpose of Study Genetic testing is becoming ubiquitous in the field of medicine and is often ordered or requested by primary care providers, non-genetics subspecialists, and patients themselves. Previous studies have demonstrated that primary care providers are often not comfortable ordering genetic testing and counseling families about genetic testing results. There have been initiatives to teach these concepts to practicing physicians via continuing medical education, however, there is no standardized training to teach resident physicians about genetic testing.

Methods Used This was an IRB-exempt study. In September-October 2020, we recruited all active Stanford pediatrics residents via email (n=102). Participant residents completed an investigator-developed Qualtrics electronic survey that addressed self-perceived current level of knowledge about fundamentals of genetic testing, as well as confidence discussing these fundamental issues with families. The survey included 19 items, with Likert scale and open-text response formats. Likert scale responses (Not at All/Slightly and Moderately/Very/Extremely) were combined into two response categories (Insufficiently and Adequately). Chi-Square test of independence was performed using IBM SPSS Statistics software.

Summary of Results Response rate was 46/102 (45%); all training levels were represented (PGY1 33%, PGY2 24%, PGY3+ 42%). Proportions of respondents who were 'Adequately' knowledgeable about fundamentals of genetic testing ranged from 20% to 72%. Smaller proportions of respondents were 'Adequately' confident discussing these concepts with families, ranging from 7% to 65%. Self-reported knowledge and confidence were highest across items for the PGY1 group, with statistically significant differences in proportions of PGY1, PGY2, and PGY3+ describing themselves as 'Adequately' knowledgeable of basic concepts (93%, 73%, and 53%, respectively; p=0.034) and in proportions describing themselves as 'Adequately' confident during discussions of genetic testing limitations (64%, 9%, and 27%, respectively; p=0.045). The majority (89%) of pediatrics residents agreed that a curriculum teaching basics of genetic testing would be helpful to them. Desired curricular topics included:
indicators/limitations of genetic testing, procedure for testing, and counseling families.

Conclusions Despite its increasing importance in medicine, medical genetics education is lacking in pediatrics residency programs. This single-institution educational needs assessment suggests that more advanced pediatrics residents have a greater awareness of deficits in genetics-related knowledge and confidence. Residents agree that they would benefit from a curriculum that teaches fundamentals of genetic testing during training. Future studies should evaluate these themes across multiple institutions and implement a curriculum that assesses pre- and post- intervention knowledge and confidence.

Purpose of Study Neonatal COVID-19 encephalitis is a disease that has been rarely reported during the COVID-19 pandemic. Given the rarity of the disease the signs associated with neonatal COVID-19 encephalitis may easily be confused with other encephalopathies. Here we describe a case of neonatal COVID-19 encephalitis that prompted genome sequencing due to findings on a brain MRI. Genome sequencing discovered a pathogenic variant in PTEN, which is likely unrelated to the COVID-19 encephalitis but has serious lifelong health implications.

Methods Used Chart review.

Summary of Results The patient was a 3 week-old female who was admitted to the PICU for seizures and upper respiratory tract infection signs in the setting of COVID-19 positive contacts at home. An MRI of the head demonstrated patchy white matter degeneration interpreted by radiology as consistent with viral encephalitis or a metabolic disorder. The genetics service was then consulted and determined that the patient likely had COVID-19 encephalitis. However, out of an abundance of caution rapid trio genome sequencing was sent. The genome demonstrated a de novo pathogenic missense variant in PTEN (chr10–89720679 C>T). This variant is well known to be pathogenic of PTEN hamartoma tumor syndrome (PHTS). This variant was determined to be unlikely relevant to her inpatient care and the baby discharged home on Keppra five days later.

Conclusions Rapid genetic testing is a powerful tool for clinical decision support, though incidental findings may complicate genetic counseling and clinical care. There is no evidence supporting pathogenic variants in PTEN increasing the susceptibility to COVID-19 encephalitis. Neonatal onset PHTS would also be unusual and macrocephaly would be expected and the patient was actually microcephalic. The parents opted not to learn about incidental findings, which further complicated the disclosure of results as the PTEN variant was reported by the lab despite the fact that it was an incidental finding unrelated to the patient's current illness. However, the parents were thankful to learn about the PTEN variant given the risks for neurodevelopmental delay and neoplasms associated with PHTS. PHTS is associated with a variety of neoplasms, most of which are adult onset. However, follicular or papillary thyroid cancer can occur as young as seven years of age in PHTS. Thus the patient will follow up with genetics for evaluation of neurodevelopment and thyroid cancer screening.

Poster session
Healthcare delivery research
6:00 PM
Thursday, January 20, 2022

#168 TRUST IN OREGON HEALTH AUTHORITY POLICIES DURING COVID-19

H Graham*, M Gentry, M Jette, E Kim, J Spaan. Western University of Health Sciences College of Osteopathic Medicine of the Pacific-Northwest, Lebanon, OR

Purpose of Study The purpose of this project was to discern trust in the healthcare system during the COVID-19 pandemic in Oregon communities. Our hypothesis was that if a person trusted the health care system, then they would agree with the statements and follow the COVID-19 policies put forth by the Oregon Health Authority (OHA).

Methods Used We measured trust by analyzing when behavior matched opinion in accordance with the research by Lazarus et al, 2019 that said higher trust in a healthcare system can be demonstrated by higher compliance with policies. To assess this, we conducted a survey that asked people about their behavior during the timeframe of April 2020 through November 2020 with a yes or no question. We then asked them to answer if a specific OHA policy would help prevent the spread of COVID-19 on a Likert scale. Our survey was distributed via email and over social media from the timeframe of April 4th, 2021, through April 7th, 2021. We closed our survey on April 7th, 2021, after we realized we had aberrant data most likely due to a computer-generated survey hack that began to occur on April 8th, 2021. After closing the survey, we were left with a total sample size of 124 responses from 12 different Oregon counties aged 18 years and older.

Summary of Results In order to analyze the data, we gave a numerical value to each question response so that the higher values would be awarded to the responses that aligned with the OHA policies. In the policy agreement section, the scores were: strongly disagree = 1, disagree = 2, agree = 3, strongly agree =4. If they answered with a ‘neutral’ regarding their feeling of OHA policies protecting against COVID, they were assigned a score of 0. The lowest scores correlated with people that never followed OHA recommended behavior and who strongly disagreed that the OHA policies would prevent infections. The highest possible score was 38 which represented the most trust in OHA. When analyzing the results, we found our respondents demonstrated a spectrum of trust in OHA with the total lowest score being 13 and our highest score...
being 38 with an average score of 31.4. This average score correlates with people having COVID conscious behaviors and agreeing or strongly agreeing with most of the OHA statements.

**Conclusions** In conclusion, these data points align with our hypothesis because they illustrate an agreement with OHA policies and a behavior that acts in accordance with that policy. Further research should include a larger sample size to better represent the population of Oregon, as well as measures to prevent survey hacking. Results could also be collected at multiple time points as the status of the pandemic is constantly in flux. When applied on a larger scale, this data could help public health officials gauge the trust of the populations they serve and lead to better health outcomes.

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#169 DOES INSURANCE STATUS INFLUENCE MATERNAL MORTALITY: A SCOPING REVIEW

1) Hannan*, 1,2A Bertoni Metoyer. 1University of Washington, Seattle, WA; 2Gonzaga University, Spokane, WA

**Purpose of Study** Maternal mortality in the US is the highest of all developed countries. Change in health insurance policy, such as extension of postpartum Medicaid, is frequently proposed as a possible solution to reduce maternal deaths. The goal of this review is to identify and summarize literature describing association of health insurance status on maternal mortality.

**Methods Used** We performed a scoping search of the databases PubMed, Web of Science Core Collection, Embase, and CINAHL using keywords related to maternal mortality and health insurance. Inclusion criteria required studies to be within the US and use empirical data from 1990 to present. Studies were excluded if they did not compare more than one insurance type or did not examine mortality separately from morbidity. Any numerical data related to mortality was extracted. Search results and articles were screened by two researchers.

**Summary of Results** Our database search yielded 894 results, of which 38 proceeded to full text screening and 17 were included in this review. Our study found that literature directly examining the association of insurance on maternal mortality is not very prevalent. Among the literature that is available, study design is varied. Units of analysis are either state aggregate data or individual mortality data. Additionally, some studies focus only on in-hospital mortality, while others include postpartum deaths within six weeks to one year of delivery. Three included studies examine state level policy, and within aggregated data, generally show increased insurance coverage rates are correlated with lower maternal mortality rates. The remaining 14 included studies use individual mortality data. Only two specifically examine insurance type as a causal factor. The other 12 examine insurance as one of many factors related to mortality and within a subsection of larger analyses, such as investigations of specific causes of maternal deaths or regional mortality case reviews. Two case control studies show a non-statistically significant effect of insurance, and one showed a higher proportion of private insurance among women who died. The remaining nine individual data studies show varying degrees of mostly beneficial correlation between being insured and reduced maternal mortality, with private insurance being more protective than Medicaid when examined. Effect size varies significantly by racial group both in state and individual data.

**Conclusions** Despite significant commentary on the topic, literature examining the effect of health insurance status on maternal mortality is scant. However, state-level research suggests that health insurance access may decrease probability of maternal mortality. This trend requires further examination at the individual level to determine mechanism and degree of effect. Understanding this effect would aid development of future health insurance policy accordingly.

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#170 INTERVENTIONS TO ADDRESS HEALTH DISPARITIES IN COLORECTAL CANCER SCREENING AMONG THE MINORITY POPULATIONS IN THE UNITED STATES

1) Hosseinian*, 1S Afzal, 1E Bolt, 1J Gu, 1D Lee, 1P Pragash, 1S Sacchetto, 1B Afghani. 1UC Irvine school of medicine, Orange, CA; 2Children’s Hospital of Orange County, Orange, CA

**Purpose of Study** Colorectal cancer in underserved or minority populations is associated with increased morbidity and mortality. Factors related to this disparity are complex and multifactorial but one of the reasons is due to lower participation rates in colorectal cancer (CRC) screening. The purpose of this study is to review the interventional studies and compile the best practices to increase CRC in minority populations.

**Methods Used** A literature review was conducted through PubMed, Google Scholar, and Sci-Hub databases using keywords: ‘colorectal cancer’ ‘screening’ ‘minority’ ‘African-American’ ‘Hispanic’ and ‘intervention.’ Inclusion criteria included interventional studies: that included a control group, consisted of a population of at least 50% minority, participants over the age of 50, and published after 2005.

**Summary of Results** A total of 8 studies satisfied our inclusion criteria (see table 1). The majority of the studies showed an improvement in CRC screening rates but the outcomes measured varied, and included fecal occult blood test, sigmoidoscopy, or colonoscopy. Interventions also varied, and included patient navigation in most of the studies and/or physician-level education in a few studies. Therefore, it was difficult to compare the interventions in different studies because of different study designs and outcomes. After intervention, the CRC screening rate ranged from 27% to 94% in different studies. It appears that interventions that were more personalized and required repetitive reminders or contact seemed to be more effective.

**Conclusions** Our literature review suggests interventional strategies are useful in increasing colorectal cancer screening in the minority population. However, even after intervention, CRC screening remained suboptimal. Larger studies are needed to identify barriers at the individual, health provider and community level and measure the impact of targeted intervention to overcome those barriers.
Abstract #170 Table 1 Interventional studies to improve colorectal cancer (CRC) screening among the minority population

<table>
<thead>
<tr>
<th>First author, year published</th>
<th>Control and intervention definition</th>
<th>Total number of subjects (intervention group)</th>
<th>Total number of controls (standard or no intervention)</th>
<th>% Minority: African-Americans and Hispanics</th>
<th>CRC screening completion rate: Controls vs. Intervention, p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Myers, 2007 Control: No intervention</td>
<td>Group 1: N=387</td>
<td>N= 387</td>
<td>58%</td>
<td>Controls: 32.56%</td>
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<tr>
<td></td>
<td>Group 1: Standard letter</td>
<td>Group 2: Standard letter and tailored messages</td>
<td>Group 2: N=386</td>
<td></td>
<td>Group 1: 45.74%</td>
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<tr>
<td></td>
<td>Group 3: Standard, tailored message and phone call</td>
<td></td>
<td>Group 3: N=386</td>
<td></td>
<td>Group 2: 43.78%</td>
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<tr>
<td>Home, 2014 Control: Educational Material</td>
<td>N=578</td>
<td>N=642</td>
<td>&gt;50%</td>
<td>Controls vs intervention: 91% vs 94%, P = 0.04</td>
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<tr>
<td>Intervention: Patient Navigator</td>
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<td>Cole, 2017 Control: Patient Motivational Interview for Blood Pressure</td>
<td>N1=234</td>
<td>N = 238</td>
<td>100%</td>
<td>Control vs intervention 1 vs intervention 2: 8.4% vs 17.5% vs. 17.8, P &lt; 0.01</td>
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<tr>
<td>Intervention1: Patient Navigation</td>
<td>N2=254</td>
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<td></td>
<td>Intervention 2: Patient Navigation and Motivational Interview</td>
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<tr>
<td>DeGroff, 2017 Control: Standard Care</td>
<td>N=429</td>
<td>N=427</td>
<td>&gt;50%</td>
<td>Control vs intervention: 53.2% vs 61.1%, P = 0.02</td>
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<tr>
<td>Intervention: Patient Navigation</td>
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<td>Ford, 2006 Control: Patients did not receive monthly communication from case managers</td>
<td>N=352</td>
<td>N = 351</td>
<td>100%</td>
<td>Control vs intervention: 51.3% vs 68.9%, P = .10</td>
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<tr>
<td>Intervention: Patients received monthly communications from case managers</td>
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<tr>
<td>Basch, 2006 Control: patients were mailed printed materials</td>
<td>N = 226</td>
<td>N = 230</td>
<td>&gt;50%</td>
<td>Control vs Intervention: 6.1% vs 27.0%, 95% CI: 2.6, 7.7</td>
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<tr>
<td>Intervention: Patients received a tailored telephone outreach</td>
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<tr>
<td>Khankari, 2007 Control: baseline screening rate</td>
<td>N = 154</td>
<td>N = 174</td>
<td>&gt;95%</td>
<td>Control vs intervention: 11.5% vs 27.9%, P &lt; .001</td>
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<tr>
<td>Intervention: Mailing screening-eligible patients a physician letter. Physicians were also trained to review health literacy and ‘best practices.’</td>
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<tr>
<td>Friedman, 2007 Control = Rate of CRC screening by residents 6 months prior to education</td>
<td>N = 132</td>
<td>N = 116</td>
<td>100%</td>
<td>Control vs intervention: 26.7% vs 59.1%, P &lt; 0.001</td>
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<tr>
<td>Intervention = Rate of CRC screening by residents 6 months post education</td>
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#171 A COMPREHENSIVE REVIEW OF WOMEN’S QUESTIONS FOLLOWING MISCARRIAGE ON DIFFERENT SOCIAL MEDIA PLATFORMS

1EG Ong*, 1L Davis, 1A Sanchez, 1H Cunel, 1H Stohl, 1A Nelson, 1N Robinson. 1Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA; 2Harbor-UCLA Medical Center, Torrance, CA

10.1136/jim-2022-WRMC.168

Purpose of Study
The purpose of this study was to investigate common questions and the accuracy of advice related to miscarriage that women post on social media to identify any gaps in miscarriage care, counseling and give learners insight into the experience of miscarriage.

Methods
Used Public social media posts from January 1st 2019 to June 30th 2021 were searched using the keywords of ‘miscarriage’, ‘preterm birth’, ‘pregnancy loss’, ‘early pregnancy loss’, ‘early fetal loss’, ‘spontaneous abortion’, ‘spontaneous pregnancy loss’ and '#IHadAMiscarriage'. Only public posts in English and originating from the United States were included. No private identifiable information was collected. The inclusion criteria for posts varied by site: all posts in public Facebook Miscarriage groups, YouTube videos with ≥1000 views, Reddit r/Miscarriage with ≥175 upvotes, Instagram photos with ≥50 likes and Tweets with ≥1 interaction were analyzed for mentions of questions, advice, and conflicts of interest. Each theme in a post was counted. Advice was classified into categories of ‘accurate’ if it was supported by current professional association guidelines, including the American College of Obstetricians and Gynecologists (ACOG), ‘Insufficient evidence’ was advice only supported by published reports, and ‘inaccurate’ if any part was against clinical guidelines or was unsupported.

Summary of Results
For women and families experiencing miscarriage, social media has become a popular outlet and resource for support. 103 posts were identified for common themes. The most common themes identified included: questions on grief (20.4%), blame (19.4%), quality of post-partum counseling (14.6%), and lack of medical support/follow-up (13.6%). 82.9% of advice mentioned the emotional consequences of loss. 17.1% of posts offered medical advice eligible for evaluation, with the most accurate platforms being YouTube (66.6%), Instagram (100%) and Reddit (100%). The least accurate platform was Facebook (33.3%). Twitter yielded the fewest results that met the inclusion criteria.

Conclusions
Many women who post on social media struggle with grief, blame and look for better support from the medical community. The majority of the online community of women who have had miscarriages offer advice related to grief and coping with loss rather than questions regarding medical information. Thus, clinicians may advise women to use social media as a resource to connect with others in their loss and grief. Social media may also be used as an invaluable resource for learners to empathize with women and their families after miscarriage. The ability to better understand the experience and impact of miscarriage will hopefully improve the quality of miscarriage care.
Abstracts

#172 MEDICAL LEARNERS VIEWS ON LEARNER FEEDBACK TO BUILD PROGRAM CAMARADERIE
1R Buller, 2AG Raja*, 3P Poysophon, 1T Allison-Aipa. 1Riverside University Health System, Riverside, CA; 2Western University of Health Sciences, Pomona, CA

Purpose of Study There is little focus on how residents and fellows can give feedback to program faculty in residencies and fellowships in the United States. This study aims to determine if learners feel comfortable giving feedback, the preferred method of feedback, and whether this feedback correlates with a culture of camaraderie between residents and faculty. It has been shown that building camaraderie improves health outcomes for patients and improves physician wellness by creating an overall positive atmosphere.

Methods Used A questionnaire using both multiple choice and open response questions was sent to all 125 residents and fellows at the Riverside University Health System medical program. The survey was composed of questions regarding honesty in giving feedback, preferred feedback method, and if learners thought increased feedback helped build camaraderie. Cronbach’s alpha was calculated to determine internal validity. Descriptive statistics and a correlation coefficient were calculated from survey responses.

Summary of Results From the 26 responses, results showed that 54% of learners felt comfortable giving feedback to their program and 50% of learners preferred individual, anonymous, written feedback as their method of choice. There were strong correlations between the use of feedback and feelings of camaraderie. Learners who were able to provide feedback had a strong correlation with learners who felt camaraderie with their faculty (p<.001). 75% of learners who preferred one-to-one verbal feedback in general also preferred this method to improve camaraderie, compared to 14% of learners who chose group written feedback. When they differed, students chose group verbal feedback to improve camaraderie.

Conclusions Our study shows that allowing residents to give feedback to their mentors can increase camaraderie, which is known to improve outcomes: patient health outcomes and physician wellness. Furthermore, this study provides a look into learners perspective and leads the way for future studies to provide more causal links between learner feedback to physician wellness and patient health.

#173 OPTIONS TO EXPAND HOME HEALTHCARE IN CORDOVA, AK
1J Bramante*, 2University of Washington, Seattle, WA; 2University of Alaska Anchorage, Anchorage, AK

Purpose of Study Patients in Cordova, Alaska lack access to standard home healthcare services such as home health agencies (HHAs) or assisted living facilities (ALFs). Patients with complicated medical issues who are homebound and often do not have robust social support in the community must rely on physician home visits or be admitted to the hospital’s long term care facility. In order for either of the primary care clinics to bill medicare for home visits, a physician must be present, which significantly limits availability of home visits.

Methods Used Patient and provider interviews were conducted at the Cordova Community Medical Center and Ilanka Community Health Center (Ilanka). An asset-based research approach was used to identify other community entities that could potentially impact health for homebound individuals. An interview was conducted with the Cordova Police Department, which provides wellness checks for individuals in the community. A literature review was conducted to evaluate the efficacy of home health intervention programs in addressing needs present in the community such as hospice and wound care. State and national regulatory institutions were then contacted to ascertain the options for expanding Centers for Medicare & Medicaid Service (CMS) coverage of the local primary care centers to include visiting nurse services.

#174 THE IMPACTS OF COVID-19 ON THE SOCIAL PERCEPTIONS OF BREASTFEEDING
1A Mehta*, 2EA Wright, 1AL Nelson, 2H Stohl, 2M Economids. 1Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA; 2University of California Los Angeles David Geffen School of Medicine, Los Angeles, CA

Purpose of Study There is a growing concern amongst physicians that the increased stress of COVID-19 has significantly impacted the number of women choosing to breastfeed. However, not many qualitative studies address whether COVID-19 has impacted breastfeeding perceptions. This study aims to identify hidden challenges women face while breastfeeding and to determine if those changed with the onset of the pandemic.

Methods Used This research was conducted manually across two social media sites, Reddit and TikTok, using the search terms ‘breastfeeding,’ ‘breastisbest,’ and ‘fedinbest.’ Posts with the most comments from pre-pandemic times [Jan 1, 2018 to March 15, 2020] were compared to the posts with the most comments from [March 15, 2020 to June 15, 2020]. Each post and comment was categorized into a theme, and thematic saturation was achieved after three comments mentioned the same theme. In order to establish a theme, there must be at least five posts speaking about that topic.

Summary of Results 248 posts were analyzed with a total of 433 mentions of the selected themes. Fifteen themes were noted across both Reddit and TikTok, and the top 2 themes across both platforms were ‘providing medical advice’ and...
FUNCTION OF SOCIAL MEDIA ON PREMENSTRUAL DYSPHORIC DISORDER

N Poladian*, A Maron, T Ghazarian, AL Nelson, Y Fernandez-Sweeney. Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA

10.1136/jim-2022-WRMIC.172

Purpose of Study Premenstrual Dysphoric Disorder (PMDD) is a common and debilitating disorder that affects reproductive age women. With society’s growing reliance on social media and the emergence of various social platforms, many patients turn to social media outlets for information. This practice also holds true for women seeking information surrounding PMDD, and those investigating their symptoms. This study was designed to provide clinicians with a better understanding of the content, support, as well as wealth and accuracy of information surrounding PMDD found in a popular social media site, Reddit. We hope that this study encourages physicians to engage in deeper conversations with their breastfeeding patients to help them better critically analyze information online but still receive the positive community support they need.

Summary of Results We analyzed a total of 232 posts that fulfilled the inclusion criteria. The most prevalent theme in posts regarding PMDD was expression of feelings (n=116), centered around the topics of depression, anxiety, identity crisis, rejection, and frustration about the lack of support and understanding from both their community and healthcare providers. The second most common theme was sharing of experience-based insight (n=56), which contained advice suggesting varying therapeutics and coping strategies to decrease PMDD symptom severity. While therapeutic suggestions of SSRIs, NSAID’s, and lifestyle modifications have clinical relevance, user misconceptions were noted regarding contraceptive use and hysterectomy for PMDD symptom relief. The third most common theme was relationship complications due to PMDD (n=31). Health-related questions (n=28), such as whether a new or recurring symptom was associated with PMDD, were the least frequent. Analysis of commentary on user posts led to further identification of four themes: offering support, sharing stories/advice, validating one another’s feelings, and showing appreciation for one another’s insights.

Conclusions This study provides clinicians with insight on the interactions between patients, not observed in clinic. Reddit primarily serves for PMDD as a positive space for promoting belonging, and in doing so, may be beneficial to patients. Furthermore, identification of user frustration and discussion pertaining to failed therapies may set realistic expectations of treatment efficacy. Clinicians may inform patients about specific therapeutic misinformation seen online, promoting further open discussion between patient and physician.

INTERNAL MEDICINE EDUCATION FOR MEDICAL TRAINEES IN NAIVASHA, KENYA

1LV Savochka*, 1D Li, 1LJ Onchey, 1D Bosibori, 1J Beste, C Farquhar, 1University of Washington School of Medicine, Seattle, WA; 2Harborview Medical Center, Seattle, WA; 3Naivasha County Referral Hospital, Naivasha, Kenya

10.1136/jim-2022-WRMIC.173

Purpose of Study In Kenya, there has been a decrease in government funding for medical education which has led to outdated educational guidelines to treat common diseases. In this project, the Clinical Education Partnership Initiative (CEPI) between the Naivasha County Referral Hospital (NCRH) and University of Washington (UoW) was used to strengthen the medical education curriculum for internal medicine trainees and pharmacy students from the University of Nairobi (UoN) who are obtaining their medical training at NCRH.

Methods Used The NCRH internal medicine trainees and pharmacy students were surveyed via Zoom on what internal medicine curriculum topics they would like updated over the course of this 8-week long project. The most popular topics were updated using current medical guidelines and recommendations from World Health Organization, UpToDate and Kenya Ministry of Health. The community CEPI partners at the NCRH helped tailor the guidelines to their resource availability. The updated curriculum was then presented at weekly seminars to the medical trainees and was also shared with the medical community to be used as a reference in the future.

Summary of Results A total of six internal medical education curriculums were updated and six seminars were delivered on topics such as asthma, chronic obstructive pulmonary disease, COVID-19, menitis, liver failure, acute coronary syndrome and acute kidney injury. The attendance of each seminar ranged from 5–10 students and included internal medicine residents, as well as pharmacy students. Throughout the project, positive feedback regarding the sessions was given by the site coordinator, as well as the attendees. In the end-of-project survey, 83% of the attendees strongly agreed that they felt more up-to-date with the topics presented and have been using the information presented in clinical settings. Additionally, 63% of
the attendees strongly agreed that they felt more confident in diagnosing and 83% felt more confident treating based on those topics. Participant feedback also revealed that case-based questions and Zoom polling were an effective way to engage attendees.

Conclusions This project successfully updated six outdated internal medicine curriculum didactic sessions for internal medicine trainees and pharmacy students at NCRH. Participants reported to have an improved confidence in diagnosis and treatment on the presented topics. The plan is for future lectures to continue incorporating case-based questions and polling to emphasize teaching points using Zoom. The presentations were also sent to the site coordinator for future use and reference by NCRH trainees.

#177 DEVELOPMENT OF A WHATSAPP-BASED PILOT PROGRAM TO IMPROVE STROKE EDUCATION AND RISK FACTOR REDUCTION IN PATIENTS WITH ARTERIAL HYPERTENSION IN HUALAR, PERU

K Turk*, J Zunt, C Abanto, A Sanchez. University of Washington School of Medicine, Seattle, WA; Instituto Nacional De Ciencias Neurologicas, Lima, Peru; Universidad Peruana Cayetano Heredia, Lima, Peru

Purpose of Study Cerebrovascular disease and stroke are the second-leading cause of death in Peru. Despite the high burden of stroke, Peru lacks a national stroke program, and there is a shortage of neurologists, stroke units, and thrombolytic therapy, particularly in rural areas. It is imperative to reduce patients’ risk factors for stroke, especially hypertension since it is the most significant modifiable risk factor in preventing stroke. WhatsApp, one of the most popular communication platforms in Peru, provides an exciting opportunity to disseminate health information to patients. The aim of this project was to develop materials for a WhatsApp-based pilot program to improve stroke education and risk factor reduction in patients with hypertension living in rural areas of Hualar, a province in the Lima Region of Peru.

Methods Used A stakeholder analysis was conducted to identify key partners whose input was essential in creating this project. Guidelines from the National Institute of Neurologic Sciences (INCN) in Peru and the American Heart Association/ American Stroke Association were used to create infographics addressing stroke awareness and risk reduction. An informative outline of the pilot program was created for Hualar Hospital partners. A cohort of patients with hypertension was identified by the Hualar Hospital Teaching and Training Unit Director. Surveys addressing patients’ stroke knowledge and comfort using WhatsApp were generated. To ensure content was medically accurate and culturally appropriate, all materials were first edited by a medical student in Peru and subsequently by Peru’s Chief of the Center for Research in Cerebrovascular Disease at the INCN.

Summary of Results Materials created for the pilot program include a detailed program outline, an informed consent adapted to WhatsApp, pre- and post-program surveys, and five educational infographics. These materials are ready to be sent to patients with hypertension in Hualar via WhatsApp. There is enthusiasm and optimism from hospital staff about the potential impact of the pilot program.

Conclusions This project will increase access to accurate stroke education for patients with hypertension living in Hualar. Now that the materials have been created, the next step is to pilot their use in the selected cohort of patients. The program’s success will be determined by comparing participants’ pre- and post-program survey responses. By using WhatsApp, this program could provide a sustainable means to improve stroke education while limiting travel and financial burdens for patients living in rural areas. If successful, this program could be replicated in other rural areas in Peru.

Poster session
Infectious diseases
6:00 PM
Thursday, January 20, 2022

#178 A CASE OF PSEUDOMONAS AERUGINOSA ASSOCIATED DIARRHEA IN A LONG-TERM HOSPITALIZED PATIENT

C Besmanos*, N Raza, H Lai, S Mishra, S Ragland, A Heidari. Kern Medical Center, Bakersfield, CA

Purpose of Study Pseudomonas aeruginosa associated diarrheal disease is not common in adults and if seen, is mostly reported in pediatric population. It has been classified into Shanghai fever, enterocolitis and antibiotics associated diarrhea in pediatric literature. In adults, immunocompromising conditions such as malignancies, neutropenia, and admission to long-term care and intensive care unit (ICU) are known risk factors. Here we describe a case of Pseudomonas aeruginosa associated diarrhea in a long-term hospitalized patient who had rectal tube.

Methods Used Retrospective case review

Summary of Results A 33-year-old Caucasian man with a history of alcohol use disorder, hypertension, and hypothyroidism presented with myxedema coma with TSH 148 requiring intubation. He had a complicated hospital course with ventilator-associated pneumonia with MRSA, sepsis with candida, and abdominal compartment syndrome requiring decompressive laparotomy. The patient slowly recovered despite 43 days of hospitalization. Before leaving his 31 days of admission to ICU, a flexi-seal rectal tube was placed due to fecal incontinence. Five days later when he was already transferred out of the ICU, he started having loose watery stools with leukocytosis and left shift. Clostridioides difficile (C.diff) colitis was suspected and he was placed on oral vancomycin empirically. His stool test for C.diff came back negative, and his rectal tube was subsequently removed. Imaging did not show any abscess, perforated viscus, or fistula formations. His stool culture grew heavy colony numbers of Pseudomonas aeruginosa. Vancomycin was stopped and he was started on oral ciprofloxacin 750 mg twice a day with complete resolution of his diarrhea and leukocytosis.

Conclusions Pseudomonas aeruginosa associated diarrheal disease is not common in adults. Complicated prolonged hospitalization, administration of broad-spectrum antibiotics, and immunocompromising conditions perhaps play role in colonization and eventually infection in the right setting. The role of rectal tube is unknown.
HHV-8 ASSOCIATED MULTICENTRIC CASTLEMAN DISEASE: A DIAGNOSTIC CHALLENGE IN A PATIENT WITH ACQUIRED IMMUNODEFICIENCY SYNDROME AND FEVER

R Dunn*, F Verter, R Janiwal, S Mishra, J Bhandohal, E Cobos, A Heidari. Kern Medical Center, Bakersfield, CA
10.1136/jim-2022-WRMC.176

Purpose of Study HHV-8 associated Multicentric Castleman Disease (MCD) is an angiofollicular lymphoproliferative disorder that affects multiple regions of lymph nodes simultaneously. Incidence of Castleman Disease is about 6500 to 7700 in the United States, 75% of them being associated with HHV-8. All cases of MCD in HIV patients are HHV-8 associated. Common symptoms include fever, lymphadenopathy, Hepatosplenomegaly, pulmonary involvement, edema, and ascites. Here we describe a case of HIV with fever of unknown origin in that after extensive work up diagnosed with MCD.

Methods Used This is a single case review after IRB approval.

Summary of Results A 28 year old male with history of HIV, non-alcoholism with antiretroviral therapy (ART) presents with constant abdominal pain, vomiting, and fevers over 5 days. CBC showed pancytopenia with a CD4 count of 27 cells/mcL and an HIV viral load of 95,200 copies/mL. Work up for Syphilis, Gonorrhea, Chlamydia, Hepatitis A/B/C, Cocci, Cryptococcus, TB, Toxoplasma, Histoplasmosis, Bartonella, Giardia, Brucella, CMV PCR, Parvo B19 PCR, Coxiella burnetii, Cryptosporidium and all of his cultures were negative. Bone marrow biopsy and culture were negative. Imaging showed diffuse lymphadenopathy in the mediastinum, hilar, axillary, retroperitoneum, iliac and inguinal lymph nodes. IR Lymph node excision and HHV-8 staining confirmed HHV-8 associated MCD with serum HHV-8 levels of 2,288,277 copies/mL. Patient was given Rituximab infusions and started on ART.

Conclusions Diagnosis of HHV-8 Multicentric Castleman Disease in HIV host could be challenging and usually is delayed. Starting antiretroviral and Rituximab is the most agreed upon therapeutic modality. This approach almost doubles the survival rate at 2 years from 42% to 90%.

ELEVATED ADENOSINE DEAMINASE LEVELS IN A CASE OF COCCIDIOIDOMYCOsis

C Salib*, L Moosavi, K Radicic. Kern Medical Center, Bakersfield, CA
10.1136/jim-2022-WRMC.177

Purpose of Study Elevated adenosine deaminase (ADA) levels have been a useful diagnostic tool associated with tuberculosis (TB). Diagnostic testing for Coccidiiodymycosis (Cocci) involves serologic testing of IgM and IgG antibodies. Though there are a variety of diagnostic testing options for TB, in a case of high clinical suspicion, pleural fluid analysis for ADA is often completed as an additional confirmatory test, due to its high sensitivity and specificity. This case demonstrates a patient who presented with elevated ADA levels despite having negative sputum cultures for TB and with elevated serological titers for Cocci.

Methods Used A single patient case report was conducted after IRB approval.

Case Presentation A 26-year-old male with history of uncontrolled Type 1 diabetes presented to the ED with shortness of breath and productive cough of two day duration. The patient denied history of travel and reported a nonspecific amount of weight loss over the past few months. He presented febrile, tachycardic, and hypoxic. CT scan found a lung abscess and empyema. He was seen by Interventional Radiology for placement of a pigtail catheter for drainage of pleural fluid. He was initially treated with Fluconazole but then transitioned to Ambisome.

During his admission, patient remained consistently tachycardic and tachypneic with worsening hypoxia. Requiring ICU admission, he was intubated three times and was subsequently extubated within a few days. The patient underwent bronchoscopy and was found to have pleural ADA level of 468. Despite having negative Quantiferon results, the patient was started on RIPE therapy due to suspected TB infection.

Results from Mycobacterial sputum culture returned negative weeks after patient was discharged from the hospital, confirming that the patient did not have an active TB infection. However, serologic test was confirmatory for Cocci.

Conclusions ADA levels have been known to have both a high sensitivity and specificity for diagnosing TB, which makes it an ideal diagnostic tool. In this instance, there was the unusual presentation of increased levels of ADA along with a negative AFP culture. The role and expected actions of the ADA enzyme can be indicative of T-lymphocyte activity. When thought of more generally, this diagnostic tool can be used to diagnose disease conditions other than TB, such as Cocci.

HYDROXYCHLOROQUINE PRE-EXPOSURE PROPHYLAXIS FOR COVID-19 IN HEALTHCARE WORKERS FROM INDIA: A META-ANALYSIS

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Purpose of Study While vaccines have taken center stage in the battle against COVID-19, alternate approaches for prevention of SARS-CoV-2 infection have been ignored. Hydroxychloroquine pre-exposure prophylaxis (HCQ PrEP) has been used to prevent COVID-19 in high-risk healthcare workers (HCWs) in a number of studies from India. We have performed a meta-analysis to evaluate the safety and efficacy of a standard HCQ PrEP regimen in HCWs enrolled in these studies.

Methods Used We performed a search of the medical literature based on the PRISMA checklist (http://www.prisma-statement.org/) using PubMed, Google Scholar, medRxiv and ResearchGate to obtain all relevant publications and preprints. Using this method we obtained eleven nonrandomized controlled trials of weekly HCQ PrEP involving 7,616 HCWs in India (3,489 treated, 4,127 controls). The HCQ PrEP regimen consisted of an 800mg loading dose in the first week, then 400mg weekly thereafter according to the guidelines of the Indian Council of Medical Research. SARS-CoV-2 infection was documented by seroconversion or positive polymerase chain reaction (PCR) testing. We used random-effects meta-analysis to summarize the risk ratio (RR) of infection across the studies.

Summary of Results Sex distribution was available for nine studies and age distribution was available for eight studies.
A UNIQUE CASE OF COVID-19 PRESENTED AS FOCAL SEIZURES WITH IMPAIRED AWARENESS

R Sharma*, S Ratnayake, H Lai, S Mishra, A Heidari. Kern Medical Center, Bakersfield, CA

Purpose of Study Severe acute respiratory syndrome coronavirus 2 (SARS-CoV2) has rapidly become a global pandemic with millions of confirmed cases worldwide. Encephalitis and seizure associated with COVID-19 has been seen and reported. Here described is a unique case of SARS-CoV2 infection presented with focal seizure with impaired awareness.

Methods Used A retrospective review following IRB approval.

Summary of Results A 54-year-old man unvaccinated for COVID-19 with no known past medical history presented to the emergency department with altered mental status. Three days prior to presentation he complained of frontal headaches and blury vision and one day prior his son noticed he was unable to speak. Minutes upon arrival he suffered a focal seizure for which levetiracetam was administered. He was found to be oriented to self and age with difficulty finding words answering only with ‘yes’. EEG was performed and he was diagnosed with focal seizure with impaired awareness. His chest x-ray showed multifocal bilateral hazy infiltrations. SARS-CoV2 PCR test came back positive. His brain MRI revealed 2 acute subcortical superior right frontal lobe lacunar infarcts. His lumbar puncture was negative. He suffered nine seizures of left occipital origin despite being treated with levetiracetam and valproic acid. Phenytoin was added which eventually controlled his seizure and he became fully oriented.

Overcoming a tumultuous hospital course, he was also found to have newly diagnosed poorly controlled diabetes and pulmonary coccidioidomycosis. He was discharged stably on hospital day 16.

Conclusions Focal seizure with impaired awareness associated with SARS-CoV2 has not been reported. Further studies are warranted to understand the pathophysiology and definitive treatment.
uncontrolled diabetic presenting with hemoptysis and mycetoma.

Methods A single patient case report was conducted after approval from IRB.

Case Presentation A 48-year-old Hispanic male with diabetes, untreated pulmonary coccidioidomycosis, and history of COVID-19 infection one year prior presented with sudden hemoptysis, night sweats, and a 45-lb weight loss. He was diagnosed with coccidioidomycosis seven years prior but did not start treatment. Three years prior he was admitted elsewhere with shortness of breath. Serum coccidiodal complement fixation titer was 1:32. Imaging found a right lower lobe cavity measuring 4 x 3 cm with right sided pneumothorax and bronchopulmonary fistula. He underwent video-assisted thoracotomy with pleurodesis and treated with fluconazole for 4 months, improving his titers to 1:4. He was then lost to follow.

Upon presentation to our facility with worsening hemoptysis, he had a larger cavity measuring 10 x 8 x 7 cm in the right lower lobe with central filling mass. He hemoptysized daily while admitted. Bronchoscopy confirmed coccidioidal mycetoma by direct stain and fungal cultures. Interventional radiology performed arterial embolization of right tracheobronchial and intercostal bronchial arteries to control hemoptysis. He was restarted on fluconazole and discharged home.

One week later he returned in respiratory distress with fever, shortness of breath, and hypoxemia. Imaging revealed new left lower lobe and unilateral consolidations. He was thought to have aspiration of right sided cavitary material to the left lung. His oxygen requirements increased significantly, concerning for severe pulmonary coccidioidomycosis and he was placed on steroids and liposomal amphotericin B. His symptoms improved and steroid was stopped. His antifungal was switched to Posaconazole and was able to go home on room air.

Discussion Management of giant pulmonary cavitary coccidioidomycosis with mycetoma and bleeding is challenging. Poorly controlled diabetics are already at increased risk for cavitation. In a patient presenting with hemoptysis and a cavitary lesion, a multidisciplinary team consisting of pulmonology, infectious diseases, thoracic surgery and interventional radiology is essential.

#185 A CASE OF PSEUDOTERRANOVA, HAVING CEVICHE WITH SPECIAL FLAVOR

Prasad, C’; McPheeters, R’; Stull, A’; Heidari, A’; Adtakem Global Education Inc, Downers Grove, IL; Kern Medical Center, Bakersfield, CA

Methods Used Retrospective case study.

Summary of Results A 17-year-old male with no significant medical history presented to our hospital after he coughed up a worm earlier that morning. He also complained of ongoing rhinorrhea and sore throat for the past four days. He denied nausea, vomiting, rash, diarrhea, fever, chills, night sweats, hematochezia, hematemesis, abdominal pain, abdominal bloating, headache, weight loss, or change in appetite. Patient stated that he was from Mexico but had been living in the United States for the past two years. A dietary history revealed that one week ago he had eaten his favorite ceviche made from fresh fish brought by a family member visiting from Ensenada, Mexico. Patient’s physical examination, lab values, and imaging were all unremarkable. He had brought the worm, which he had coughed up, to the hospital. The worm was sent to the pathology lab and identified as Pseudoterranova species. Patient was discharged with instructions to return if symptomatic. He was referred for follow-up in the outpatient setting.

Conclusion Anisakiasis is rare with current United States food handling regulations. The clinical suspicion is raised when raw fish from alternative sources of fresh seafood is consumed. Removal of the worm via endoscopy or even surgery might be necessary and is considered therapeutic. Diagnosis is made by direct visualization of the nematode.

#186 MANDIBULAR OSTEOMYELITIS DUE TO AGGREGATIBACTER ACTINOMYCETECOMITANS

Ratnayake*, H’; Sidhu, CD’; Petersen, G’; Heidari, A’; Medical Center, Bakersfield, CA; Bass University School of Medicine, Miramar, FL

Case Report Aggregatibacter actinomycetcomitans is frequently associated with localized aggressive periodontitis. A. actinomycetcomitans is a Gram-negative facultative anaerobe that is a member of the HACEK group of fastidious Gram-negative bacteria that can rarely cause endocarditis. We report a 21-year-old Hispanic male with osteonecrosis of the mandible from a tooth infected with A. actinomycetcomitans requiring antibiotics and surgical intervention.

Case Description A 21-year-old Hispanic male with no known past medical history presented to the emergency department with a purulent right jaw and neck abscess. Two weeks prior he began having right lower tooth pain. Over the next 10 days his pain and swelling progressed, limiting his ability to open his mouth and to eat and drink. He then noticed an enlarging mass over the right jaw and neck with purulent discharge. Intolerable pain brought him to the ED. Admission CT of the soft tissue of the neck found right mandibular angle and ramus osteomyelitis, adjacent masticator and sternocleidomastoid infectious myositis, reactive right parotiditis, and severe right cervical cellulitis. There were also small periapical abscesses of the right mandibular first molar, and small pockets of localized edema in the right suprathyroid neck without organization and right upper cervical adenopathy. He was started on vancomycin and piperacillin-tazobactam. Ear nose and throat surgeon aspirated the abscess. Aspirated abscess culture grew A. actinomycetcomitans. Antibiotics were narrowed to ceftriaxone and metronidazole. ENT then performed incision and drainage of the right jaw and neck abscess and extracted two infected teeth. He was
successfully discharged on post-operative day one on moxi-floxacin with goal of 6 weeks of therapy with close ENT follow up.

Conclusion Early recognition and treatment of periodontal infections is important to prevent complications such as abscess formation and osteonecrosis from osteomyelitis. We report a prototypical example of an acute progression of a simple tooth ache developing into severe osteonecrosis by a rare HACEK organism not commonly encountered requiring aggressive antibiotics and surgical management.

#187 MALLET FINGER COMPPLICATED BY DISTAL PHALANX OSTEOMYELITIS

D Deng*, N Hatchard, R Patel. Western University of Health Sciences College of Osteopathic Medicine of the Pacific-Northwest, Lebanon, OR; Philadelphia College of Osteopathic Medicine, Philadelphia, PA; Inspira Health Network, Vineland, NJ

Case Report A 24-year-old male presented with a right index finger injury which occurred during a soccer game. The mechanism of injury was the ball hitting the top of the second digit and lacerating the most distal portion of the digit. A diagnosis of mallet finger was made, and later on osteomyelitis of the distal phalanx.

The patient was initially evaluated in the emergency room where he was diagnosed with mallet finger and placed in a distal interphalangeal (DIP) extension QuickCast splint. A follow-up visit revealed five degree DIP flexion posture and ten degree proximal interphalangeal extension posture. DIP active range of motion flexion was noted at 30 degrees and only 45 degrees and painful for PIP active and passive range of motion. As X-ray of the right index finger revealed no evidence of fracture, dislocation, or degenerative changes, surgical intervention was not indicated at the time. However, patient was showing persistent edema, erythema, and pain 8 weeks after the initial accident. Follow-up MRI imaging identified fluid collection in the DIP joint. With suspicion for infectious process, surgical debridement of the finger was performed which revealed necrotizing tissue at the mid dorsal metaphysis of the distal phalanx spreading through a crevice to the germinal matrix. This impacted the patient’s recovery and rehabilitation of the joint along with the structural and functional outcome. The patient was ultimately diagnosed through MRI and bone biopsy and found to have Enterobacteriaceae skeletal osteomyelitis of the distal interphalangeal joint. The patient was treated with oral ciprofloxacin post-surgical debridement and was able to fully recover from the osteomyelitis infection. However, continued and permanent droop of the DIP with inability to regain full ROM, strength, and utility in the right index phalanx was seen during the patient’s follow-up appointment.

Our aim with this case study is to demonstrate the possible complications of open mallet finger injury and its rehabilitation considerations. In this case, the patient’s lacerated mallet finger led to osteomyelitis of the digit. While mallet finger itself is a very common injury, its recovery is rarely complicated by infection in previously healthy individuals.

This case highlights the importance of considering bone infection in non-healing joint injuries such as mallet finger and subsequently appropriately adjusting therapies to include prophylactic antibiotics and splinting to facilitate the structural and functional rehabilitation.

#188 PREAXIAL POLYDACTYLY: CHARACTERIZING SONIC HEDGEHOG REGULATION

M Malone*, K Ball, C Pira, K Oberg. Loma Linda University School of Medicine, Loma Linda, CA; Loma Linda University Departures of Basic Sciences, Loma Linda, CA; Loma Linda University, Loma Linda, CA

Purpose of Study Preaxial polydactyly is associated with ectopic Sonic hedgehog (Shh) expression in the presumptive thumb. Shh is secreted from the zone of polarizing activity (ZPA), the signaling center that directs limb development along the radial-ulnar (anterior-posterior) axis. Shh is necessary for development of the ulna and posterior four digits, while the thumb develops in the absence of Shh. A limb-specific enhancer, the ZPA regulatory sequence (ZRS), is necessary for Shh expression, although little is known about how the ZRS restricts Shh expression to the ZPA. Single nucleotide variations (SNVs) within the ZRS cause ectopic Shh expression resulting in preaxial polydactyly. These SNVs are spread across three highly conserved regions: peaks 1, 2, and 3. We hypothesize that one or more of these peaks is necessary for ZRS activity.

Methods Used Plasmids containing full-length ZRS, or each peak respectively were electroporated into Hamburger-Hamilton stage 14 chicken embryo presumptive limb buds. The embryos were incubated for 48 hours then observed by fluorescence microscopy. Fluorescent activity of each peak was then compared to that of full-length ZRS.

Summary of Results Constructs containing peak 1 and peak 2 have no detectable activity in the limb. Peak 3, however, maintains some activity, but markedly less than full-length ZRS.

Conclusions Peak 3 is necessary and sufficient for activity while peaks 1 and 2 are not, suggesting peak 3 contains core sequences necessary for ZRS activity. It is surprising that peaks 1 and 2 lack activity as many clinically relevant SNVs occur in these two regions and the reported Hand2 binding site is in peak 2. Since peak 3 activity is less than that of full-length ZRS, peaks 1 and 2 likely enhance the activity of peak 3. Further studies will assess the quantitative difference in ZRS peak activity and determine how the ZRS localizes Shh expression to the ZPA. Mapping the ZRS will clarify the role of critical sequences in Shh regulation that contribute to limb development and malformation.

#189 VARIATIONS IN THE PERONEUS TERTIUS MUSCLE: EVOLUTIONARY AND CLINICAL PERSPECTIVES

K Kay*, J Scott, M Wedel. Western University of Health Sciences, Pomona, CA

Purpose of Study Peroneus tertius (PT) is a muscle in the anterior compartment of the leg that functions in dorsiflexion and eversion of the foot at the ankle. PT was long thought to be unique to humans, but it is now known to be variably present...
in many other primates, including bonobos, chimpanzees, gorillas, and several species of monkeys. However, there is still disagreement between anthropologists and comparative anatomists regarding the evolutionary origin of the muscle. PT is highly variable in origin, insertion, size, and number of musculotendinous slips. Previously published studies have also reported PT to be extremely variable in terms of prevalence, ranging between 38% and 100% in different populations. Our goals are to summarize the evolutionary origin, anatomical variations, and clinical implications of the muscle.

Methods Used We synthesized information from the literature on the evolution, prevalence, variability, and clinical correlations of the PT using Google Scholar, NCBI, and ResearchGate. For statistical analysis, we measured prevalence using the number of limbs with PT present per number of legs total. In particular, we compared studies that observed the muscle in human cadavers (n=11) versus studies that used palpation on living subjects (n=11).

Summary of Results Numerous variants of PT have been identified, including an origin from the extensor hallucis longus. The muscle is occasionally doubled, in both humans and gorillas. Peroneus tertius is associated with multiple causes of ankle laxity, and can be used in surgery for repair of other tendons. We found that the prevalence of PT varies depending on the method of data collection. Studies that use cadaveric dissection consistently report a significantly (p < 0.0001) higher prevalence of PT (86–100%) than studies that rely on palpation of the muscle in living subjects (38–84%).

Conclusions Although the higher prevalence of PT in humans than in other primates has been linked to the evolution of bipedality, PT is not crucial for an erect bipedal gait. Recent studies have found no significant difference in ROM or in the strength of dorsiflexion or eversion among patients with and without PT. PT has been used in tendo-capsular, tendon transfer, resection surgeries, transposition to correct ankle laxity, and transplantation surgeries for foot drop. The significant difference between dissection and palpation studies in the reported prevalence of PT suggests that palpation consistently understimates the true prevalence of the muscle. We suggest that the variability in the size and course of PT, as well as with the potential for a small PT to be difficult to palpate through the extensor retinaculum, combine to make the muscle difficult to diagnose via palpation. Although palpation studies are potentially faster and less expensive than dissection-based studies, we urge caution in interpreting their results.

Poster session
Neonatology general
6:00 PM
Thursday, January 20, 2022

#190 GENETIC ASSOCIATIONS WITH PREECLAMPSIA, INTRAUTERINE GROWTH RESTRICTION, AND SPONTANEOUS PRETERM BIRTH

AL Baranoff*, 1, 2A Paquette, 1Seattle Children’s Research Institute, Seattle, WA; 2University of Washington School of Medicine, Seattle, WA

10.1136/ijim-2022-WRMC.187

Purpose of Study Pregnancy-related conditions are attributable to a combination of factors and may increase one’s risk for later disease. Three of the most common conditions are preeclampsia (PE), intrauterine growth restriction (IUGR), and spontaneous preterm birth/delivery (SPTB/D). The genetic basis of these conditions is illustrated through the findings that women who experience SPTB/D are at increased risk of subsequent SPTB/D of the same gestational age, and women who experienced PE or IUGR are also more likely to experience these same complications in subsequent pregnancies. We investigated the genetic profiles of each outcome and analyzed which maternal and fetal genomic variants are most associated with multiple pregnancy complications. This knowledge will increase our ability to better treat and/or prevent adverse pregnancy outcomes by identifying persons who may be at higher risk of complications prior to their manifestation in disease.

Methods Used We conducted a literature review of studies that assessed the association between fetal and/or maternally-based single nucleotide polymorphisms (SNPs) and PE, IUGR, and SPTB/D. We selected genetic variants that were significantly associated with these pregnancy outcomes in an initial candidate gene study, genome-wide association study (GWAS), or summarized in a meta-analysis (MA). The significance of association between SNP and pregnancy outcome was reported at either a study-wide significance level (for GWAS and MA studies) or p < 0.05 for candidate gene studies.

Summary of Results Our search yielded 103 articles, of which 59 investigated genetic associations in PE, 27 in IUGR, and 18 in SPTB. We identified five variants found in both PE and IUGR: rs1918975, rs10774624, rs3184504, rs4769612, rs1884082 which were in regulatory regions of four genes: FLT1, SERPINA3, MECOM, and SH2B3. We identified no variants found in both IUGR and SPTB. We identified five genes that contained SNPs in both IUGR and SPTB (ADCY5, WNT4, IGF1R, EBF1, IGF1), although none of the SNPs overlapped. No genetic variations or gene profiles were found to be shared amongst all adverse pregnancy outcomes.

Conclusions We identified five genetic variants associated with both PE and IUGR which correlated to four genes as well as five genes associated with both IUGR and SPTB. Several of these gene variants are also risk factors for the development of diseases that impact health throughout life, such as cardiovascular and kidney disease as well as neurological delay, revealing that adverse pregnancy outcomes and adult disease have shared and complex genetic risk factors. The contribution of the identified shared genetic variants in the pathogenesis of PE, IUGR, and SPTB/D should be the focus of future studies.

#191 FACTORS ASSOCIATED WITH HYPOTERMIA AND HYPOGLYCEMIA ON ADMISSION IN VERY LOW BIRTH WEIGHT INFANTS

C.Marquez*, K. Ram, Y. Shao, NS Nanduri, A. Hisey, L. Barton, R. Ramanathan, M. Biniwale.

LAC + USC Medical Center, Los Angeles, CA

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Purpose of Study Maintaining temperature and glucose in preterm infants are vital as these abnormalities can predispose them to many undesirable complications in early neonatal period. The present study was conducted to identify the

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Abstracts

Abstract #191

<table>
<thead>
<tr>
<th>Birth weight (g)</th>
<th>Hypothermia on admission</th>
<th>Normothermia on admission</th>
<th>P value</th>
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</thead>
<tbody>
<tr>
<td>942</td>
<td>991</td>
<td>0.116</td>
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</tr>
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</table>

Factors associated with hypothermia and hypoglycemia in very low birth weight (VLBW) preterm infants.

Methods Used The data consisting of VLBW infants born at our hospital was collected retrospectively from 2009 through 2021. Maternal factors, delivery room events and early neonatal morbidities were analyzed against infants’ first temperature and glucose done on admission to NICU. Hypothermia was defined as temperature <36.5°C. Hypoglycemia was defined as blood glucose <45 mg/dl on admission checked by point of care testing. IRB approval was obtained to review the data from electronic medical records. SPSS version 28 statistical software was used to analyze the data.

Summary of Results From all VLBW infants born during this period 152/642 (23.7%) had temperature below 36.5°C on admission to NICU while 104/652 (15.9%) were diagnosed to have hypoglycemia. Birth weight or gestational age had no impact on either hypothermia or hypoglycemia on NICU admission. Infants needing resuscitation including chest compressions and epinephrine administration were at highest risk for hypothermia. These infants were also noted to have metabolic acidosis and low 5 min apgar scores. Infants presented with hypoglycemia were small for gestational age (57%) vs 24% p=0.009. Maternal medical conditions including diabetes did not put these infants at additional risk for hypoglycemia. Hypoglycemia on admission was also associated with additional risk of requiring higher ventilation as well as oxygen requirement in the first 24 hours of NICU stay.

Conclusions VLBW Infants needing resuscitation in the delivery room are at risk for hypothermia. These infants may present with metabolic acidosis on admission to NICU. Hypoglycemia on admission may predispose VLBW infants for more respiratory support.

### 192 THE EFFECT OF SARS-COV-2 ON THE RATES OF BREASTFEEDING IN THE NEWBORN NURSEY

1 Wang*, 1AF Ahmed, 2R Ramanathan, 3A Yeh. 1LAC+USC Medical CenterUCSC, Los Angeles, CA, 2LAC+USC Medical Center, Keck School of Medicine of USC, LA, CA

Purpose of Study Exclusive breastfeeding for the first six months of life is recommended by the American Academy of Pediatrics and the Centers for Disease Control for its benefits to infant immunity, maternal-child bonding, and long-term health. While these benefits are well studied, the SARS-CoV-2 pandemic raises questions about the safety of breastfeeding among SARS-CoV-2-positive mothers. In addition, the pandemic’s effects on hospital staffing, patient-provider face-time, and healthcare access may impact breastfeeding rates. This study aims to explore the effect of the SARS-CoV-2 pandemic on breastfeeding in the newborn nursery.

Methods Used This is a retrospective cohort study comparing breastfeeding rates between neonates at LAC+USC Medical Center Newborn Nursery from January 2019 to April 2021. We defined the pre-SARS-CoV-2 group as all neonates born prior to April 2020, and the during-SARS-CoV-2 group as those born from April 2020 to April 2021. Maternal data gathered included gravidity and parity, ethnicity, age, mode of delivery, and pregnancy complications. Infant data gathered included gestational age, birth weight, sex, and hyperbilirubinemia requiring intensive phototherapy. Neonates with maternal contraindications to breastfeeding, such as positive toxocology screen, positive HIV status, incarceration, and placement in foster care were excluded. Rates of exclusive breastfeeding and any breastfeeding were calculated for each month within this time period and compared using T-test. P value less than 0.05 was considered significant.

Summary of Results Of the 964 newborns screened in the pre-SARS-CoV-2 cohort, 913 were included. Of the 800 screened during-SARS-CoV-2 cohort, 763 newborns were included. There were no significant differences in the demographics between the two cohorts (table 1). We found a 11% decrease in the rate of exclusive breastfeeding (p<0.05) and a 4% decrease in any breastfeeding (p<0.05) during the SARS-CoV-2 period (Image 1).

Conclusions The SARS-CoV-2 pandemic had a negative impact on the rates of both exclusive breastfeeding and any

### Abstract #192 Table 1 Maternal and neonatal demographics and characteristics

<table>
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<th>Study Subjects</th>
<th>Pre-SARS-CoV2</th>
<th>During-SARS-CoV2</th>
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<tbody>
<tr>
<td>Total Newborns Born</td>
<td>964</td>
<td>800</td>
</tr>
<tr>
<td>Total Newborns Included In This Study</td>
<td>913</td>
<td>763</td>
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<td>Exclusion Criteria</td>
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<tr>
<td>Drug Screen Positive%</td>
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<td>Maternal Incarceration%</td>
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<td>DCF Case%</td>
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<td>Baby Demographics</td>
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<tr>
<td>Hispanic or Latino%</td>
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<td>Average Maternal Para</td>
<td>2.2</td>
<td>2.3</td>
</tr>
<tr>
<td>Maternal Preeclampsia%</td>
<td>4.4</td>
<td>6.0</td>
</tr>
<tr>
<td>Infant of Diabetic Mother%</td>
<td>10.5</td>
<td>13.1</td>
</tr>
<tr>
<td>Baby Characteristics</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Average Baby Gestational Age</td>
<td>38.9</td>
<td>38.9</td>
</tr>
<tr>
<td>Preterm Baby%</td>
<td>8</td>
<td>8.4</td>
</tr>
<tr>
<td>Birthweight &lt; 2.5 kg%</td>
<td>4.1</td>
<td>4.1</td>
</tr>
<tr>
<td>Birthweight &gt; 4kg%</td>
<td>7.1</td>
<td>6.2</td>
</tr>
<tr>
<td>Baby Underwent Intensive Phototherapy%</td>
<td>10.3</td>
<td>10.5</td>
</tr>
<tr>
<td>Breastfeeding Rate</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Exclusive Breastfeeding Rate%</td>
<td>73.2</td>
<td>62.4</td>
</tr>
<tr>
<td>Any Breastfeeding Rate%</td>
<td>94.6</td>
<td>91.2</td>
</tr>
</tbody>
</table>

Abstracts

Abstract #191 Table 1

<table>
<thead>
<tr>
<th>Birth weight (g)</th>
<th>Hypothermia on admission</th>
<th>Normothermia on admission</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>942</td>
<td>991</td>
<td>0.116</td>
<td></td>
</tr>
</tbody>
</table>

Factors associated with hypothermia and hypoglycemia in very low birth weight (VLBW) preterm infants.

Methods Used The data consisting of VLBW infants born at our hospital was collected retrospectively from 2009 through 2021. Maternal factors, delivery room events and early neonatal morbidities were analyzed against infants’ first temperature and glucose done on admission to NICU. Hypothermia was defined as temperature <36.5°C. Hypoglycemia was defined as blood glucose <45 mg/dl on admission checked by point of care testing. IRB approval was obtained to review the data from electronic medical records. SPSS version 28 statistical software was used to analyze the data.

Summary of Results From all VLBW infants born during this period 152/642 (23.7%) had temperature below 36.5°C on admission to NICU while 104/652 (15.9%) were diagnosed to have hypoglycemia. Birth weight or gestational age had no impact on either hypothermia or hypoglycemia on NICU admission. Infants needing resuscitation including chest compressions and epinephrine administration were at highest risk for hypothermia. These infants were also noted to have metabolic acidosis and low 5 min apgar scores. Infants presented with hypoglycemia were small for gestational age (57%) vs 24% p=0.009. Maternal medical conditions including diabetes did not put these infants at additional risk for hypoglycemia. Hypoglycemia on admission was also associated with additional risk of requiring higher ventilation as well as oxygen requirement in the first 24 hours of NICU stay.

Conclusions VLBW Infants needing resuscitation in the delivery room are at risk for hypothermia. These infants may present with metabolic acidosis on admission to NICU. Hypoglycemia on admission may predispose VLBW infants for more respiratory support.
breastfeeding among newborns in the normal nursery from a single center in Los Angeles. These results prompted the creation of a specific task force to counter the detrimental effect of the pandemic on breastfeeding. prospective studies would be useful in assessing the long-term effects of the SARS-CoV-2 pandemic on breastfeeding rates and associated effects on infant immunity, maternal-child bonding, and long-term health.

Abstract #193
EXPECTED GROWTH TRENDS IN A LARGE COHORT OF ALMOST 7000 PRETERM INFANTS FROM BIRTH TO EIGHTEEN YEARS

1-2 J Barnard, 2 A Defante, 1-2 J Ryu. 1 University of California San Diego, La Jolla, CA; 2 Rady Children’s Hospital San Diego, San Diego, CA

Purpose of Study Despite numerous studies about the growth of preterm infants (PI) postnatally, there is still no consensus on expected growth rates for PI through childhood. The standard of care is to correct for gestational age (GA) until age two years during which time PI are expected to ‘catch up.’ In addition to correcting for GA, there may be a need to account for growth restriction. There are conflicting studies on whether infants born small for gestational age (SGA) ‘catch up’ by age two years. However, to the best of our knowledge, there are no studies with this cohort size following the growth of SGA PI over 18 years.

Methods Used This retrospective cohort study of 6916 followed infants born between 23-32 weeks over 18 years. Data was pulled from Rady Children’s Hospital electronic medical record system which includes specialists as well as over 30 general pediatrician offices. Being the main institution for follow-up of premature infants in a very large catchment area allowed for longitudinal follow-up of a large cohort. Infants were categorized as SGA if their birthweight was ≤ tenth percentile birthweight for their GA, AGA if tenth to ninetieth percentile and LGA if ≥ 90th percentile based on the WHO Fetal Growth Charts. Their weights and BMIs at ages two through 18 years were categorized as above the tenth percentile, between the tenth and ninetieth percentiles and above the ninetieth percentile by CDC standards.

Summary of Results Using a Chi Squared test, preliminary results show statistically significant differences (p-values all < 0.001) in the counts of PI who are below the tenth percentile, between the tenth and ninetieth percentiles and above the ninetieth percentile by CDC standards at ages two through 18 years based on whether they were born SGA, AGA, or LGA. Those born SGA had more infants remain ‘small’ than those born AGA or LGA. But for all groups, we observed significantly more PI stay ‘small,’ less than the tenth percentile for weight, at ages two through 18 years than expected. This held true for BMI as well for ages two through 18 years for PI born SGA, AGA, or LGA with p-values all <0.001.

Conclusions As more extremely premature and very low birth weight infants are surviving, there is a need for further assessment of this subpopulation’s expected postnatal growth. The observed distribution of PI over two through 18 years differed significantly by their size at birth. Infants born SGA may continue below the tenth percentile for weight for several years and this may not be ‘abnormal’ growth for them. These infants may be seen by various specialists for failure to thrive but might just need different standards. This study validates the need for different expectations of growth for infants born growth-restricted and very premature.

Abstract #194
GENERAL MOVEMENT ASSESSMENTS IN THE SURGICAL GASTROINTESTINAL NEONATAL POPULATION

S Bell, S Espinosa*, K Kesavan, KL Calkins. University of California Los Angeles, Los Angeles, CA

Purpose of Study Infants with gastrointestinal disorders (GD) have increased survivorship with advances in neonatal medicine and surgery. GD infants are at risk for neurodevelopmental impairment; they require surgery and are at high risk for sepsis, growth failure, and prolonged hospital stays. There is evidence that the General Movement Assessment (GMA) is an early biomarker of motor delays, including cerebral palsy. Most studies have focused on extremely low birth weight infants (ELBW) and have neglected GD infants.

Methods Used In this retrospective single-site study, GD infants (i.e., gastroschisis, omphalocele, atresias) who underwent surgery within the first 90 days of life were compared to ELBW (2/20/18 – 5/01/2021). The primary outcome was GMA results during the writhing stage at 36–49 weeks corrected gestational age (normal or abnormal (poor repertoire (PR), cramped synchronous (CS), or chaotic)) and fidgety stage at 3–4 months corrected gestational age (normal or abnormal). Abnormal fidgety was defined as fidgety movements that were not observed or movements that have exaggerated amplitude, speed, or jerkiness.

Summary of Results There were 55 GD infants; 31 (56%) had at least one GMA. There were 33 ELBW; 28 (85%) with one GMA. Gestational age and birth weight were significantly different when the cohorts were compared. However, the number of surgeries were similar (1.5 (0.8) vs. 1.7 (0.9), p=0.3) for the GD an ELBW cohort, respectively (table 1).

Of the GD infants, 60% had an abnormal writhing stage GMA (0% CS, 60% PR) and 20% had an abnormal fidgety
Abstracts

#194 Table 1 Characteristics of GD and ELBW infants

<table>
<thead>
<tr>
<th>GD (n=31)</th>
<th>ELBW (n=28)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maternal age (years)</td>
<td>Mean (SD),%</td>
</tr>
<tr>
<td>Maternal age (years)</td>
<td>30 (7)</td>
</tr>
<tr>
<td>Caealread%</td>
<td>48</td>
</tr>
<tr>
<td>Maternal gravida</td>
<td>2 (2)</td>
</tr>
<tr>
<td>Maternal parity</td>
<td>2 (1)</td>
</tr>
<tr>
<td>Chorionamniotitis%</td>
<td>10</td>
</tr>
<tr>
<td>Female%</td>
<td>39</td>
</tr>
<tr>
<td>Gestational age (weeks)</td>
<td>36 (4)</td>
</tr>
<tr>
<td>Birth weight (kg)</td>
<td>45 (7)</td>
</tr>
<tr>
<td>Birth head circumference (cm)</td>
<td>32 (4)</td>
</tr>
<tr>
<td>Small for gestational age%</td>
<td>29</td>
</tr>
<tr>
<td>Maternal tobacco use</td>
<td>7</td>
</tr>
<tr>
<td>Maternal illicit drug use</td>
<td>17</td>
</tr>
<tr>
<td>Length of stay (days)</td>
<td>44 (42)</td>
</tr>
<tr>
<td>Necrotizing enterocolitis</td>
<td>3</td>
</tr>
<tr>
<td>Late onset sepsis</td>
<td>3</td>
</tr>
<tr>
<td>Age of first feed (days)</td>
<td>11 (12)</td>
</tr>
<tr>
<td>Total number of surgeries</td>
<td>1.5 (0.8)</td>
</tr>
</tbody>
</table>

*P

Stage GMA. Of the ELBW infants, 35% had abnormal writhing stage GMA (15% CS, 20% PR) and 22% had an abnormal fidgety stage GMA. Rates for abnormal GMAs were similar when the groups were compared (p= 0.1, 0.9).

Conclusions In this study, GD and ELBW infants were at high risk for abnormal GMAs. GMAs maybe a helpful tool to ensure that GD infants receive long-term follow-up and resources required to reach their developmental potential. Further longitudinal research is required to determine the accuracy of GMAs in the GD population.

#195 ABNORMAL BLOOD GAS AND OXYGEN REQUIREMENTS IN FIRST 24 HOURS OF LIFE AS INDICATORS OF MORBIDITY IN VERY LOW BIRTH WEIGHT INFANTS

1,4,5 S Nanduri*, 1,2 A Hisey, 1 Y Shao, 1 K Ramm, 1 L Barton, 1 R Ramanathan, 1 M Brinwale, 1 Los Angeles County University of Southern California Medical Center, Los Angeles, CA; 2 Drexel University College of Medicine, Philadelphia, PA; 3 Loyola University Chicago Stritch School of Medicine, Maywood, IL

Purpose of Study Oxygen requirement as well as abnormal blood gas values have often been used as indicators of morbidity in the premature neonate, however, there is mixed evidence regarding its utility to predict short and long term morbidity. This study aims to assess the impact of abnormal pCO2, pH, and FiO2 values in the first 24 hours of life of very low birth weight (VLBW) infants on short and long term outcomes.

Methods Used Data on all VLBW infants was retrospectively gathered from the electronic medical record between the years of 2009 and 2021. IRB approval was obtained prior. For blood gas levels, significance was calculated based on pH of 7.15 and CO2 of 50 mm Hg. FiO2 was considered significant if the infant required 50% in the first 24 hours of life. Each category was statistically analyzed against common neonatal outcomes including intraventricular hemorrhage (IVH), bronchopulmonary dysplasia (BPD), and retinopathy of prematurity (ROP).

Summary of Results Of 564 VLBW infants studied 46.6% had an abnormal pCO2 and 53.4% had a normal pCO2. 78.7% of these infants had an abnormal pH and 19.5% had a normal pH. 87.2% had a normal FiO2 and 12.8% had an abnormal FiO2. Infants with elevated pCO2, decreased pH, and high FiO2 were significantly more likely to need intubation in the delivery room, emergently, remained intubated at 24 hours and had significantly higher days of invasive mechanical ventilation (table 1). These infants were more likely to need surfactant and had higher mortality (table 1). Lastly, these infants were at increased risk of IVH and ROP (table 1). BPD was correlated with higher FiO2 as well as low pH. Neonates with abnormal FiO2 also showed an increased risk for having abnormal MRI before discharge (51.6% vs 69.0% p = 0.034).

Conclusions Abnormal blood gas and higher FiO2 in the first 24 hours of life is associated with increased need for intubation as well as invasive mechanical ventilation in the delivery room and NICU. VLBW infants showed increased likelihood of developing long term complications including BPD, severe IVH, and severe ROP.

#196 INTUBATION IN THE FIRST 24 HOURS AS AN INDICATOR OF NEONATAL MORBIDITY

1 N Nanduri*, 1,2 A Hisey, 1 Y Shao, 1 K Ramm, 1 C Marquez, 1 L Barton, 1 R Ramanathan, 1 M Brinwale, 1 Los Angeles County University of Southern California Medical Center, Los Angeles, CA; 2 Loyola University Chicago Stritch School of Medicine, Maywood, IL

Abstract #195 Table 1

<table>
<thead>
<tr>
<th>pCO2</th>
<th>pH</th>
<th>FiO2</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 50</td>
<td>&gt; 50</td>
<td>p-value</td>
</tr>
<tr>
<td>&gt; 7.15</td>
<td>&lt; 7.15</td>
<td>p-value</td>
</tr>
<tr>
<td>&lt; 50</td>
<td>&gt; 50</td>
<td>p-value</td>
</tr>
</tbody>
</table>

- Death: 3.0% 13.7% < 0.001 3.1% 29.8% < 0.001 4.1% 29.0% < 0.001
- Duration of invasive ventilation: 8.8 26.4 < 0.001 13.9 32.8 0.003 27.0 29.3 0.010
- Intubation in delivery room: 37.0% 54.9% < 0.001 40.4% 65.9% < 0.001 35.6% 72.5% < 0.001
- Emergency intubation: 32.1% 55.0% < 0.001 37.5% 65.5% < 0.001 36.9% 59.0% 0.008
- Intubated at 24 hours: 26.6% 67.9% < 0.001 37.7% 82.8% < 0.001 36.9% 88.6% < 0.001
- Sulfactant: 49.8% 64.3% 0.002 53.4% 69.8% 0.006 48.4% 59.2% 0.089
- Severe IVH: 2.3% 9.8% 0.01 2.4% 20.4% < 0.001 3.6% 20.6% 0.001
- Severe ROP: 5.5% 21.9% < 0.001 8.5% 36.6% < 0.001 10.3% 30.4% 0.011
- BPD: 48.4% 59.4% 0.055 50.8% 65.5% 0.043 4.1% 29.0% < 0.001
Purpose of Study: Delivery room intubation in the premature neonate has been previously correlated with an increased risk for neonatal morbidities such as bronchopulmonary dysplasia (BPD). Invasive ventilation for longer duration also puts these infants at similar risk. This study further assessed relationships between infants needing invasive ventilation at 24 hours of life and short term neonatal outcomes.

Methods: Used Retrospective data of preterm VLBW infants born between 2009 and 2021 at LAC + USC Medical Center was evaluated for invasive ventilation at 24 hours of life and common neonatal morbidities. Institutional IRB approval was obtained. Maternal factors, delivery room interventions and standard neonatal outcomes were analyzed.

Summary of Results: Out of the 313 infants meeting inclusion criteria, 136 (43.5%) required invasive ventilation beyond 24 hours of life. Infants born to mothers with histologic chorioamnionitis were more likely to need invasive respiratory support at 24 hours. These infants had lower birth weight as well as lower gestational age. Resuscitation in the delivery room including intubation and chest compressions were also strongly correlated. These infants had a higher incidence of patent ductus arteriosus (PDA) as well as severe intraventricular hemorrhage (IVH). Bronchopulmonary dysplasia (BPD), severe retinopathy of prematurity (ROP) and MRI abnormalities by term gestation were also more often seen.

Conclusions: VLBW infants who had invasive ventilation in the first 24 hours of life were more likely to have BPD, PDA, abnormal brain MRIs, and severe ROP during their NICU stay.

Abstract #197 Table 1: Factors associated with intubation at 24 hours

<table>
<thead>
<tr>
<th></th>
<th>Invasive ventilation in the first 24 hours (%)</th>
<th>Non invasive ventilation in the first 24 hours (%)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth Weight</td>
<td>749.7 ± 279 g</td>
<td>1100 ± 253 g</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Gestational Age</td>
<td>25.9 ± 2.3 weeks</td>
<td>28.1 ± 2.3 weeks</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Histologic</td>
<td>18.7</td>
<td>6.8</td>
<td>0.002</td>
</tr>
<tr>
<td>Chorioamnionitis</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Intubation in the delivery room</td>
<td>76.5</td>
<td>24.3</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Chest compressions</td>
<td>33.8</td>
<td>4.5</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Severe IVH</td>
<td>10.2</td>
<td>1.1</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>PDA</td>
<td>75.2</td>
<td>31.4</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>PDA requiring surgery</td>
<td>26.5</td>
<td>2.9</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Abnormal brain</td>
<td>61.0</td>
<td>33.7</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>MRI</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BPD</td>
<td>63.3</td>
<td>25.1</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Severe ROP</td>
<td>82.9</td>
<td>17.1</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Purpose of Study: Hepatitis C virus (HCV) is the leading cause of blood-borne infection globally with an associated increase from the ongoing opioid epidemic. Current recommendations call for antibody screening of HCV-exposed infants after 18 months of age or RNA testing after 2 months of age, however, studies have shown low compliance. We hypothesize that many women with HCV do not receive appropriate screening during pregnancy resulting in gaps in infant care. We seek to identify factors associated with suboptimal pediatric HCV screening that could improve screening and subsequent treatment in perinatally exposed and chronically infected children.

Methods: Used A retrospective chart review was completed using data obtained by Tricore Laboratories. The data assessed the yearly proportion of HCV in pregnant women who were tested from 2014–2019 at our institution, characterize their demographic and health information, and identify their infants and HCV testing status. Demographics of mothers and infants with HCV testing were compared to those without testing to determine if certain demographics portend a greater probability of follow up care for infants with possible congenital HCV infection.

Summary of Results: From 2014–2019, a total of 14,709 women delivered at our institution with 63% (n=9,310) receiving HCV testing. Of these women, 139 (1.5%) were antibody positive; 107 mother-infant pairs were included in our analysis. Only 29 infants (27%) had antibody testing and 4 infants (3.7%) received viral load testing. One child was found to be antibody and viral load positive. The majority of these infants (n=81) were discharged to their birth parent from the nursery or neonatal ICU regardless of testing status. Mean infant gestational age, mothers’ gravidity/parity, liver enzyme levels, time between initial positivity of the mother and birth of infant, and maternal viral load at prenatal care onset did not differ significantly between the infant groups. However, urine positivity for opioid replacement therapy (ORT; methadone or buprenorphine) appeared to approach significance (p=0.08) for mothers whose infants were tested. Maternal and infant ALT levels showed a 0.34 correlation.

Conclusions: Mothers receiving ORT were more likely to have infant testing completed. This could be partly due to involvement of these mothers in programs which subsequently screen their infants. Additionally, a correlation between maternal and infant ALT levels seemed to exist however it is known that LFTs can fluctuate in those with HCV and in newborns. Thus, we are unable to conclude that this finding was clinically significant especially given the small number of infants who had LFT testing.

Abstract #198 Table 1: PAID FAMILY LEAVE AND VERY LOW BIRTHWEIGHT INFANT HEALTH OUTCOMES

<table>
<thead>
<tr>
<th></th>
<th>Paid Family Leave (n=345)</th>
<th>Low Birthweight (n=209)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth Weight</td>
<td>749.7 ± 279 g</td>
<td>1100 ± 253 g</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Chorioamnionitis</td>
<td>75.2 ± 31.4</td>
<td>61.0 ± 33.7</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Birth Weight</td>
<td>749.7 ± 279 g</td>
<td>1100 ± 253 g</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Chorioamnionitis</td>
<td>75.2 ± 31.4</td>
<td>61.0 ± 33.7</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Purpose of Study: Paid family leave (PFL) is associated with improved infant health, potentially through increased breastfeeding. Little is known regarding the effects of PFL on very low birthweight (VLBW) infants, a population in which human breast milk (HBM) is critical. California (CA) was the first state to implement PFL in 2004. The primary aim of this
study was to determine the impact of California’s PFL program on use of HBM at discharge, necrotizing enterocolitis (NEC), and in-hospital mortality in VLBW infants.

Methods Used We conducted a quasi-experimental study by employing a difference-in-differences design using data from Vermont Oxford Network. PFL was defined as the intervention with CA as the exposed group and the rest of the Western U.S. (WUS) as the unexposed group. Singleton infants with birthweight <1500g cared for at VON participant hospitals in CA & rest of the WUS from 2001–2010 were included. Infants with early mortality or congenital anomalies were excluded. Relative risk (RR) and adjusted RR (aRR) of each outcome for infants born pre 2004 and post 2004 (the year of PFL implementation) were calculated using multivariate regression models, controlling for maternal race/ethnicity, birthweight, mode of delivery, and antenatal steroids. Trends pre/post PFL in CA vs. the WUS were compared to identify the effect of PFL on the outcomes of interest.

Summary of Results Of 41,633 infants who met inclusion criteria, both CA and WUS infants were more likely to receive HBM at discharge post vs pre PFL enactment in CA (CA RR 1.19 [95% CI 1.15–1.23], aRR 1.17 [1.11–1.24]; WUS RR 1.03 [0.99–1.07], aRR 1.04 [0.98–1.10]). Both groups had higher incidence of NEC post vs pre PFL (CA RR 1.32 [1.18–1.47], aRR 1.29 [1.15–1.45]); WUS RR 1.43 [1.25–1.64], aRR 1.43 [1.24–1.64]). Incidence of NEC was lower in CA compared to WUS throughout the study period. There were no significant differences in mortality in either group pre vs post PFL. There was a trend of increasing use of HBM at discharge in CA but not WUS both pre and post PFL. Trends in NEC and mortality did not differ between CA and WUS. Overall, no statistically significant effect of PFL on the outcomes of interest was found when comparing difference in differences.

Conclusions Implementation of PFL legislation in CA did not have a clear, significant impact on use of HBM at discharge, NEC, and mortality in VLBW infants. Inadequate duration or utilization of PFL may account for the lack of observed impact. PFL may also not be sufficient for families who spend long periods for NICU hospitalizations. Further research investigating individual level effects of PFL on VLBW infants and patterns of PFL utilization by families with infants in the NICU is warranted.
Methods Used Following Institutional Review Board (IRB) approval (#5190190), this prospective study was performed at two large academic Neonatal Intensive Care Units (NICU) in southern California. Our recruitment targeted mothers of very low birth weight infants(<1500 grams). After informed consent, they completed a questionnaire including demographics, health, substance usage and socioeconomic status. The infant data was collected from birth until NICU discharge and monitored if the development of NEC.

Descriptive statistics and qualitative analysis were performed as appropriate. P values <0.05 were considered statistically significant.

Summary of Results Seventy infants were enrolled: 37(57.9%) male and 33(47.1%) female. Only 12(17.1%) developed NEC ≥ Bell stage 2. NEC infants had a lower gestational age than infants without NEC 25.9 vs. 29.0 weeks (*p<0.05), and lower birth weight at 752.8 vs. 1082.3 grams (*p<0.05). There were no significant differences in mechanical ventilation, vasopressors, intracranial hemorrhage, infection, or feed initiation.

Mothers of NEC infants reported, on average, more overall stressors 1.5 vs 0.5 and specifically more emotional stressors 10/12 (83.3%) vs. 20/58 (35.7%) during pregnancy (*p<0.05) (table 1). Other maternal factors were not significantly associated with NEC: age at conception, pregnancy complications, smoking history, alcohol and drug usage, household income, education level, first born child and family history of prematurity.

Conclusions Very low birth weight infants (<1500 grams) with necrotizing enterocolitis (NEC) were smaller and born earlier. Our findings suggest that specifically emotional stressors and overall number of maternal stressors during pregnancy may be risk factors for developing necrotizing enterocolitis.
Abstracts

VARIATIONS IN PARENT PARTICIPATION IN NURSING CARE SESSIONS IN THE NICU BY MEDICAID STATUS

S Takamatsu*, AM Cunningham, J Dempsey, J Kelleher, AG Dempsey. University of Colorado – Anschutz Medical Campus, Aurora, CO

Purpose of Study Parents are faced with the challenge of navigating other external responsibilities (e.g., parenting of other children, work) while their infant is hospitalized in the NICU. Families may have financial barriers impacting stability of housing, childcare, and transportation. As a result, increasing and stabilizing engagement is a common research interest, as it is influential in skill development and caregiver efficacy. Cares sessions with nursing occur in the NICU throughout the day, offering a structured time to observe and partake in the care of the infant. Participation in cares lends itself to demonstration of skills, explanation of care, and can be helpful for parents to feel competent interacting and caring for their baby. The present study collected data on the frequency of parents’ attendance of cares sessions in the NICU. To begin to understand differences in potential social barriers to engagement, differences were compared based on mother’s Medicaid status.

Methods Used Our sample included 122 premature infants in the NICU who were part of a larger quality improvement study to enhance family engagement. Attendance of four or more cares sessions with nursing per seven days was set as the target goal. Bedside nurses entered data into the infant’s medical record. A $X^2$ test was performed to detect a difference in goal attainment by Medicaid status.

PREVALENCE OF NEONATAL INTENSIVE CARE UNIT ADMISSION AMONG PATIENTS WITH GENETIC TESTING

SB Zoucha*, J Jensen, JL Bonkowsky. University of Utah Health, Salt Lake City, UT

Purpose of Study Genetic disease is estimated to affect many critically ill neonates, but an unbiased determination of genetic disease prevalence in Neonatal Intensive Care Units (NICU) has been unclear. Since rapid and extensive genetic testing is increasingly available and practical and can impact outcomes, there is a need for identifying best clinical practices for testing use. We hypothesized that a history of NICU admission is more common in patients with known or presumed genetic disease; and that neonatal NICU characteristics can guide best use of genetic testing.

Methods Used We performed a retrospective population-based cohort analysis of children on whom genetic testing was performed at a tertiary children’s hospital. The hospital and specialists are the only providers of pediatric sub-specialty care in a 500-mile radius. We analyzed the cohort for a history of NICU admission, and clinical characteristics of the admission. We identified 3894 patients with a history of genetic testing with birthdates between 1/1/09 and 6/1/21 to identify a final cohort of 1611 patients.
Summary of Results Of this cohort, 132 patients with a history of NICU admissions were identified (8.2% of the cohort). Of patients with a history of NICU admission; 36% had a positive (diagnostic) genetic test result, 30% had a negative test, and 34% had an uncertain (VUS) result. Compared to children without a history of NICU admission; 38% had a positive test result, 26% negative, and 36% uncertain; differences between these groups was not significant (p = 0.34). The age at testing was lower in those with a history of NICU admission (3.4 years vs 4.6 years; p < 0.001).

Conclusions Our study suggests that a genetic condition is present in a minority of children with a history of NICU admission. Further work into clinical characteristics of NICU children in whom genetic diagnoses are considered will help prioritize use of genetic testing.

Purpose of Study Contemporary research has increasingly explored the possibility that preterm births (PTBs), the live-birth of babies prior to 37 weeks of completed pregnancy, are associated with first trimester vaginal microbiota abnormalities; yet the association between the two remains unclear.

Methods Used A systematic literature review was conducted to assess current evidence for the role of first trimester vaginal microbiota abnormalities driving PTBs. Following the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) 2009 guidelines, scientific databases, including Medline, Embase, and the Maternity & Infant Care Database (MIDIRS), were searched from January 2009 to March 2019. The search terms used were (vagina* microb* OR vagina* bacteria OR vagina* flora OR vagina* microflora* OR vaginal dysbiosis OR bacterial vaginosis) AND (preterm OR preterm OR premature OR early term OR early birth). Details on population/sample, study design, method of microbiota determination, measures of microbiota associations with PTBs, and Lactobacillus prevalence were extracted.

Summary of Results Thirteen studies (nine cohort, four randomized control trial) were included in the review. The results provided strong evidence for an association between abnormal first trimester vaginal microbiota and PTBs. Specifically, low-Lactobacillus, high-diversity microbial eoniches were found to be associated with an increased likelihood of PTB outcomes. Notably, two studies reported contradictory findings showing a negative association between first trimester AVF and PTBs, and three studies reported no significant association. Possible explanations for the negative association reported in Farr et al., 2015 include regional bias and inclusion of women with chronic conditions. Selection bias was also of concern in Koumans et al., 2010, which also reported a negative association, as participant recruitment was not randomized, possibly shifting the baseline health demographic of recruited women. No definite taxa-specific trends associated with PTB were identified; however, different studies reported Mycoplasma and Ureaplasma parvum increase the odds of PTB manifestation.

Conclusions Associations were largely consistent and strong, suggesting vaginal flora measurements might hold the potential to enable early prediction of PTBs. It is important to note that causality and a biological mechanism for AVF-associated PTBs is not yet proven, with more research being recommended.
illness, and is associated with adverse neurodevelopmental outcomes. Although delirium is recognized in pediatric ICUs, it is not commonly diagnosed in the neonatal population. The NICU at Rady Children’s Hospital is comprised of medically complex patients that are often on multiple medications for pain and sedation. Early recognition and treatment of delirium in the NICU may be helpful in improving clinical outcomes. We conducted a QI project to implement screening for neonatal delirium in high-risk patients. Our objective is to increase delirium screening (RASS/CAPD scores) from 0% to 85% in eligible NICU patients by 10/2021. Inclusion criteria are defined as NICU patients > or = 38 weeks corrected gestational age who are mechanically ventilated > 7 days and who are receiving any sedatives or opiates.

Methods Used Multiple interdisciplinary meetings were initiated with key stakeholders to develop an algorithm for the evaluation of neonatal delirium. Completion of the RASS (Richmond Agitation and Sedation Scale) and age-adjusted CAPD (Cornell Assessment of Pediatric Delirium) scores were used as the objective tool for delirium screening. Weekly nursing compliance with RASS/CAPD score documentation (figure 1) is the primary process measure. Outcome measures include child psychiatry consultations and a diagnosis of delirium.

Summary of Results Implementation of screening and data collection began in October 2020. After implementation, data from 10/2020 through 2/2021 showed an average weekly screening compliance of 76%. Our data shows a sustained shift with an average compliance of 88%, placing us above our goal of 85% compliance. Targeted future interventions to sustain our goal include: creation of an order set in the medical record and required documentation.

Conclusions Through this QI project, we have increased awareness of neonatal delirium as a diagnosis in our NICU. Our expectation is that early recognition of delirium in our chronic patients will lead to more timely management of symptoms and decreased use of narcotic and sedative medications. This early recognition will be important to these patients’ overall recovery.

#206 IMPROVING ELECTROLYTE AND MINERAL HOMESTASIS IN EXTREMELY PREMATURE INFANTS
S Markee, J Fuller, A Yaroslavski*, E Shenk, J Maxwell. University of New Mexico Hospital, Albuquerque, NM

Purpose of Study Infants ≤ 28 weeks gestational age (GA) are at increased risk for developing electrolyte and mineral abnormalities due to reduced baseline bone mineral content. ~80% of fetal calcium stores are obtained in the third trimester and ~54% of extremely preterm infants have metabolic bone disease. These infants rely heavily on early parenteral nutrition (PN) to provide appropriate electrolytes, but there is limited evidence on how to optimize calcium and phosphorus. Our primary outcome is to improve calcium and phosphorus by day of life (DOL) 7 in infants ≤ 28 weeks GA.

Methods Used This Quality Improvement project is currently in the fourth Plan-Do-Study-Act (PDSA) cycle. We reviewed electrolyte and PN data in infants ≤ 28 weeks GA in 2019 (cohort 0; n=16). In March 2020, we obtained daily serum calcium and phosphorus levels in infants ≤ 28 weeks GA during the first postnatal week (n=13). Upon review, a custom PN form was designed for the next cohort (n=9), introducing calcium and phosphorus in PN earlier. In our third PDSA cycle we implemented new stock fluids, which included calcium gluconate 0.5 mEq/dL, dextrose, and amino acids for immediate use following birth for those born ≤ 28 weeks GA (n=10). A comparison was made between all three prospective cohorts. Additionally, serum creatinine and ionized calcium (ical) levels were compared. A Grubb’s analysis was used followed by a student’s t-test.

Summary of Results Mean serum calcium on DOL1 between cohort 1 and 2 were similar (6.57 ± 0.25 mg/dL and 6.74 ± 0.12 mg/dL, respectively (p=0.61)). In cohort 3, the calcium level was noted to be lower on DOL3 compared to cohort 1 (9.6 ± 0.24 mg/dL and 9.03 ± 0.09 mg/dL, respectively (p=0.08)). Interestingly, mean serum creatinine on DOL1 nearly significantly decreased in cohort 2 (0.76 mg/dL ± 0.05 mg/dL), compared to cohort 1 (0.90 mg/dL ± 0.05 mg/dL), p=0.07. Mean serum creatinine remained < 1 mg/dL throughout the first week of life for infants in cohorts 2 and 3, while it trended higher in cohorts 0 and 1 over the first week of life.

Conclusions The new PN allowed a more consistent and gradual increase in serum calcium levels, while remaining in normal limits, during the first postnatal week. Additionally, creatinine levels were lower and ical levels remained in goal range, requiring less therapeutic intervention. Addition of calcium gluconate to stock fluids for infants ≤ 28 weeks GA on DOL0, as seen in cohort 3, augmented the improvement and stabilization of serum calcium and creatinine levels. Surprisingly, we have found that although we have been unable to show an improvement in our serum phosphorus for this patient population, we have seen improvement in calcium and creatinine levels in the first postnatal week.

#207 IMPLEMENTATION OF A NEONATAL MASSIVE TRANSFUSION PROTOCOL
1H Ko*, 1R Griggs, 1J Raval, 2T Zamora. 1University of New Mexico Health Sciences Center, Albuquerque, NM; 2Regents of the University of Minnesota, Minneapolis, MN

Purpose of Study A Massive Transfusion Protocol (MTP) is an institutional plan created to facilitate communication, ensure timely lab monitoring and reduce delays/errors when ordering multiple blood products. Though adult and pediatric (MTPs) exist, they are ill-suited for use in the neonatal population. Given the unique nature of neonates, this population could benefit from a neonatal-specific MTP.

Methods Used A pre-implementation survey was sent out to physicians, nurse practitioners, physician assistants, and nurses in the NICU. A multi-disciplinary team was then put together, involving clinical staff from the NICU, transfusion medicine, and the transfusion committee in order to develop a neonatal-specific MTP.

Summary of Results Fifty responses were recorded to 5 questions. 64% were aware of MTPs in general. Almost all participants (96%) correctly identified that MTPs were generally used for any patients requiring large blood volume replacements and/or multiple blood products. Forty-six percent responded that they had previously experienced a clinical situation in which there was difficulty obtaining blood products. Of the 46% who experienced difficulty getting blood...
products, 67% identified that the delay was mostly attributed to waiting for blood products to arrive.

Conclusions Our multi-disciplinary group collectively developed a neonatal-specific MTP to allow for a safer, timelier, and standardized approach to administering multiple blood products. Massive transfusions are rare but high-risk events in the NICU setting thus this study will hopefully improve patient outcomes related to transfusions. Our pre-implementation survey shows educational and practical barriers exist when attempting to order multiple blood products and further work will need to be done to address these barriers.

#208 HOSPITAL VARIATION IN EXTREMELY PRETERM BIRTH

GP Goldstein*, P Kan, C Phibbs, E Main, GM Shaw, H Lee. Stanford University, Stanford, CA

10.1136/jim-2022-WRMC.205

Purpose of Study To assess between-hospital variation in extremely preterm birth (EPTB) frequency when stratifying by hospital level of care, and determine the proportion of variance explained by differences in maternal and hospital factors.

Methods Used We assessed 7,072,562 births in California from 1997 to 2011, using hospital discharge, birth and death certificate data. We estimated the association between maternal and hospital factors and EPTB using multivariable regression, calculated hospital-specific EPTB frequencies and estimated between-hospital variances, intra-class coefficients, and median odds ratios stratified by hospital level of care.

Summary of Results Hospital frequencies of EPTB ranged from 0.01% to 3.00%. Between-hospital EPTB frequencies varied substantially, despite stratifying by hospital level of care and accounting for confounding factors. This variation appeared to be related to differences in a collection of hospital, maternal sociodemographic and medical factors, and other factors not accounted for in our study, such as barriers to maternal transfer prior to delivery.

Conclusions Our results demonstrate differences in EPTB frequency among hospitals when stratifying by hospital level of care. Proportion of EPTBs at level 1 and 2 NICUs should be further investigated and considered as a hospital quality measure.

#209 PERCEIVED MOTIVATIONS AND BARRIERS OF NEONATAL NURSE PRACTITIONERS IN FELLOW EVALUATION: A PILOT CASE STUDY

N Dyess*, University of Colorado Denver School of Medicine, Aurora, CO

10.1136/jim-2022-WRMC.206

Purpose of Study Fellowship programs must provide objective performance evaluations of trainees that are formative and come from multiple evaluators including nonphysicians. The utility of multisource feedback has been well demonstrated in the literature; however, it is difficult to ensure nonphysician staff fill out evaluations. The purpose of this study is to elicit the neonatal nurse practitioner’s (NNP’s) perceived motivations and barriers to completing fellow evaluations. As a pilot study, an additional purpose was to assess feasibility and refine methods.

The research questions guiding this study are: how do NNP’s view their role in the trainee evaluation process, what motivates NNP’s to fill out evaluations, what are the self-perceived barriers to evaluations, and how can we increase NNP evaluation of trainees.

Methods Used I performed a pilot study of a phenomenological, qualitative case study of NNP’s at the University of Colorado. A convenience sample of 3 NNP’s, selected via purposive sampling, participated in semi-structured, one-on-one interviews to explore the perceived motivations and barriers to evaluating fellows. A constructivist epistemological framework guided the study. The data was viewed through a theoretical framework inspired by Maslow’s motivation theory. Interview transcripts were coded inductively via the constant comparative method and then clustered into emergent themes using phenomenological reduction, horizontalization, imaginative variation, and thematic analysis.

Summary of Results A conceptual framework emerged from the data, consisting of five themes of driving and restraining forces to completing evaluations which interact in a process akin to a neuronal cell’s action potential. The
framework describes the implicit weighing of these forces to determine if a threshold for activation is reached to complete an evaluation. Themes are supported by Maslow’s motivation theory, with each of Maslow’s levels of need equating to a NNP’s level of need for completion of fellow evaluations.

The Fellow Characteristics theme describes how extremes of behavior, repetitive behavior, and a fellow’s value of feedback affect evaluation completion. The NNP-Fellow Relationship theme describes how increased exposure and knowledge of fellow repercussion affect evaluation completion. The Evaluation Characteristics theme illustrates how anonymity, specificity, feasibility, and timeliness affect evaluation completion. The NNP-Evaluation Relationship theme describes how knowledge of evaluator role/value and the evaluation process is critical to evaluation completion. The NNP Characteristics theme describes how alignment with preferred feedback strategies and seniority affect evaluation completion.

Conclusions The conceptual framework provides insights into the motivations and barriers to completion of fellow evaluations by NNPs that can inform measures to increase completion rates of trainee evaluations by nonphysicians.

### Abstract #209 Table 1 Themes of driving and restraining forces and representative quotes

<table>
<thead>
<tr>
<th>Themes</th>
<th>Driving Forces</th>
<th>Restraining Forces</th>
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<tbody>
<tr>
<td>Fellow Characteristics</td>
<td>* extremes of behavior; pattern/repetitive behavior; values feedback</td>
<td>* don’t think I’ve known that people want feedback necessarily ‘most experiences are kind of like somewhere in the middle…doesn’t stick out to you in the same way’ what if they’re having a bad week, what if they’re having a bad service, what if they’re having a bad day’</td>
</tr>
<tr>
<td>NNP-Fellow Relationship</td>
<td>* continuity/exposure; perceived hierarchy</td>
<td>* feel like there’s still a sense of hierarchy…it can feel like intimidating or strange to be critical of someone that you maybe see as your superior in some way or leader ‘it’s very difficult to know where your boundaries are and where the line is in as far as giving negative feedback’</td>
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<tr>
<td>Evaluation Characteristics</td>
<td>* perceived feasibility; confidentiality/anonymity; type of feedback; degree of specificity; degree of timeliness</td>
<td>* questions where it’s like give me an example…then I have to rack through my memory of the past month that you’ve been on service ‘I would be hesitant to give that exact situation because then they would know who gave that feedback’ ‘you’re thinking of one specific scenario but I don’t know where that fits in these like 3 questions that they ask about…It’s like you’re trying to fit your feedback into a mold that didn’t really go together’ you are given a random evaluation and you haven’t worked with that person in like a month’</td>
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<tr>
<td>NNP-Evaluation Relationship</td>
<td>* knowledge of evaluator role and value (to trainee and to program); knowledge of evaluation process and outcomes; time constraints, interruptions, unprotected time</td>
<td>* ‘I just think that our feedback because we are in a different role is also invaluable because you will be working with other nurse practitioners in community hospitals, other nurses in community hospitals, rather than other attendings 24–7’ ‘It’s very important to have the bedside nurse and NNP…join in on that conversation because you’re not just communicating with consultants and you’re not just communicating with other attendings…you speak to an attending differently than how you speak to a NNP or bedside RN…so I feel like having a bedside RN and NNP weigh in on what you’re doing is important’</td>
</tr>
<tr>
<td>NNP Characteristics</td>
<td>* preferred evaluation and feedback strategies; new to role, knowledge of own role, knowledge of culture/system</td>
<td>* ‘If it were simplified and we can just…have a place where there is eval forms…like ok I really have something to say about this person, I don’t want to wait for an evaluation to come out, I can go online and I can click on this tab and I can fill [it] out ‘more years into the role, like I can see the differences in where a fellow stands and where they are in their career and how well they do their job’</td>
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**Purpose of Study** The incidence of neonatal opiate withdrawal syndrome (NOWS) in the US has grown dramatically over the past two decades. Many rural hospitals are not equipped with resources and materials to manage best practices of these patients resulting in transfers to hospitals in bigger cities. The purpose of this study is to evaluate a curriculum we created to support rural hospitals to keep healthy infants with NOWS for observation instead of transferring them.

**Methods Used** The curriculum was used for quality improvement at a rural hospital and shared with providers in another state that expressed interest. To evaluate the curriculum, we conducted pre- and post-surveys of NOWS knowledge, attitudes, and care practices, plus post-curriculum interviews and focus groups.
EFFECTS OF POSTNATAL GLUCOCORTICOIDS ON BRAIN STRUCTURE IN PRETERM INFANTS, A SCOPING REVIEW

I Robles*, MA Eidsness, HM Feldman, SE Dubner. Stanford University School of Medicine, Stanford, CA

Purpose of Study Postnatal GCs (GC) are given for many indications in infants, including for the reduction in incidence and severity of bronchopulmonary dysplasia, a major risk factor for morbidity, mortality, and neurodevelopmental disability in children born preterm. Variation exists in medication, dosing, timing, and reported outcomes. Clinical neurodevelopmental outcomes after GC administration may be due, in part, to GC induced alterations in neonatal brain development. The objective of this scoping review is to identify what is known about the effects of GC treatment on brain structural development in preterm human infants in order to identify potential mechanisms by which GCs may affect later clinical neurodevelopmental outcomes and to identify gaps in the literature.

Methods Used A search query was developed to search online databases for original research on human infants, GCs, and brain structure. Potential article titles and abstracts were screened by two reviewers to identify papers for full-text review.

Summary of Results 6565 titles were identified based on the search query for title and abstract review. Inclusion and Exclusion criteria are shown in the table. 70 were included for full-text review. Multiple imaging modalities and outcomes were reported.

Conclusions GC effects on brain are of interest to a wide audience of researchers across the lifespan and across many clinical conditions. Relatively few human studies have directly assessed the effect of this intervention on early brain structural development. This study highlights the need for additional research on neonatal GCs and their potential effects on brain development.

Abstract #211 Table 1 Inclusion and exclusion criteria for title and abstract review

<table>
<thead>
<tr>
<th>Inclusion Criteria</th>
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<tbody>
<tr>
<td>1. Papers from 1990 and later</td>
<td>1. Conference abstracts, case studies/ reports if less than 10 subjects,</td>
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<tr>
<td>2. English language</td>
<td>2. Endogenous GC exposure</td>
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<tr>
<td>3. Published and peer reviewed</td>
<td>3. GCs only administered antenatally</td>
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<tr>
<td>4. Contains empirical data</td>
<td>4. No structural brain outcomes reported</td>
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<tr>
<td>5. Human subjects study</td>
<td>5. Animal study with no human participants</td>
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<tr>
<td>6. Infants born before 37 weeks gestation</td>
<td>6. Topical GC application only</td>
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<tr>
<td>7. Postnatal GC exposure (exogenous)</td>
<td>7. GCs only after birth hospitalization</td>
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<tr>
<td>8. GCs administered systemically - enteral, intravenous, intramuscular, inhaled, nebulized, intranasally administered, sublingual, or subcutaneous</td>
<td>8. GCs only after birth hospitalization</td>
</tr>
<tr>
<td>9. GCs administered to infants during birth hospitalization</td>
<td>9. Reports on at least one structural brain outcome measured after postnatal GC exposure</td>
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DOES MATERNAL AGE IMPACT FEEDING OUTCOMES IN PRETERM INFANTS?

A Patel*, C Bradley. 1University of California Irvine School of Medicine, Irvine, CA; 2University of California Irvine Department of Pediatrics, Irvine, CA

Purpose of Study For preterm infants in the Neonatal Intensive Care Unit (NICU), successful and safe oral feeding is a requirement for discharge from the hospital. There are a variety of factors that contribute to feeding outcomes as neonates learn to coordinate breathing during oral feeding. While studies have detailed the effects of advanced maternal age on various neonatal morbidities, there is a dearth of literature exploring the association between maternal age and feeding outcomes in neonates. This analysis sought to examine whether a correlation exists between maternal age at delivery and feeding outcomes in preterm infants.

Methods Used A retrospective chart review was conducted for 12 healthy infants born at less than 37 weeks gestational age who were admitted to the NICU and were transitioning from gavage to bottle feeding. Data was collected on maternal age at delivery, rate of milk transfer (in milliliters per minute), oral feeding performance (in percentage of milliliters taken during the feeding per milliliters prescribed), and feeding proficiency (in percentage of milliliters taken during the first five minutes of the feeding per milliliters prescribed). A correlation matrix was then developed via a linear regression model using these, and other, maternal and infant health factors to assess for possible correlations.

Summary of Results Maternal age showed a moderately positive correlation with two of the three feeding outcomes studied. The Pearson correlation coefficient (R) for the association between maternal age and feeding performance was +0.66. The R for the association between maternal age and feeding proficiency was +0.48. Maternal age was not associated with rate of milk transfer, with an R of +0.04.

Conclusions This preliminary analysis shows that increased maternal age may serve as a protective factor for the complex oral feeding process that is challenging for preterm infants due to immature neurodevelopment. Maternal age at delivery may be valuable to consider in studying progression to full...
We will continue education, data collection and analysis with the hope to achieve our measures and aims.

Adolescent medicine and general pediatrics II
Concurrent session
8:00 AM
Friday, January 21, 2022
LIMPING THROUGH A DIFFERENTIAL: AN UNCOMMON PRESENTATION TO A COMMON PEDIATRIC DIAGNOSIS

J Smith*, J Gardner, K Dreher, S Sanders. University of Arkansas for Medical Sciences, Little Rock, AR

10.1136/jim-2022-WRMC.212

Case Report A 4-year-old previously healthy boy presented to the pediatric emergency department (ED) with a one-month history of a progressively worsening limp with concern for osteomyelitis. He was recently evaluated by an orthopedic surgeon with x-rays of the ankle showing medial metaphyseal lucency and joint effusion concerning for infection as well as distal fibular sclerosis concerning for a healing fracture. His limp progressed to the inability to bear weight due to worsening left lower extremity pain. There was no history of trauma to the area. He complained of no other bone pain.

Physical Exam Exam was notable for left ankle swelling with tenderness to palpation and pain with passive movement. The left lower extremity had normal sensation and perfusion. The liver edge was palpated several centimeters below the costal margin. No lymphadenopathy was appreciated. No rashes or bruising noted.

Diagnostic Evaluation Laboratory evaluation was significant for elevated inflammatory markers. A complete blood count revealed a borderline microcytic anemia, but was otherwise normal. Magnetic resonance image (MRI) of the ankle showed diffuse abnormal bone marrow enhancement consistent with an infiltrative process. Further questioning revealed a history of night sweats, increasing fatigue, and intermittent fevers over the past month. Lactate dehydrogenase and uric acid were elevated and a peripheral smear showed atypical lymphocytes. The patient underwent bone biopsy. No drainable fluid collection was found. Cultures did not grow bacteria. Bone marrow aspirate with flow cytometry showed 60% B-lymphoblasts consistent with B-cell acute lymphoblastic leukemia (ALL).

Discussion Acute onset limping and acute lymphoblastic leukemia are commonly encountered problems in the ED. Extremity pain with limping carries a wide differential diagnosis in a child. Studies have shown that the initial diagnosis in a patient with limp is only correct 42% of the time. ALL often presents with diffuse bone pain; however, presentation with limp as the primary problem is rare. This case highlights the importance of a thorough history and physical exam. It was not until they were asked specifically about night sweats, weight loss, and fevers that the patient’s guardians responded in the affirmative. In the workup of the limping child, it is imperative to have malignancy on the differential even for localized bone pain.

REFERENCES

PROTEIN CONCENTRATION OF HUMAN MILK VARIES WITHIN A FEED AND OVER A 24-HOUR PERIOD

KA Bull*, AC Gogel, IE Williams, Y Bonney, MA McGuire, MK McGuire. University of Washington School of Medicine, Moscow, ID; University of Idaho College of Agricultural and Life Sciences, Moscow, ID

Purpose of Study Fundamental to understanding human milk composition is determining what is a representative milk sample. Many factors can influence the concentration of some milk components, and controlling for these variables is essential to documenting ‘typical’ milk composition. While variation in lipid content has been widely studied, there is less research on macronutrients. The purpose of this project was to determine if total protein concentration varies with time of day and during a feed. We hypothesized that we would detect no differences in protein based on these factors.

Methods Used This was an epidemiologic, repeated-measures trial involving 17 healthy women from Moscow, ID and surrounding areas who were ≥18 y of age, not taking antibiotics, and nursing healthy infants. Participants were 31.4 ± 3.6 y old and 223 ± 175 d postpartum. Milk collection kits were delivered to each mother’s home with detailed sampling instructions, and study personnel were available (via phone) during most of the sample collection times. During d 1, participants collected milk representing the foremilk, mid-milk, and hindmilk of a feed during a single breast expression. On d 2, milk was collected 4 times (T1-T4), 6 ± 1.5 h apart, the first being between 0500–0900 hr. Protein concentrations in milk were measured using a colorimetric method (Pierce BCA Protein Assay Kit) with human serum albumin (Sigma-Aldrich) as the standard. Data were analyzed using linear mixed models with participant as a random variable while assuming an autoregressive correlation structure for the repeated measures within participant as implemented in SAS (v 9.4).

Summary of Results Contrary to our hypotheses, protein concentration varied within a feed (p = 0.0058) and over a 24-hr period (p = 0.0046). Protein concentrations of foremilk and mid-milk were lower than that of hindmilk (17.66 ± 0.46, 17.80 ± 0.46, and 18.56 ± 0.57 g/L, respectively; p = 0.0210 and p = 0.0015, respectively). Protein concentrations increased in a stepwise fashion over the 24-hr period (17.31 ± 0.51, 17.83 ± 0.44, 18.37 ± 0.45, and 19.15 ± 0.44 g/L at T1, T2, T3, and T4, respectively).

Conclusions Our results demonstrate that human milk protein concentration increases from the beginning to the end of a feed and over a 24-hr period. These findings contradict the majority of published studies and bring into question current published values of ‘typical’ human milk protein concentration. Our data suggest that time within a feed and time of day must be considered when collecting a representative milk sample for protein analysis.
Case Report Autoimmune (AI) diseases have been shown to be more common in patients with Klinefelter Syndrome (KS) than the general population. To our knowledge, there are no known records of autoimmune scleritis in Klinefelter Syndrome. We report a unique case of a 13-year old male with KS and AI scleritis.

Methods The medical records of a patient with KS and autoimmune scleritis were reviewed over a 3-year period.

Results A 13-year old male with KS presented 1–2 week of right eye edema, erythema, decrease in vision, and mild ophthalmalgia and ophthalmodynia, worsened with eye movement. Patient reported concurrent pain in knees and shoulders. Patient was noted to have ‘red eyes’ as a child and diagnosed with keratitis. Family history was positive for arthritis, unexplained deep vein thrombosis in his mother, and antiphospholipid syndrome in grandfather. Physical exam was notable for the right eye with red, injected sclera and conjunctiva.

Further studies included MR of the right orbit showing mild right proptosis, as well as right peribulbar, intrabulbar and extracanal intraorbital edema and enhancement. Right uveal thickening and enhancement was seen and the extracocular muscles were normal in caliber and symmetry. The globes were otherwise unremarkable and the optic nerve/sheath complexes were normal in configuration. Final MRI impression was significant for right orbital and peribulbar cellulitis with retinitis and mild proptosis. Patient had a mildly elevated white blood cell count at 12.7 K/ul and was started on IV Vancomycin and Unasyn. There was no clinical response after 2 days of treatment. Ophthalmology determined that presentation was most consistent with a likely autoimmune etiology, after which prednisone treatment was started at 80 mg daily. The patient showed immediate improvement in symptoms. Primary immune screen included: Antinuclear antibodies (ANA) was positive, but titers <1:40 (titers in the range of 1:40 to 1:60 are considered low), anti-neutrophil cytoplasmic autoantibody (ANCA), rheumatoid factor (RF), Immunoglobulin (Ig) G subclasses, IgM, and IgA were all normal, SSA/SSB and HLA-B27 were negative. Initial erythrocyte sedimentation rate (ESR) was 31 and normalized to 2 (normal ESR in males ≤15 mm/hr) after being on prednisone treatment. There are no specific tests available to confirm autoimmune scleritis; however, patient was unresponsive to appropriate antibiotic therapy and highly responsive to prednisone treatment, implying an autoimmune etiology.

At the age of 14, gonadal failure and delayed puberty were diagnosed.

Conclusions Appropriate autoimmune screening should be part of the medical management of patients with KS. As presented in this case report, it is also important to be aware of eye symptoms, particularly scleritis, as the first symptom of autoimmunity in KS patients.

Purpose of Study Social media sites, such as Twitter, have been used to sample informal attitudes and messages shared about birth control between users. TikTok is a newer platform that appeals to younger users, that has not been used to study this topic. On TikTok, content creators openly share their experiences with birth control through dialogue and humorous interpretations. TikTok’s largest age group is 18–24, while Twitter’s is 30–49. Identification of sentiment about hormonal birth control in the previously untapped TikTok population could provide insight into younger patients’ attitudes towards birth control.

Methods Used In this qualitative IRB-approved study, 100 videos per hormonal birth control method (oral contraceptive pills [OCP], injections, intrauterine devices [IUD] and implant) were identified through hashtags. Given interest in user experience, we excluded videos posted by self-identified medical providers or commercial agents. Videos were then analyzed for sentiment, factual accuracy, and theme saturation.

Summary of Results 60% of the content we surveyed on TikTok was negative, while 8% was positive. A previous study on Twitter had found a majority of birth control content was neutral or positive.

TikTok content most commonly highlighted side effects (59.5%), experiences with providers (19.5%), and humor (18.5%). OCP videos discussed side effects the least (37%) and were more likely to include humor (37%). Implant (70%), IUD (67%) and injection (64%) videos focused on side effects much more than OCP videos. IUD (18%) and OCP (13%) videos were most likely to contain misinformation, IUD contained the most videos mentioning complications (17%).

We found that videos discussing short-acting methods were twice as likely to be positive (12%) compared to long-acting methods (5%). A previous study found that Tweets mentioning long-acting methods were more likely to be positive than short-acting methods.

Humor was utilized most in OCP (37%) and IUD (16%).

Many of the humorous videos joked about the onset and severity of side effects, or used humor to make fun of children’s misbehavior as their motivation for using birth control. One content creator stated, ‘I take the pill so my boyfriend doesn’t have to be a teen dad’. Much of the humorous content on TikTok relied on trending audio tracks, self-deprecating jokes, and facial expressions.

Conclusions Attitudes and themes towards hormonal birth control options presented on TikTok differ from previous studies on public platforms, and introduce humor and use video formats that both resonate with the application’s younger users. Recognition of different perceptions of birth control by younger TikTok users in comparison to the older Twitter users can be used to strategically target misinformation and potentially identify generational differences in perception of birth control.
Purpose of Study: Inequalities in educational opportunities and lack of STEM identity have contributed to underrepresentation of various minority backgrounds in medical professions. We developed Kidz MedEd, an innovative pipeline program to increase STEM identity in impoverished minority youth interested in health careers through the creation of animated medical education videos and medical infographics.

Methods Used: High school students were recruited from the UCSF Fresno Doctors Academy Summer Program while premedical college students volunteered to provide mentorship to the high school students. Learners were divided into 3 groups containing medical, college, and high school student(s). Each group was headed by a pediatric subspecialist and met weekly via Zoom for mentorship and medical education. Animated medical education videos were created with Powtoon, with Canva used to create visually appealing medical infographics.

Summary of Results: Students learned digital literacy skills and improved their fund of medical knowledge while collaborating with both peers and near-peers to make informative infographics and videos on a variety of topics. An emphasis was placed on copyright literacy, gender and racial neutrality, and provision of medically accurate information to those with low literacy levels. Students were able to make informative infographics and videos on a variety of topics including asthma, constipation, diabetes, hepatic steatosis, hormones, obesity, and thyroid disease.

Conclusions: The Kidz MedEd project has provided underrepresented minority high school and college students from California’s impoverished Central Valley with an opportunity to gain insight into the medical field, learn valuable digital literacy skills, while simultaneously receiving mentorship from healthcare professionals. It demonstrates that educational technology can be used to create innovative medical learning experiences for high school and college students interested in health careers. Collaborations such as this can benefit both disadvantaged students who wish to gain greater exposure to the medical field and physicians who lack the time to create educational tools such as animated medical videos, infographics, and other social media content.

Cardiovascular III
Concurrent session
8:00 AM
Friday, January 21, 2022

#220 SMIDT HEART INSTITUTE TAKOTSUBO REGISTRY: ADJUDICATION METHODS

1 J Maughan, 1 O Obrutu, 1 B Tjoe, 1 R Herscovic, 1 P Moy, 1 N Rojas, 1 P Marano, 1 J Wei, 1 C Shufelt, 1 C Baiery Merz. Cedars-Sinai Medical Center, Los Angeles, CA; 2 Sheba Medical Center, Tel Hashomer, Israel
10.1136/jim-2022-WRMC.217

Purpose of Study: Takotsubo syndrome (TTS) is an acute transient left ventricular dysfunction often triggered by emotional or physical stressors and seen predominantly in post-menopausal women. The Smidt Heart Institute Takotsubo Registry at Cedars-Sinai Medical Center aims to understand the prevalence, recurrence and prospective status of TTS. Rigorous medical record review is needed to adjudicate the diagnosis of TTS according to the International Takotsubo (InterTAK) Diagnostic Criteria.

Methods Used: Once enrolled into the Takotsubo Registry, research staff obtain medical records of all prior TTS events from the enrollees directly (upload via HIPAA secure BOX), through medical record request using a signed medical release of Information Authorization form, or via the electronic health record EPIC and Care Everywhere system. Once medical records are obtained, study staff detail the completion of the records and note the TTS clinical picture, hospitalization labs, TTS triggers, electrocardiograms (EKG), echocardiograms (ECHO) during admission and recover, coronary angiography and ventriculogram findings, past medical history, and past and present medications. The minimum records needed for adjudication are troponin levels with ECGs during event, ECHOs, angiography and hospitalization summary. Once medical records are complete, they are sent to be adjudicated by two board-certified cardiologists in consensus using the following InterTAK Diagnostic Criteria: 1) transient left ventricular dysfunction with or without right ventricular dysfunction presenting as apical ballooning or midventricular, basal, or focal wall motion abnormalities, typically extending beyond a single epicardial vascular distribution, 2) absence of culprit atherosclerotic coronary artery disease 3) new and reversible ECG abnormalities, 4) positive elevation in cardiac troponin, 5) absence of infectious myocarditis. Events are adjudicated on REDCap as definite TTS, probable TTS, probable not TTS and definite not TTS based on the medical record review. Enrollees receive written adjudication results.

Summary of Results: From January 2019 to July 2021, 104 of the 131 participants enrolled in the registry have been adjudicated. Of the baseline events, 101 events were adjudicated as definite TTS, 2 probable TTS, 10 probable not TTS and 8 definite not TTS. Furthermore, 32 enrollees reported a total of 53 recurrent events (between 2–5 events per enrollee) which include 14 definite TTS, 11 probable TTS, 7 probable not TTS and 23 definite not TTS events.

Conclusions: The Smidt Heart Institute Takotsubo Registry aims to investigate TTS pathophysiology and gain an accurate estimation of TTS reoccurrence. Limitations of the adjudication process include time intensive collection of medical records from institutions which is a focus for improvement.

#221 RISK FACTORS FOR SEIZURES FOLLOWING HEART TRANSPLANTATION

1 L Mishalani, 1 N Patel, 1 T Singer-Englar, 1 J Kim, 1 M Hamilton, 1 J Kobashigawa. Columbia University, New York, NY; 2 Cedars-Sinai Smidt Heart Institute, Los Angeles, CA
10.1136/jim-2022-WRMC.218

Purpose of Study: Seizures following heart transplantation is not uncommon. Patients who have had previous strokes have a higher propensity to develop seizures after open heart surgery. Patients who undergo heart transplant may also have underlying atherosclerotic vascular disease and have a history of stroke or cerebral microvascular disease. Both of these entities may be a risk for developing seizures after heart transplant. It has not been well established what the frequency of seizures is and whether or not we can identify risk factors as to who will develop these seizures.
Abstracts

Abstract #221 Table 1  Univariate analysis

<table>
<thead>
<tr>
<th></th>
<th>Seizures Post-HTx</th>
<th>No Seizures Post-HTx</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male Gender</td>
<td>63.5%</td>
<td>67.8%</td>
<td>0.271</td>
</tr>
<tr>
<td>History of Diabetes</td>
<td>37.5%</td>
<td>43.0%</td>
<td>0.750</td>
</tr>
<tr>
<td>History of Hypertension</td>
<td>55.0%</td>
<td>63.2%</td>
<td>0.012</td>
</tr>
<tr>
<td>History of Stroke</td>
<td>22.5%</td>
<td>21.7%</td>
<td>0.388</td>
</tr>
<tr>
<td>History of Atherosclerosis</td>
<td>67.5%</td>
<td>50.2%</td>
<td>0.035</td>
</tr>
<tr>
<td>History of Smoking</td>
<td>40.0%</td>
<td>39.6%</td>
<td>0.962</td>
</tr>
<tr>
<td>History of Previous Seizures</td>
<td>10.0%</td>
<td>2.9%</td>
<td>0.017</td>
</tr>
</tbody>
</table>

Abstract #221 Table 2  Multivariate Analysis

<table>
<thead>
<tr>
<th></th>
<th>Odds Ratio (95% CI)</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>History of Hypertension</td>
<td>2.193 (1.130-4.259)</td>
<td>0.020</td>
</tr>
<tr>
<td>History of Atherosclerosis</td>
<td>1.786 (0.891-3.581)</td>
<td>0.102</td>
</tr>
<tr>
<td>History of Previous Seizures</td>
<td>3.924 (1.204-12.791)</td>
<td>0.023</td>
</tr>
</tbody>
</table>

Methods  Used Between 2015 and 2020, we assessed 560 patients undergoing heart transplantation and found that 40 patients developed seizures within the first month postoperatively. We identified risk factors for these patients such as previous stroke, atherosclerotic vascular disease, previous history of seizures, history of diabetes, history of smoking, and male gender. The main endpoint was to establish the frequency of seizures following heart transplant surgery and to assess risk factors to develop seizures. These patients with seizures were compared to patients without seizures.

Summary of Results The incidence of seizure was 7.1% following heart transplant surgery. Univariate analysis found that hypertension, history of atherosclerosis, and history of seizures were significant risk factors to developing seizures immediately postoperatively. Multivariate analysis found that only hypertension and history of previous seizures were significant for the development of seizures post-heart transplant. Patients who were on tacrolimus appear to have a lower threshold to experience seizures and these patients were subsequently switched to cyclosporine.

Conclusions Patients undergoing heart transplant with hypertension and history of seizures are at risk to develop seizures postoperatively. These patients may be considered for prophylactic anti-seizure therapy for the first 30 days following heart transplant surgery.

Case Report Systemic lupus erythematosus (SLE) is an autoimmune disease where the immune system can attack its own tissue and causes damage to many organs, even the heart. This case report shows a rare presentation of SLE myocarditis in an 18 year old female.

Methods Approval was obtained from IRB. A single patient case report was conducted.

Case Presentation A 18-year-old Filipino female with no past medical history presented to the emergency department with bilateral extremities and periorbital swelling for 10 days with recent development of oral mucosal ulcers. Patient was found to have hyponatremia and nephrotic range proteinuria; in addition to a positive family history of autoimmune disorder, an autoimmune workup was ordered. The work up showed high antinuclear antibodies titer of 1:1280 and high double-stranded DNA antibody of 6 IU/mL which was highly suspicious of SLE related cause of patient’s symptoms. Further workup with ultrasound-guided renal biopsy resulted in lupus podocytopathy. Chest x-ray showed cardiomegaly so transthoracic echocardiogram (TTE) was ordered to rule out pericarditis and pericardial effusion. The TTE showed left ventricular ejection fraction (LVEF) 60%. Eight days later, the patient was found to be tachycardic to 130–140s with oxygen saturation of 88% so computerized tomography angiogram of the chest was obtained. It showed no evidence of pulmonary emboli but there was right lower lobe pneumonia, right sided small pleural effusion, and borderline cardiomegaly. Electrocardiogram showed no abnormalities but BNP was elevated to 3608 pg/mL and repeat TTE showed concentric left ventricular hypertrophy with LVEF 20% and small circumferential pericardial effusion. Guideline-directed medical therapy (GDMT) with lisinopril and carvedilol initiated. The patient was discharged with outpatient follow up.

Conclusion SLE can affect virtually all organs by the immune system attacking its own tissue and causing widespread inflammation and tissue damage. The heart is a common affected organ usually presenting with pericarditis or premature coronary artery disease in patients with long standing disease while myocarditis is less common.

This case shows a rare presentation of SLE myocarditis in an acute setting seen with a huge decline in cardiac function over 8 days. One can argue that this could be a case of stress myocarditis due to the acute presentation. A better understanding of patients with SLE and the risk for specific cardiac manifestations of SLE should be researched for improvement of clinical outcome for future patients.

SMIDT HEART INSTITUTE TAKOTSUBO REGISTRY – STUDY DESIGN

1 0 Brutu*, 1 J Maughan, 1 B Tjo, 1 R Herscovici, 1 P Hoy, 1 N Rhojas, 1 P Marano, 1 J Wei, 1 C Shufelt, 1,2 H Rutledge, 1 C Bainey Merz. 1 Cedars-Sinai Smidt Heart Institute, Los Angeles, CA; 2 Chaim Sheba Medical Center, Tel-Hashomer, Israel; 2 VA San Diego Healthcare System, San Diego, CA; 3 University of California, San Diego, CA

10.1136/jim-2022-WRMC.220

Purpose of Study Takotsubo syndrome (TTS) is an acute form of transient systolic heart failure that occurs predominantly in women, often triggered by emotional or physical stressors. The Smidt Heart Institute Takotsubo Registry at Cedars-Sinai Medical Center aims to establish a database for deep phenotyping of this syndrome.

MYOCARDITIS AS AN EARLY MANIFESTATION OF SYSTEMIC LUPUS ERYTHEMATOSUS IN A YOUNG FEMALE

1P Chan*, 1VK Narang, 1T Joolhar, 1T Win. 1Ross University School of Medicine, Miramar, RI; 2UCLA-Kem Medical, Bakersfield, CA

10.1136/jim-2022-WRMC.219
Abstract #223 Figure 1 Recruitment and enrollment flowchart

Methods Used Registry participants are recruited from multiple sources: Deep 6-based medical records review, physician referrals, peer and self referrals (Abstract #223 figure 1). Deep 6 is an artificial intelligence software that analyzes clinical data to identify patients that match complex clinical trial criteria. Peer and self referrals are sourced through social media advertisements on a Facebook support group for TTS survivors. Recruited participants remotely sign consents and provide detailed information via questionnaires on REDCap. Medical records of TTS events are adjudicated by cardiologists using the International Takotsubo Diagnostic Criteria. MITRA kits are used to remotely collect blood samples for further analysis.

Summary of Results From January 2019 to July 2021, 131 participants (99% female, mean age: 61.2±9.8 years) enrolled in the registry across 25 US states and 3 other countries. About 41% of enrollees are self or peer referrals. Overall, 89% completed all baseline questionnaires and 78% returned filled MITRA kits. Completed questionnaires provide information about participants’ general health status, physical function, psychosocial history, and TTS events.

Conclusions The Takotsubo Registry will reach a large and diverse participant base by using a multifaceted approach to recruitment as well as tools to facilitate remote enrollment and study participation. Study questionnaires, blood samples and medical record review will allow for deep phenotyping of participants in order to deepen our understanding of TTS pathophysiology.

Purpose of Study Electronic cigarettes (e-cigarettes) have increased in popularity and pose a public health crisis, demanding further research on their cardiovascular health effects to better inform future interventions. We have shown that mice treated with e-cigarettes containing nicotine develop increased serum free fatty acid levels (FFAs) and systolic dysfunction associated with inflammation. As such, we aim to study the differentially expressed genes between the hearts of e-cigarette and saline-treated mice and to define the genes controlling the physiological normalization produced by acipimox (a lipolysis inhibitor). We hypothesize that acipimox will reverse the transcriptomic changes associated with e-cigarette-induced cardiomyopathy.

Methods Used C57BL/6J wild type mice were exposed to saline, e-cigarettes, and e-cigarettes plus acipimox for 12 weeks. Left ventricular RNA was sequenced for differential gene expression and analyzed using Ingenuity Pathway Analysis (IPA) and Gene Set Enrichment Analysis (GSEA).

Summary of Results Our preliminary echocardiographic data showed that acipimox abrogates the e-cigarette-induced increases in serum FFAs and cardiomyopathy. RNA-sequencing analysis showed that 79 genes were dysregulated by e-cigarette treatment, and IPA showed that these changes are associated with inflammation. 30 of these genes were normalized by acipimox treatment and are implicated in inflammation, atherogenesis, and cardiac function. GSEA revealed that acute myocardial infarction, circadian rhythm, and G2/M DNA damage checkpoint genes are enriched in e-cigarette-treated mice compared to e-cigarette plus acipimox and saline-treated mice.

Conclusions Our RNA-sequencing analysis offers a mechanistic insight into how acipimox prevents cardiomyopathy in e-cigarette-treated mice. These findings provide gene expression evidence suggesting that lipolysis is necessary for e-cigarette-induced cardiac dysfunction. This study and future studies will not only illuminate the harms involved with e-cigarettes, but will also aid in the identification of targets for intervention to address complications associated with e-cigarette usage.

Abstract #224 Cardiac transcriptome analysis of the electronic cigarette-induced cardiac dysfunction mouse model treated with acipimox

Purpose of Study Due to the novelty and severity of the COVID-19 pandemic, the long-term cardiac outcomes of children diagnosed with multi-system inflammatory syndrome (MIS-C) have not been extensively studied. The purpose of this study was to evaluate cardiac outcomes, comparing echocardiogram and lab results at admission and follow-up periods.

Methods Used A literature review using Google Scholar and Pubmed was conducted, utilizing keywords such as ‘long-term cardiac outcomes’, ‘multi-system inflammatory syndrome in children’, and ‘Coronavirus Disease 2019’. We included studies of MIS-C patients that reported echocardiographic data and outcomes, troponin levels, and a minimum follow-up period of two weeks. Studies only evaluating electrocardiogram results were excluded.

Summary of Results Left ventricular (LV) dysfunction or decreased LV ejection fraction (LVEF) were the most common

#225 Long-term cardiac outcomes of multi-system inflammatory syndrome in children (MIS-C)


1University of California Irvine School of Medicine, Irvine, CA; 2Children’s Hospital of Orange County, Orange, CA

10.1136/jim-2022-WRMIC.222

Purpose of Study Due to the novelty and severity of the COVID-19 pandemic, the long-term cardiac outcomes of children diagnosed with multi-system inflammatory syndrome (MIS-C) have not been extensively studied. The purpose of this study was to evaluate cardiac outcomes, comparing echocardiogram and lab results at admission and follow-up periods.

Methods Used A literature review using Google Scholar and Pubmed was conducted, utilizing keywords such as ‘long-term cardiac outcomes’, ‘multi-system inflammatory syndrome in children’, and ‘Coronavirus Disease 2019’. We included studies of MIS-C patients that reported echocardiographic data and outcomes, troponin levels, and a minimum follow-up period of two weeks. Studies only evaluating electrocardiogram results were excluded.

Summary of Results Left ventricular (LV) dysfunction or decreased LV ejection fraction (LVEF) were the most common...
initial finding and seen in up to 60% of patients with MIS-C. Upon follow-up, these abnormalities had resolved in the vast majority of patients, with residual dysfunction seen in a few patients. Coronary artery dilations and aneurysms were seen in up to 33% of patients initially, and resolved in almost all patients, especially when follow-up period was longer. Elevated troponin levels had normalized at follow-up. Other common MIS-C-associated cardiac abnormalities included myocarditis, mitral valve regurgitation, and pericardial effusion, which were often resolved by the follow-up period.

Conclusions Our review suggests that patients with MIS-C have favorable long-term cardiac outcomes. In general, the initial cardiac complications and findings substantially improve or resolve by follow-up. Persistent cardiac abnormalities are rare, and often mild in severity. Further research with larger sample sizes and longer follow-up periods must be evaluated to better understand the long-term cardiac outcomes of MIS-C.

#226 WET BERIBERI IN A PATIENT WITH ISCHEMIC CARDIOMYOPATHY AND GASTRIC BYPASS: A CASE REPORT

T Azenkot*, OM Campa. University of California Davis Health System, Sacramento, CA

Case Report A 49-year-old woman with hypertension and Roux-en-Y gastric bypass 9 years prior presented for subacute exertional dyspnea and chest pain. Her home medications were multivitamins. On admission, blood pressure was 163/123 mmHg and heart rate 96 bpm. Hemoglobin was 12.9 g/dL. Troponin was 22 and unchanged on repeat. B-type natriuretic protein was 276 pg/mL. An electrocardiogram was without changes concerning for acute coronary syndrome. An echocardiogram showed global hypokinesis with an estimated left ventricular ejection fraction (LVEF) of 25%. Coronary angiography demonstrated complete total occlusion of the right coronary artery, though her global akinesis was out of proportion to this finding. She was discharged on guideline directed medical therapy for heart failure (HF).

Four months later, the patient returned to the emergency department for dizziness. Her blood pressure was 53/43, with improvement to 91/53 after fluids. Creatinine was 2.29 and granular casts seen on urinalysis. Her alkaline phosphatase and aspartate and alanine transaminases were 1,213 U/L, 615 U/L, and 338 U/L, respectively, a pattern consistent ischemic hepatopathy. An echocardiogram showed improved LVEF to 32% with no focal wall motion abnormalities. Thiamine level was 51 (normal 70–180) nmol/L, concerning for component of thiamine deficiency-induced cardiomyopathy, or wet beriberi. Supplementation was initiated. HF medications were held. Her dizziness and lab abnormalities resolved with fluids.

The prevalence of thiamine deficiency in patients admitted for HF is estimated to be 30% (Hanninen et al., 2006). This may be in part due to the association of diuretics, including furosemide, with increased renal excretion and decreased intestinal absorption of thiamine. Post gastric bypass patients also have a prevalence of thiamine deficiency up to 30% due to disruption of thiamine absorption in the duodenum and jejunum (Wilson, 2020). This renders patients with HF and gastric bypass, such as the present case, dually vulnerable to thiamine deficiency. Several small studies have demonstrated improved HF outcomes (i.e., improved LVEF and decreased hospital admission rates) with thiamine supplementation, though further studies are needed (Jain et al., 2015).

Abstract #225 Table 1 Cardiac characteristics and long-term outcomes in patients with multisystem inflammatory syndrome in children (MIS-C)

<table>
<thead>
<tr>
<th>Author, Location, Year</th>
<th>Mean/Median Age (years)</th>
<th>Initial LV Dysfunction/Decreased LVEF (%)</th>
<th>Initial Dilations (%)</th>
<th>Initial Aneurysms (%)</th>
<th>Elevated Troponin (%)</th>
<th>Myocarditis, MR, PCE (%)</th>
<th>Follow-up LV Dysfunction/Decreased LVEF (%)</th>
<th>Follow-up Dilations (%)</th>
<th>Follow-up Aneurysms (%)</th>
<th>Follow-up Period</th>
</tr>
</thead>
<tbody>
<tr>
<td>Minocha et al., NYC, 2021</td>
<td>2.8</td>
<td>4/33 (13%)</td>
<td>2/33 (7%)</td>
<td>0/33 (0%)</td>
<td>7/32 (22%)</td>
<td>1/33 (5%), 5/33 (15%), 1/33 (3%)</td>
<td>0/33 (0%)</td>
<td>0/33 (0%)</td>
<td>0/33 (0%)</td>
<td>14 days</td>
</tr>
<tr>
<td>Galtonde et al., Georgia, 2020</td>
<td>8</td>
<td>8/12 (67%)</td>
<td>1/12 (8%)</td>
<td>0/12 (0%)</td>
<td>-</td>
<td>MR: 12/12 (100%), PCE: 5/12 (42%)</td>
<td>1/12 (8%)</td>
<td>2/12 (17%)</td>
<td>0/12 (0%)</td>
<td>Median = 45 days</td>
</tr>
<tr>
<td>Closer et al., USA, 2021</td>
<td>7.3</td>
<td>7/18 (39%)</td>
<td>0/18 (0%)</td>
<td>0/18 (0%)</td>
<td>-</td>
<td>PCE: 2/18 (11%)</td>
<td>1/11 (9%)</td>
<td>1/11 (9%)</td>
<td>0/11 (0%)</td>
<td>Median = 29 days</td>
</tr>
<tr>
<td>Dionne et al., USA, 2020</td>
<td>9.7</td>
<td>15/25 (60%)</td>
<td>3/25 (12%)</td>
<td>2/25 (8%)</td>
<td>2/25 (8%)</td>
<td>-</td>
<td>2/15 (13%)</td>
<td>-</td>
<td>-</td>
<td>Median = 51 days</td>
</tr>
<tr>
<td>Penner et al., UK, 2021</td>
<td>10.2</td>
<td>15/46 (33%)</td>
<td>3/45/46 (84%)</td>
<td>0/46 (0%)</td>
<td>1/46 (2%)</td>
<td>0/46 (0%)</td>
<td>1/46 (2%)</td>
<td>1/46 (2%)</td>
<td>6 months</td>
<td></td>
</tr>
<tr>
<td>Klosy et al., Boston, 2020</td>
<td>3.5</td>
<td>7/12 (58%)</td>
<td>1/12 (8%)</td>
<td>1/12 (8%)</td>
<td>9/12 (75%)</td>
<td>MR: 3/12 (25%), PCE: 4/12 (33%)</td>
<td>1/8 (13%)</td>
<td>2/8 (25%)</td>
<td>1/8 (13%)</td>
<td>Median</td>
</tr>
<tr>
<td>Jhaveri et al., NYC, 2021</td>
<td>11.5</td>
<td>8/15 (53%)</td>
<td>-</td>
<td>4/15 (33%)</td>
<td>-</td>
<td>MR: 8/15 (53%), PCE: 2/15 (13%)</td>
<td>3/13 (23%)</td>
<td>-</td>
<td>1/13 (8%)</td>
<td>Median</td>
</tr>
<tr>
<td>Feldstein et al., US, 2021</td>
<td>9.7</td>
<td>172/393 (34%)</td>
<td>-</td>
<td>57/424 (13%)</td>
<td>-</td>
<td>PCE: 125/393 (25%)</td>
<td>1/172 (0.58%)</td>
<td>-</td>
<td>12/172 (7%)</td>
<td>2–20 weeks</td>
</tr>
<tr>
<td>Kobayashi et al., Canada, 2021</td>
<td>11.4</td>
<td>9/25 (36%)</td>
<td>2/25 (8%)</td>
<td>4/26 (16%)</td>
<td>9/23 (39%)</td>
<td>MR: 1/25 (4%), PCE: 0/25 (0%)</td>
<td>2/9 (22%)</td>
<td>-</td>
<td>-</td>
<td>1–2 months</td>
</tr>
<tr>
<td>Totals</td>
<td>2.8–11.5</td>
<td>13–60%</td>
<td>0–12%</td>
<td>0–33%</td>
<td>8–84%</td>
<td>-</td>
<td>0–23%</td>
<td>0–25%</td>
<td>0–13%</td>
<td>2 weeks–6 months</td>
</tr>
</tbody>
</table>

Abstracts

Abstracts

Abstract #224 cardiomyopathy and gastric bypass: a case report

WET BERIBERI IN A PATIENT WITH ISCHEMIC CASE REPORT

T Azenkot*, OM Campa. University of California Davis Health System, Sacramento, CA

10.1136/jim-2022-WRMC.223

Case Report A 49-year-old woman with hypertension and Roux-en-Y gastric bypass 9 years prior presented for subacute exertional dyspnea and chest pain. Her home medications were multivitamins. On admission, blood pressure was 163/123 mmHg and heart rate 96 bpm. Hemoglobin was 12.9 g/dL. Troponin was 22 and unchanged on repeat. B-type natriuretic protein was 276 pg/mL. An electrocardiogram was without changes concerning for acute coronary syndrome. An echocardiogram showed global hypokinesis with an estimated left ventricular ejection fraction (LVEF) of 25%. Coronary angiography demonstrated complete total occlusion of the right coronary artery, though her global akinesis was out of proportion to this finding. She was discharged on guideline directed medical therapy for heart failure (HF).

Four months later, the patient returned to the emergency department for dizziness. Her blood pressure was 53/43, with improvement to 91/53 after fluids. Creatinine was 2.29 and granular casts seen on urinalysis. Her alkaline phosphatase and aspartate and alanine transaminases were 1,213 U/L, 615 U/L, and 338 U/L, respectively, a pattern consistent ischemic hepatopathy. An echocardiogram showed improved LVEF to 32% with no focal wall motion abnormalities. Thiamine level was 51 (normal 70–180) nmol/L, concerning for component of thiamine deficiency-induced cardiomyopathy, or wet beriberi. Supplementation was initiated. HF medications were held. Her dizziness and lab abnormalities resolved with fluids.

The prevalence of thiamine deficiency in patients admitted for HF is estimated to be 30% (Hanninen et al., 2006). This may be in part due to the association of diuretics, including furosemide, with increased renal excretion and decreased intestinal absorption of thiamine. Post gastric bypass patients also have a prevalence of thiamine deficiency up to 30% due to disruption of thiamine absorption in the duodenum and jejunum (Wilson, 2020). This renders patients with HF and gastric bypass, such as the present case, dually vulnerable to thiamine deficiency. Several small studies have demonstrated improved HF outcomes (i.e., improved LVEF and decreased hospital admission rates) with thiamine supplementation, though further studies are needed (Jain et al., 2015).
Case Report Identify clinical manifestations of regional cardiac tamponade

Case Presentation The patient is a 31-year-old female with stage IV ER+, PR+, Her2- breast cancer complicated by recurrent malignant pleural effusions who presented with worsening dyspnea. Vital signs on admission included BP 121/91, HR 124, RR 32 and oxygen saturation of 99%. Shortly after admission, blood pressure dropped to 92/68 and oxygen saturation was 97% on 2L nasal cannula. On exam, patient had increased work of breathing with elevated neck veins. EKG showed low voltage QRS complexes with electrical alternans. CTA chest revealed a large pericardial effusion with no evidence of pulmonary embolus (PE). Given the pericardial effusion, a manual pulsus was checked and was negative. TTE showed a large pericardial effusion with moderate right ventricular (RV) collapse, dilated inferior vena cava and normal left ventricular systolic function. Based on the echocardiographic evidence of pericardial effusion with RV collapse in the context of worsening hypotension and tachycardia, the diagnosis of regional cardiac tamponade was made. Pericardiocentesis was performed with an opening pericardial pressure of 23mmHg and 680mL of fluid was drained with cytology confirming malignant pericardial effusion. Following the pericardiocentesis, the patient’s dyspnea and tachycardia resolved. Significant post-procedure pericardial drainage prompted placement of a pericardial window and the patient was discharged to hospice.

Discussion This patient’s presentation of dyspnea, tachycardia, progressive hypotension and distended neck veins fits multiple illness scripts including PE and cardiac tamponade. When the CTA was negative for PE, the team pursued tamponade as a possible diagnosis. Normally the lack of a pulsus paradoxus would argue against clinical tamponade physiology. However, this case is particularly unique for the echocardiographic finding of isolated RV collapse, leading to clinically significant regional tamponade. Collapse of a cardiac chamber occurs when pericardial pressure is greater than chamber pressure. Regional tamponade is often seen after procedures (pericardiectomy, cardiac surgery) or myocardial infarction where a loculated effusion or pericardial hematoma forms in a particular region. Regional cardiac tamponade fails to produce pulsus physiology because the increased pericardial pressure is localized to the RV. The RV’s inability to expand prevents interventricular septum from bowing into the left ventricle (LV) on inspiration. As a result, there is no compression of the LV, a phenomenon which is required for the clinical manifestation of pulsus paradoxus. This patient’s presentation of dyspnea, hypotension, and tachycardia with negative pulsus paradoxus but echocardiographic findings of pericardial effusion merited a high level of suspicion for regional cardiac tamponade.

Abstracts

Case reports I

Concurrent session
8:00 AM

Friday, January 21, 2022

#227 MALIGNANT EFFUSIONS: A RARE CASE OF SUBACUTE, SEVERE REGIONAL CARDIAC TAMPOANDE
N Shamapant*, C Duarte. University of Colorado, Denver, CO
10.1136/jim-2022-WRMC.224

Case Report Identify clinical manifestations of regional cardiac tamponade

Case Presentation The patient is a 31-year-old female with stage IV ER+, PR+, Her2- breast cancer complicated by recurrent malignant pleural effusions who presented with worsening dyspnea. Vital signs on admission included BP 121/91, HR 124, RR 32 and oxygen saturation of 99%. Shortly after admission, blood pressure dropped to 92/68 and oxygen saturation was 97% on 2L nasal cannula. On exam, patient had increased work of breathing with elevated neck veins. EKG showed low voltage QRS complexes with electrical alternans. CTA chest revealed a large pericardial effusion with no evidence of pulmonary embolus (PE). Given the pericardial effusion, a manual pulsus was checked and was negative. TTE showed a large pericardial effusion with moderate right ventricular (RV) collapse, dilated inferior vena cava and normal left ventricular systolic function. Based on the echocardiographic evidence of pericardial effusion with RV collapse in the context of worsening hypotension and tachycardia, the diagnosis of regional cardiac tamponade was made. Pericardiocentesis was performed with an opening pericardial pressure of 23mmHg and 680mL of fluid was drained with cytology confirming malignant pericardial effusion. Following the pericardiocentesis, the patient’s dyspnea and tachycardia resolved. Significant post-procedure pericardial drainage prompted placement of a pericardial window and the patient was discharged to hospice.

Discussion This patient’s presentation of dyspnea, tachycardia, progressive hypotension and distended neck veins fits multiple illness scripts including PE and cardiac tamponade. When the CTA was negative for PE, the team pursued tamponade as a possible diagnosis. Normally the lack of a pulsus paradoxus would argue against clinical tamponade physiology. However, this case is particularly unique for the echocardiographic finding of isolated RV collapse, leading to clinically significant regional tamponade. Collapse of a cardiac chamber occurs when pericardial pressure is greater than chamber pressure. Regional tamponade is often seen after procedures (pericardiectomy, cardiac surgery) or myocardial infarction where a loculated effusion or pericardial hematoma forms in a particular region. Regional cardiac tamponade fails to produce pulsus physiology because the increased pericardial pressure is localized to the RV. The RV’s inability to expand prevents interventricular septum from bowing into the left ventricle (LV) on inspiration. As a result, there is no compression of the LV, a phenomenon which is required for the clinical manifestation of pulsus paradoxus. This patient’s presentation of dyspnea, hypotension, and tachycardia with negative pulsus paradoxus but echocardiographic findings of pericardial effusion merited a high level of suspicion for regional cardiac tamponade.

Case Report Thrombotic events are a common complication of cancer but it is rare to have arterial thrombosis. This is an unusual case of left ventricular thrombus and multiple thrombi on the mitral valve while taking direct oral anticoagulants (DOAC) in a patient with stage IV gastroesophageal signet ring adenocarcinoma.

Methods Approval was obtained from the IRB at Kern Medical. A single patient chart review was conducted.

Summary A 45-year-old female with no past medical history presented to the emergency department (ED) with decreased appetite, fatigue, dysphagia, abdominal pain, and unintentional weight loss for 4 months. Initial workup with esophagogastro-duodenoscopy (EGD) showed an epigastric mass that was biopsied. The pathology came back showing stage IV gastroesophageal signet ring adenocarcinoma. During this hospitalization, the patient also had a computerized tomography (CT) of the chest done which showed pulmonary emboli (PE), and was started on apixaban for PE therapy.

Four weeks later, a CT of the chest, abdomen, and pelvis was ordered for evaluation of port-a-cath placement evaluation which demonstrated a filling defect within the left ventricular apex measuring 19 x 17 mm. The patient was referred for a transthoracic echocardiogram (TTE) which revealed a large left ventricular thrombus and multiple thrombi on the mitral valve. The patient was then admitted to the hospital for initiation of anticoagulation with therapeutic Lovenox. The patient was then discharged home with therapeutic Lovenox and instructions to follow up with cardiology outpatient.

Conclusion It is well known that there is a link between thromboembolism and cancer but the underlying mechanism is poorly understood. It is believed that there are many ways the cancer cells activate the coagulation system such as having the ability to produce and secrete procoagulant/fibrinolytic substances and inflammatory cytokines. Deep vein thrombosis (DVT) and pulmonary embolism (PE) are common complications in patients with cancer but arterial thrombosis secondary to malignancy is rare. The case highlights a rare presentation of a large left ventricular thrombus and multiple thrombi in the mitral valve in a patient with stage IV gastroesophageal signet ring adenocarcinoma that was already on a DOAC for a known PE.
The three cases at our hospital consisted of previously healthy male adolescents between the ages of 14 and 17 that presented with chest pain within 4 days of receiving their first or second mRNA covid-19 vaccine. On admission, they all had an abnormal electrocardiogram (EKG), elevated C-reactive protein (CRP), and elevated troponin I levels (table 1). Brain natriuretic peptide (BNP) levels were tested in two patients and found to be within normal limits. All patients had a normal echocardiogram, except for a residual patent foramen ovale in one patient. All patients showed down-trending troponin levels and inflammatory markers as well as complete resolution of symptoms before discharge on hospital day 4. All patients had normal troponin and CRP levels at post-discharge follow-up with an outpatient pediatric cardiologist visit. No causal relationship has been proven between the vaccine and myocarditis or pericarditis. Currently, the Centers for Disease Control and Prevention (CDC) continues to recommend the COVID-19 vaccination for anyone older than twelve years old as the known risks of illness and possible severe complications far outweigh having a possible rare adverse reaction. Suspected cases of myocarditis and pericarditis after COVID-19 vaccination should continue to be reported to VAERS.

### Abstract #229

**CARDIOVASCULAR MANIFESTATIONS IN ADOLESCENT MALES AFTER THE PFIZER COVID-19 VACCINE**

1M Arhin*, 2H Hamad, 3A Iheagwara, 4H Hales. 1University of North Carolina at Chapel Hill School of Medicine, Chapel Hill, NC; 2Geisinger Commonwealth School of Medicine, Scranton, PA; 3North Carolina State University, Raleigh, NC; 4HCA Holdings Inc, Nashville, TN

10.1136/jim-2022-WRMC.226

**Case Report**

The Food and Drug Administration (FDA) granted Emergency Use Authorization (EUA) for the Pfizer BioNTech mRNA (BNT162b2) and Moderna COVID-19 vaccines in December 2020 to address rising COVID-19 infections across the nation. Clinical trials showed the vaccines to be effective and safe. Once these vaccines became publicly available, some minor adverse events were reported to the Vaccine Adverse Event Reporting System (VAERS); they included fever, fatigue, and joint pain. Additionally, there have been cases of post-immunization myocarditis and/or pericarditis reported to VAERS.

This case series describes the clinical course of the first three adolescent patients in our hospital that presented with myocarditis or pericarditis within two to four days after administration of their first or second dose of the Pfizer mRNA covid-19 vaccine. The COVID-19 status of the patients was assessed by reverse transcription-polymerase chain reaction (RT-PCR) of nasopharyngeal swabs. Laboratory tests for all patients included a routine complete blood count, basic metabolic panel, troponin, and inflammatory markers. Two patients had testing for SARS-CoV-2/Covid-19 and other viral causes of myocarditis and pericarditis, including the Epstein-Barr virus, cytomegalovirus, parvovirus B19, respiratory syncytial virus, influenza A virus, influenza B virus, Roseola, and human herpesvirus 6. These tests were all negative.

### Abstract #230

**METASTATIC AND PERSISTENT BACTEREMIA WITH METHICILLIN-RESISTANT STAPHYLOCOCCUS AUREUS, ASSOCIATED WITH COVID-19 PNEUMONIA, A MANAGEMENT NIGHTMARE**

1C Berman*, 2C D’Asumpcao, 3J Fong, 4R Kuran, 5A Heidari. 1American University of the Caribbean School of Medicine BV, Cupecoy, Sint Maarten (Dutch part); 2Kennedy Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.227

**Case Report**

Telavancin is a vancomycin-derivative semisynthetic lipoglycopeptide that has antimicrobial activity against resistant gram-positive organisms, namely methicillin-resistant *Staphylococcus aureus* (MRSA). The purpose of this study is to describe a case in which telavancin clinically failed in treating persistent MRSA bacteremia in a patient.

**Methods Used**

Retrospective case review.

**Summary of Results**

A 61-year-old man with nasal MRSA colonization and history of previous abscesses was initially admitted for severe COVID-19 pneumonia requiring high flow nasal cannula oxygen support and dexamethasone protocol as per current guidelines. He developed MRSA pneumonia and persistent polyclonal resistant MRSA bacteremia with hematogenous seeding causing lumbar vertebral osteomyelitis despite treatment with vancomycin for 5 days follow with telavancin for 11 days. MRSA susceptibility to the glycopeptide antibiotics as well as cefaroline and daptomycin were evaluated using E-test to interpret minimum inhibitory concentration (MIC) according to the manufacturer’s instructions. Telavancin MIC was initially 0.064 but then on retest was 0.125. Since MRSA continued to grow in repeated blood cultures after 11 days, telavancin was considered to have clinically failed. He was switched to combination of cefaroline and daptomycin and subsequently developed daptomycin associated cosinophilic pneumonitis. He was started on prolonged course of prednisone. He was switched to cefaroline and rifampin to complete 6 weeks total of antibiotics after blood sterilization. Outpatient MRSA decolonization protocol was also started just prior to discharge.
Conclusion Secondary bacterial infection associated with COVID-19 is on the rise particularly after adoption of dexamethasone as standard of care in severe cases. Persistent bacteremia with MRSA complicated with metastatic seeding in this setting is not described and perhaps is due to host-pathogen mediated mechanisms. Clinical failure of telavancin in deep seeded MRSA infections has not yet been reported.

A SEVERE CASE OF RECURRENT HERPES SIMPLEX VIRUS-1 ENCEPHALITIS WITH AUTOIMMUNE COMPONENT

1VF Civelli*, 2C D’Assumpcao, 3M Kaur, 4R Kuran, 5K Sabetian, 6A Heidari. 1Clínica Sierra Vista, Bakersfield, CA; 2Kern Medical Center, Bakersfield, CA; 3Ross University School of Medicine, Miramar, FL

10.1136/jim-2022-WRMC.228

Case Report The incident of Human Herpes Simplex Virus-1 (HSV-1) encephalitis is 4 in one million cases worldwide. From that 5–27% can become recurrent a rare phenomenon with serious sequel. Differentials include induced autoimmune-mediated sequelae or paraneoplastic encephalitis. Knowing that 69–89% of first HSV-1 encephalitis will result in permanent neurological deficits, the degree of damage in recurrent cases is estimated to be higher. The pathophysiology of recurrent HSV-1 encephalitis and its aftermath are poorly understood. We report a case of recurrent HSV encephalitis with evidence of autoimmune dysfunction.

Methods Used Retrospective Study

Summary of Results A 58-year-old Hispanic female presented elsewhere for acute onset of confusion. She was diagnosed with HSV-1 encephalitis by PCR testing of the CSF and received 28 days on intravenous (IV) Acyclovir and recovered. She presented to our facility with new onset of seizures, expressive aphasia, short-term memory loss, bowel/bladder incontinence, and unsteady gait. Her mental status declined, and diagnosis of autoimmune encephalitis was made due to negative HSV-1 PCR and brain MRI findings. A 5-day course of IV Immunglobulin and high dose methylprednisolone resulted in transient relief of her symptoms. She was discharged home with prednisone for 5 weeks. Autoimmune CSF panel showed elevated acetylcholine receptor ganglioside (alpha 3) Ab at 172, and elevated VGCC Type P/Q Ab at >30, and VGCC Type N Ab at 69. She had two episodes of recurrent neurological symptoms resulted in courses of IV high dose steroid with transient relief each time CSF only was positive for mild pleocytosis with negative HSV-1 PCR. Her brain MRI continued to show progressive disease now spreading to both sides. On the third recurrence episode of rapid decline in mentation CSF came back positive for HSV-1 PCR. She was started on a 3-week course of IV Acyclovir. Follow up CSF was negative for HSV-1 PCR and she is placed on prolong course of oral acyclovir with unknown duration perhaps lifelong.

Conclusion Differential diagnosis of recurrent encephalitis after the first episode of HSV-1 infection includes immune-mediated HSV-1 viral encephalitis vs recurrence of HSV-1 itself. Residual irreversible neurocognitive deficits are expected and may result in protracted disease. The diagnosis is challenging and prognosis is poor.

HUMAN HERPESVIRUS-6 MENINGOECEPHALITIS IN AN IMMUNOCOMPETENT MALE

1VK Narang*, 2C D’Assumpcao, 3M Valdez, 4K Radić, 5I Mosavi, 6R Kuran, 7A Heidari. 1UCLA-Kern Medical, Bakersfield, CA; 2Kern Medical-UCLA, Bakersfield, CA

10.1136/jim-2022-WRMC.229

Case Report Meningoencephalitis with Human herpesvirus 6 (HHV-6) in adults is rare. It is known to occur as a result of reactivation of infected dormant brain cells during childhood, when host becomes immunocompromised in conditions such as organ or bone marrow transplantations. New primary infection in adults has also been described. Diagnosis is difficult and challenging particularly in immunocompetent adults due to low level of suspicious and undefined clinical, central spinal fluid (CSF) and neuroimaging findings. The duration and choice of antiviral also have not been well reported. We present a case of HHV-6 meningoencephalitis in an immunocompetent elderly male who presented with fever and altered mental status.

Methods A single patient case report was conducted after IRB approval.

Case Presentation Patient is a 79-year-old male with a history of dementia and diabetes mellitus who presented after being found down by a bystander. Patient arrived lethargic, tachypneic, and febrile to 39.4 °C. Physical exam was remarkable for rigid extremities with laboratory studies demonstrating a leukocyte count of 14.6x10^3 cells/mm3 without left shift and hyperglycemia to 550 mg/dL. He was empirically placed on antibiotics to cover bacterial etiologies plus acyclovir. CT of the head demonstrated moderate to severe global volume loss and periventricular leukomalacia. CSF showed WBC count of 17 with 76% monocytes and 22% lymphocytes. Extensive initial work up and all cultures came back negative. MRI brain with gadolinium found atrophic and chronic microangiopathic changes without enhancement. CSF Meningoencephalitis Panel (BioFire, BioMerieux, Salt Lake City, Utah) found HHV-6 positivity. HHV-6 serology testing found negative IgM and positive IgG. Serum HHV-6 PCR confirmed the diagnosis with > 2 million copies/mL. Patient was started on ganciclovir which resulted in significant improvement in symptoms. He was discharged to rehabilitation facility to complete a 30 day course of ganciclovir.

Conclusion HHV-6 meningoencephalitis is a serious but rare condition particularly in otherwise immunocompetent adults. Clinicians should be aware of this infection when initial work up is not diagnostic.

A CASE OF CONCURRENT DISSEMINATED COCCIDIOIDOMYCOSIS AND EMBRYONAL CARCINOMA WHEN LICE AND FLEAS COEXIST

M Ke, A Heidari, M Valdez*, A Tsyrier, R Kuran, R Johnson. Kern Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.230

Case Report Coccidioidomycosis (CM) is a fungal infection endemic to the southwestern United States with a wide range of clinical presentations depending on the infected organ systems. CM causes a primary pulmonary infection. 1 percent of cases disseminate, via hematogenous or lymphatic spread. It is in these cases, that more severe symptoms may present and potentially overlap with those characteristics of other systemic
illnesses. We report a case of CM disseminated to lymph nodes in a 24-year-old man with concomitant metastatic embryonal carcinoma. It is difficult to identify the primary etiology for many components of his presentation and the relationship between these concurrent disease processes is not entirely clear. Factors that may contribute include locus minoris resistentiae or a shared immune response between infectious organisms and malignant cells.

Case Description 24-year-old man from the central valley of California presented with a new rash. Examination showed periocular edema and diffuse raised, erythematous, hyperpigmented skin lesions. Differential included psoriasis and CM. CM serology showed nonreactive IgM, very weakly reactive IgG, and complement fixation (CF) titer <1:2. The patient was lost to follow-up but returned 6 weeks later with 17-pound weight loss, progression of the rash, and proximal muscle weakness. He had a heliotrope rash and elevated CK. He was prescribed prednisone 60 mg daily for dermatomyositis. 2 weeks later, he developed fevers and CM CF titer was 1:16. He was started on fluconazole 800 mg daily. 5 weeks later, he developed diffuse lymphadenopathy and imaging showed multiple pulmonary nodules, destructive lesion in the iliac bone, and retroperitoneal and pelvic lymphadenopathy. CM CF titers were now 1:64 and he was started on liposomal amphotericin B. New retroperitoneal and right testicular masses were then identified. Histopathology from orchectomy and retroperitoneal mass biopsy revealed embryonal carcinoma while inguinal lymph node excision showed granulomatous inflammation with endosporulating spherules diagnostic for CM. He completed 9 weeks of amphotericin and bone scan showed no foci of increased uptake. He is on track to complete 4–6 cycles of chemotherapy. On subsequent clinic visits, weight was up-trending and improvement in both rash and lymphadenopathy were noted.

Conclusion As CM and several types of malignancies may have similar or overlapping presentations, a thorough examination and tissue sampling are generally necessary to distinguish. In rare cases, coexistence may occur. ‘Läuse und Flöhe haben.’ A German phrase, which translates to ‘Having lice and fleas,’ refers to having two reasons for a problem. Understanding the etiology or identifying the relationship between the concomitant conditions is essential to formulate the most appropriate treatment plan.

#235 GASTRIC MUCORMYCOSIS IN AN ADOLESCENT WITH NEW ONSET DIABETES MELLITUS

J. Timberman, C. Kupekan. Valley Children’s Healthcare, Madera, CA
10.1136/jim-2022-WRMC.235

Case Report Mucormycosis is a rare angioinvasive fungal infection associated with a high mortality rate, especially in children and neonates. Recent data suggests the incidence is rising in the United States. Risk factors include diabetes, glucocorticoids, hematological malignancies and iron overload. The most frequent types of mucormycosis include rhinoorbitocerebral, pulmonary, and cutaneous disease.

Here we present a child with newly diagnosed type II diabetes mellitus who developed hematemesis and was found to have gastrointestinal mucormycosis.

Methods Used Case Report

Summary of Results A 17-year old obese male presented to the emergency department with a 1-day history of abdominal pain, vomiting and respiratory distress. Weight loss, polyuria and polydipsia were reported. He had no other significant medical or surgical history. On examination his temperature was 37.8 C, pulse 137, BP 116/77, RR 44, and Pox 96% in...
Endocrinology and metabolism II
Concurrent session
8:00 AM
Friday, January 21, 2022

#236 NEURAL CORRELATES OF OBESITY AND INFLAMMATION IN YOUTH WITH CLASSICAL CONGENITAL ADRENAL HYPERPLASIA

1 1MS Kim*, 2A Pickering, 3DL Cotter, 1NR Fraga, 4S Luo, 1C Won, 1M Gefney, 1M Herting, 2Children’s Hospital of Los Angeles, Los Angeles, CA; 3Children’s Hospital of Los Angeles Saban Research Institute, Los Angeles, CA; 4University of Southern California Keck School of Medicine, Los Angeles, CA

Purpose of Study Patients with classical congenital adrenal hyperplasia (CAH) due to 21-hydroxylase deficiency exhibit an increased prevalence of obesity over their lifetime. We have previously shown that CAH youth exhibit smaller gray matter brain region volumes, including the prefrontal cortex (PFC), amygdala, and hippocampus, as well as white matter microstructure abnormalities. Here, we aim to study the relationship between these patterns of altered brain structure with obesity and inflammation in youth with and without CAH.

Methods Used We studied 27 youth with CAH (12.6 ± 3.4 yr, 16 females) and 35 controls (13.0 ± 2.8 yr, 20 females) via 3-T MRI, and examined the PFC and its subregions, amygdala, and hippocampus regions of interest (ROIs), as well as white matter tracts including the fornix and stria terminalis (ST). Relaimpo (relative importance for linear regression: $R^2_{\text{pool}}$) analyses identified body composition and inflammatory markers, as well as gray and white matter ROIs most associated with CAH status. Regression analyses examined associations between brain structure, CAH, and other variables.

Summary of Results Waist-to-height ratio (WHtR; $R^2_{\text{pool}} = 0.12$) and monocyte chemotactic protein-1 (MCP-1 $R^2_{\text{pool}} = 0.21$) exhibited the strongest associations with CAH status amongst body composition variables and inflammatory markers respectively. Analysis of gray matter ROIs showed the PFC was most strongly associated with CAH status ($R^2_{\text{pool}} = 0.13$). Superior frontal (SF) was the only PFC subregion associated with all key variables: CAH status ($\beta = -0.58$, $P = 0.001$), WHtR ($\beta = -0.23$, $P = 0.009$), and MCP-1 ($\beta = -0.35$, $P < 0.001$). CAH youth exhibited significantly larger WHtR and MCP-1, and smaller SF volume than controls ($P < 0.001$ for all). White matter microstructure showed fornix and ST associations with CAH (fornix $\beta = 0.15$, $P = 0.02$; ST $\beta = 0.22$, $P < 0.001$) and WHtR (fornix $\beta = 0.36$, $P = 0.01$; ST $\beta = 0.35$, $P = 0.02$), but only ST associations with MCP-1 ($\beta = 0.30$, $P = 0.02$).

Conclusions Youth with CAH demonstrate an important relationship between altered gray and white matter brain structure, WHtR, and MCP-1 that could have implications for neuroinflammation and obesity in CAH.

#237 POPULATION-BASED ASSESSMENT OF CARDIOMETABOLIC-RELATED DIAGNOSES IN YOUTH WITH KLINEFELTER SYNDROME: A PEDSNET STUDY

S Davis*, N Nokoff, A Furniss, A Valentine, L Pyle, A Dempsey. University of Colorado, Denver, CO

Purpose of Study Diabetes and cardiovascular diseases are common among men with Klinefelter syndrome (KS) and contribute to higher morbidity and mortality. Cardiometabolic health outcomes have not been evaluated in a large population of youth with KS.

Methods Used Data from electronic health records from six pediatric institutions were used to compare the prevalence of five cardiometabolic-related outcomes among 1,080 youth with KS to 4,497 youth without KS matched for sex, age (mean ± SD), year of birth, race, ethnicity, insurance, site, and duration of care (mean ± SD). Odds ratios (OR) and 95% confidence intervals (CI) both unadjusted and adjusted for potential covariates including obesity, testosterone, and antipsychotic use were computed using generalized estimating equations.

Summary of Results The odds of overweight/obesity (OR 1.6 (95%CI 1.4–1.8)), dyslipidemia (3.0 (2.2–3.9)), and liver dysfunction (2.0 (1.6–2.5)) were all higher in KS compared to controls. While the adjusted model attenuated the effect of KS on these outcomes, boys with KS still had 45% greater odds of overweight/obesity (CI 1.2–1.7) and 70% greater odds of liver dysfunction (1.3–2.2) compared to controls, and both dyslipidemia (1.6 (1.1–2.4)) and dysglycemia (1.8 (1.1–3.2)) were higher in KS but of borderline statistical significance when accounting for multiple comparisons. The odds of
hypertension were not different between groups in unadjusted or adjusted models.

Conclusions This large, population-based cohort of youth with KS found a higher odds of many cardiometabolic-related diagnoses compared to matched controls. Investigation into the mechanisms that underlie risk for cardiometabolic dysfunction in youth with KS independent of obesity and testosterone is needed.

#238 REACTIVE HYPOGLYCEMIA FOLLOWING A SUGAR CHALLENGE IS ACCOMPANIED BY HIGHER INSULIN IN ADOLESCENT GIRLS WITH OBESITY

1,2MA Ware*, 1A Careau, 1Y Garcia-Reyes, 1,2H Rahat, 1,2Diniz Behn, 1M Cree-Green. 1University of Colorado – Anschutz Medical Campus, Aurora, CO; 2Rocky Vista University, Parker, CO; 3Univ Laval Faculté de médecine, Quebec, QC, Canada; 4Colorado School of Mines, Golden, CO

10.1136/jim-2022-WRMC.235

Purpose of Study Adolescent girls can present with postprandial, episodic symptoms of dizziness and excess sweating, which are often clinically disregarded. These symptoms can be associated with idiopathic reactive hypoglycemia (RH), reproduced with a glucose challenge, and may relate to increased risk of insulin resistance and type 2 diabetes (T2D). We sought to determine the prevalence of and glycemic measures associated with RH among adolescent girls with obesity following an oral sugar tolerance test (OSTT).

Methods Used Secondary analysis of 112 adolescent girls with obesity (age 12–21 yrs; body mass index (BMI) ≥90th percentile). Participants completed a 4-hr OSTT (75 g glucose, 25 g fructose) and were stratified by glucose nadir between 120–240 mins: ≥60 mg/dL as RH, ≥80 mg/dL as normoglycemic (NG), and 61–79 mg/dL as indeterminate. Area under the curve (AUC) of glucose, insulin, glucagon, and C-peptide and measures of insulin sensitivity and β-cell function including oral minimal model, insulinogenic index, and oral disposition index were calculated. Demographic, physical, and metabolic characteristics between RH and NG were compared with Student’s t-tests or Mann-Whitney U tests. OSTT curves were compared with repeated measures ANOVA.

Summary of Results 12% of girls had RH (n=13) and 36% had NG (n=40). Groups were similar in age, race, ethnicity, and BMI. Glucose concentrations were lower in RH than NG when fasting (p=0.033), at 210 mins (p=0.001), and 240 mins (p<0.001). In RH, glucose nadir occurred at a median time of 210 min postprandial. RH had higher glucagon AUC for 90–240 min of the OSTT (p=0.035) and a lower insulin at 240 mins (p=0.007) confirming a counterregulatory response. Insulin AUC (p=0.074) and C-peptide AUC (p=0.054) for 0–120 min trended to be higher. Groups had similar hemoglobin a1c, family history of T2D, insulin sensitivity, and β-cell function.

Conclusions RH was relatively common in this cohort of adolescent girls with obesity and occurred at 3–4 hrs. Early higher insulin secretion preceded the counterregulatory RH responses but was not related to insulin sensitivity. Future studies should aim to further understand the long-term effects of RH during adolescence and its significance in predicting future hyperglycemia.

#239 DIETARY QUALITY AND ONE-YEAR PROGRESSION OF GLUCOSE INTOLERANCE IN SUBJECTS WITH OBESITY


10.1136/jim-2022-WRMC.236

Purpose of Study Previous longitudinal cohort studies have shown that improved dietary quality is correlated with a reduction in the risk of cardiovascular disease and potentially diabetes. There has been limited research into whether a similar relationship holds for the progression of glucose intolerance. The hypothesis for this study is that better dietary quality is correlated with reduced progression in glucose intolerance over the course of one year in people with obesity.

Methods Used This was a prospective cohort study. Participants (N=38) at baseline completed three 24-hour dietary recalls. The majority (N=37) were obese and subjects with diabetes were excluded. Each participant’s Healthy Eating Index–2015 (HEI) score was calculated from the dietary recalls. The HEI is a measure of dietary quality that assigns weights to meeting adequacy targets (i.e., getting sufficient vegetables) and not exceeding moderation targets (i.e., limiting saturated fat) for a total score between 0 and 100. All participants underwent a 2-hour 75-gram glucose tolerance test at baseline and at 12 months. Glucose tolerance was calculated as the percentage change in the incremental area under the curve (iAUC) between these two time points. Linear regression analysis in STATA was used to determine the relationship between HEI score at baseline (and other dietary measures), and progression of glucose intolerance over one year. One statistically influential outlier with percentage change in glucose iAUC of >150% was excluded.

Summary of Results A trend was observed showing that higher dietary quality by HEI score was correlated with less progression of glucose intolerance (ß = -1.27, R² = 0.102, p = 0.034). When age was added as a covariate, the values were (ß = -1.21 R² = 0.103, p = 0.098). Higher saturated fat intake was correlated with greater progression of glucose intolerance (ß = 5.07, R² = 0.168, p = 0.012). There were no statistically significant relationships between vegetable, added sugar, or dietary fiber intake and progression of glucose intolerance.

Conclusions In this small study of people with obesity, higher baseline dietary quality as measured by HEI correlated with less progression of glucose tolerance with borderlineline statistically significant (p = 0.034). Higher saturated fat intake correlated with greater progression of glucose intolerance (p = 0.012). The clinical relevance of this work would be to help optimize dietary recommendations to prevent progression of glucose intolerance which can be seen before manifestation of overt disease. Limitations of this study include the small number of subjects, lack of normal weight subjects, limited follow-up period, and self-reported dietary data. Future directions of research could include increasing sample size, studying normal weight subjects over a longer timeframe, and exploring methods for collecting dietary data that do not rely on self-report.
ORAL CONTRACEPTIVE USE IN ADOLESCENTS WITH POLYCYSTIC OVARY SYNDROME AND OBESITY IS ASSOCIATED WITH ALTED FAT METABOLISM

1E Finn*, 1C Severn, 1Y Garcia-Reyes, 1MA Ware, 1H Rahat, 1M Cree-Green. 1University of Colorado, Aurora, CO; 1Children’s Hospital Colorado, Aurora, CO

Purpose of Study Polycystic ovary syndrome (PCOS) is a common endocrine condition characterized by reproductive and metabolic hormone dysregulation and increased risk for metabolic disease. Combined oral contraceptive pills (OCPs) are first-line treatment for management of hyperandrogenic symptoms and menstrual regulation, yet data on the cardiometabolic effects in youth are limited.

Methods Used Participants were enrolled in a cross-sectional trial of girls aged 12–21 years, BMI% >90th ile with PCOS per NIH 1990 criteria (NCT02157974). Measurements included anthropomorphics, fasting metabolic and hormonal assessments, 6-hour oral sugar tolerance test (OSTT), hepatic fat per MRI, DXA for body composition, and vascular function by EndoPAT and dnapulse. Area under the curve (AUC) for responses to the OSTT were calculated, as were several indices of insulin sensitivity. Participants treated with OCPs for at least 6 months were matched 1:2 with replacement with untreated participants on BMI, race, and ethnicity. The effect of treatment was estimated with paired t-tests.

Summary of Results Data from 10 participants taking OCPs (age 15.7 ± 1.3 years, BMI 34.2 ± 5.6 kg/m²) were compared to that from 20 untreated girls (age 15.6 ± 1.9, BMI 33.6 ± 5.0). The OCP group had decreased free testosterone (p=0.001), increased sex hormone binding globulin (p=0.001), and decreased free androgen index (p=0.001). The OCP group had higher total cholesterol (p=0.045), body fat percentage (p=0.028), and serum leptin (p=0.038). The OCP group also had higher free fatty acids AUC (p=0.007) and glyceral AUC (p=0.011) during the OSTT. No differences between the groups were noted in glucose metabolism including fasting glucose and insulin, HOMA-IR, c-peptide, glucose AUC, or insulin AUC. Vascular function and hepatic fat fraction were similar between groups.

Conclusions Treatment with OCPs was not associated with worsening dysglycemia or markers of vascular dysfunction. However, there was evidence of altered fat metabolism, including a shift towards greater fat mass and adipose insulin resistance with persistent lipolysis. Additionally, hepatic fat was not improved. The overall therapeutic benefit of OCP’s in girls with PCOS and obesity who are at high cardiometabolic risk needs to be carefully evaluated.

ACIPIMOX PREVENTS CARDIAC DYSFUNCTION INDUCED BY ELECTRONIC CIGARETTES IN MICE

1CJ Lao*, 1J Espinoza-Derout, 1K Hasan, 1MC Jordan, 1J Wilon, 1J Molina, 1K Luna, 1JL Arambulo, 1A Sinha-Hikim, 1K Roos, 1T Friedman. 1Charles Drew University of Medicine and Science, Los Angeles, CA; 1University of California Los Angeles David Geffen School of Medicine, Los Angeles, CA

Purpose of Study Tobacco use is a leading cause of preventable death in the United States. In recent years, electronic cigarettes (e-cigarettes) use by youth has augmented at a disturbing rate. Cardiac dysfunction can lead to heart failure, which is responsible for 287,000 deaths each year in the United States. In a mouse model, we will analyze whether acipimox treatment can rescue the cardiac dysfunction phenotype caused by e-cigarettes with 2.4% nicotine through the inhibition of lipolysis.

Methods Used For 12 weeks, mice were exposed to saline, e-cigarettes (2.4%), and e-cigarettes plus acipimox in 12-hour cycles through specialized chambers. Acipimox (0.05%) was delivered in the drinking water. Mouse weights and food consumption data were collected weekly. Echocardiograms were used to analyze heart function.

Summary of Results Weight data indicates that acipimox decreases the rate of weight change over time when compared to both saline control and e-cigarette groups. Acipimox treatment rescued e-cigarette induced phenotypes in fractional shortening, ejection fractions and velocity of circumferential fiber shortening. Triglyceride levels were increased in e-cigarette groups but were not fully rescued after acipimox treatment. Acipimox treatment reduced circulating levels of free fatty acids (FFA), Macrophage Colony-Stimulating Factor (M-CSF), and Interleukin-6 (IL-6) when compared to e-cigarette treatment alone. Gene set enrichment analysis (GSEA) indicates that genes involved in the G2/M checkpoint were upregulated by e-cigarette treatment and rescued by acipimox treatment. Heme oxygenase 1 (HO1) and Poly(ADP-Ribose) Polymerase 1 (PARP1) protein levels were rescued after acipimox treatment.

Conclusions E-cigarette use led to an increase in free fatty acids which resulted in increased inflammation and reactive oxygen species within cardiac tissue. This resulted in DNA damage in the heart and halting of the cell cycle at the G2/M checkpoint. The DNA repair pathway was activated with increased levels of PARP1, a protein known for cellular recovery. This led to the cardiac dysfunction phenotype after e-cigarette use. Acipimox rescued this phenotype by inhibiting lipolysis and preventing downstream effects of dysfunctional adipose tissue and increased FFAs on cardiac tissue. In conclusion, the data obtained so far shows that lipolysis is necessary for the e-cigarette induced cardiac dysfunction. Lipolysis may become a new therapeutic target to curve the harmful cardiovascular effects produced by e-cigarettes.

ACCESS TO CARE FOR METABOLIC SYNDROME AND CARDIOVASCULAR DISEASE AMONG IMMIGRANT POPULATIONS IN THE UNITED STATES

J Beltran*, M Shaheen, D Pan. Charles Drew University of Medicine and Science, Los Angeles, CA

Purpose of Study It is estimated that one-quarter to one-third of adults meet the Metabolic Syndrome (MetS) criteria in the United States. MetS could be responsible for approximately 7% of total mortality and up to 17% of cardiovascular disease (CVD). This research aims to determine the relationship between metabolic syndrome, cardiovascular disease, and access to care by immigration status.

Methods Used We analyzed data from the National Health and Nutrition Examination Survey (NHANES) 2017–2018 using bivariate chi-square and multiple logistic regression, considering the design and sample weight. We presented the data as an adjusted odds ratio and 95% confidence interval, and a p-value of <0.05 will be considered statistically significant.
Abstracts

Summary of Results Of the 5,265 adults, 19.3% were immigrants, 8.7% had CVD, and 34.3% had MetS. Immigrants had lower prevalence of CVD (5.4%) compared to non-immigrants (9.5%) (p<0.05). A higher percentage of immigrants were uninsured (24%) compared to non-immigrants (11%) (p<0.05). In the multiple logistic models, there was no significant difference in the odds of CVD, MetS, and access to care among immigrants relative to non-immigrants after adjusting for the confounding variables. Though, in four of the five risk factors for MetS, immigrants had higher odds of CVD than non-immigrants (p<0.05).

Conclusions Though the prevalence of CVD is not statistically different from non-immigrant, immigrants are more susceptible to the risk factors of MetS which increases the likelihood of developing CVD. Resources need to be distributed to the vulnerable immigrant population to decrease the risk factors of MetS, CVD and increase access to healthcare.

#243 CAN MEDICAL CARE PROVIDERS PREVENT HEART ATTACKS AND STROKES?

J Giannini*, J Padilla, RP Eaton, K Gonzales, DS Schade. University of New Mexico School of Medicine, Albuquerque, NM

10.1136/jim-2022-WRMC.240

Purpose of Study Cardiovascular disease prevention is an important goal of medical care providers. In the clinical setting, we addressed whether myocardial infarctions and strokes can be prevented in asymptomatic patients with proven subclinical cardiovascular disease.

Methods Used Two hundred six asymptomatic self-selected individuals requested a coronary artery calcium scan (CAC). Of these individuals, 125 had positive scans (Agatston score >1) and were recommended for medical treatment to prevent heart attacks and atherosclerotic strokes. Eighty-one individuals had zero scores and were not treated. Of the 125 individuals with positive scans, 110 followed medical treatment (rosuvastatin 10 mg/d, ezetimibe 10 mg/d, and a low cholesterol diet) and achieved a low density lipoprotein cholesterol (LDL-C) of ≤60mg/dl. The other 15 individuals did not follow recommended treatment and maintained an elevated LDL-C.

Summary of Results All patients were followed for an average of 3.5 years (minimum 1 year – maximum 5 years). No atherosclerotic cardiovascular events were observed in the group that achieved an LDL-C ≤60 mg/dl. The expected number of atherosclerotic cardiovascular events based on their calcium scores and published observational data was 12.6. Two patients died of non-atherosclerotic causes (an auto accident and Alzheimer’s disease). In the non-compliant group, two heart attacks were observed (one patient died, one had a stent). No events were observed in the zero CAC group. No adverse effects of the treatment regimen occurred.

Conclusions Medical care providers can prevent atherosclerotic heart attacks and strokes by treating asymptomatic patients who have positive calcium scores to an LDL-C goal of ≤60 mg/dl. A simple regimen of rosuvastatin, ezetimibe, and a low cholesterol diet has no major adverse effects, is inexpensive, and is acceptable by almost all patients.

Gastroenterology

Concurrent session

8:00 AM

Friday, January 21, 2022

#244 DIABETIC HYPERGLYCEMIA EXACERBATES INFLAMMATORY BOWEL DISEASE IN MICE WITH DIET-INDUCED OBESITY

1-3K Francis*, 1MC Pacheco, 1K Alonge, 2BA Phan, 2S Hu, 2N Schwartz, 1,2J Scarlett. 1Seattle Children’s Hospital, Seattle, WA; 2University of Washington School of Medicine, Seattle, WA

10.1136/jim-2022-WRMC.241

Purpose of Study Inflammatory bowel disease (IBD), obesity, and type 2 diabetes (T2D) are chronic inflammatory conditions that are increasing in prevalence. Obesogenic high-fat diets (HFD) that promote diet-induced obesity (DIO) and T2D have been shown to worsen IBD. Furthermore, comorbid T2D in patients with IBD is a predictor of poor disease-related outcomes. To determine the mechanisms whereby T2D worsens IBD pathology, we examined the impact of hyperglycemia on disease activity in a DIO mouse model of IBD, focusing on measures of intestinal barrier integrity.

Methods Used Wild-type C57BL/6J mice were fed HFD for 8 weeks to induce DIO and then given intraperitoneal injections of vehicle or low-dose streptozotocin (STZ) to induce hyperglycemia. At week 12, 2% dextran sodium sulfate (DSS) or vehicle control was administered in drinking water for 7 days to induce colitis. Outcomes included serial measures of body weights, blood glucose, and disease activity index (DAI) scores, which incorporate weight loss, rectal bleeding, and stool consistency. Animals were euthanized at the end of DSS course and colon tissue was collected and processed for immunohistochemical staining.

Summary of Results As predicted, STZ administration raised mean blood glucose levels significantly compared to vehicle-treated controls (267mg/dL vs. 159mg/dL, p=0.006). Hyperglycemic mice treated with DSS (DSS/STZ) manifested clinical signs of colitis 2 days earlier than their normoglycemic counterparts (DSS/Veh), and their DAI scores were significantly higher from Days 2–7 of DSS (p<0.0001). Colon length and histologic damage scores also demonstrated higher disease burden in DSS/STZ mice (colon length 5.93cm vs. 6.88cm, p=0.0003; histologic score 8.0 vs 6.5, p=0.03). Colonic tissues demonstrated significantly decreased quantity of the tight-junction protein E-cadherin (E-cad) in DSS/STZ mice compared to DSS/Veh (percent area 24.1 vs. 34.9, p=0.004), and...
a similar decrease in colonic mucin barrier was detected by Alcian Blue (AB) staining (percent area 55.3 vs. 71.0, p<0.0001). Notably, measures of both colonic E-cad and AB were inversely correlated with both DAI scores and blood glucose levels.

Conclusions In DIO mice, clinical and pathological IBD outcomes are exacerbated by moderate hyperglycemia in a DSS model of colitis. We identify two potential mechanisms by which hyperglycemia contributes to intestinal pathology: 1) reduced tight-junction protein quantity in the colonic epithelium and 2) a decrease in the protective colonic mucin barrier. These results support the hypothesis that hyperglycemia induces intestinal barrier dysfunction, which, in combination with an obeseogenic diet, increases the risk for IBD progression.

Purpose of Study Intraoperative cholangiography (IOC) is frequently performed during cholecystectomy and considered definitive testing for evaluation of common bile duct (CBD) stones. Previous studies evaluating the accuracy of IOC for the evaluation of choledocholithiasis are inconsistent, and false-positive testing may lead to increased utilization of diagnostic ERCP. The aim of this study is to evaluate the diagnostic property of IOC and to assess the clinical outcomes of patients who underwent ERCP with or without retained choledocholithiasis.

Methods Used Consecutive hospitalized patients who received cholecystectomy with IOC at Loma Linda University Medical Center (1/2014–6/2021) were identified. Positive IOC was defined by the presence of filling defect, meniscus sign, and/or failure to drain contrast into the duodenum necessitating additional diagnostic evaluations. Definitive diagnosis of choledocholithiasis was defined by the presence of choledocholithiasis and/or sludge on any of the definitive testing for choledocholithiasis (MRCP, EUS, and/or ERCP) performed after IOC. Patients who had prior sphincterotomy or negative IOC without documented ≥2-year follow-up were excluded.

Summary of Results The mean age of 259 patients who underwent same-day cholecystectomy and IOC was 46.0±19.8 years, 189 (73%) were female, and 144 (56%) were of Latino ethnicity/race. On presentation, 22 (9%) patients had bilirubin ≥4.0 mg/dL, 111 (43%) had a dilated CBD ≥6 mm, and 7 (3%) had choledocholithiasis detected on abdominal ultrasound. During cholecystectomy, 103 (40%) patients had abnormal IOC including the presence of filling defect in 94 (36%), meniscus sign in 2 (1%), and failure of passage of contrast to the duodenum in 7 (3%). Subsequently 122 (46%) patients had definitive testing including ERCP in 102 (39%), MRCP in 45 (17%), and EUS in 14 (5%). Diagnostic property of IOC for choledocholithiasis are shown in the table 1. ERCP was performed at a median of 2 days (range, 0–141 days) from IOC. Of 102 patients, 8 (3%) had adverse events including pancreatitis in 6 (moderate in 3, mild in 3) and perforation in 2; five patients with adverse events received diagnostic ERCP for evaluation of false-positive IOC.

Conclusions Although IOC demonstrated high sensitivity and negative predictive value of >90% for ruling out retained CBD stones, nearly a third of the patients received diagnostic ERCP. Given adverse events associated with ERCP, other less invasive definitive tests such as MRCP and EUS should be considered for evaluations of patients with positive IOC prior to ERCP.

Purpose of Study To determine whether ethnicity and other risk factors can predict the severity of fibrosis or cirrhosis in women with NAFLD applying the four predictor models FIB-4, NFS, BARD, and APRI.

Methods Used We performed a retrospective study among Hispanic (n=567) and Caucasian (n=136) women with history of NAFLD (mean age 55.7 ± 11.4 SD years) at The University of California Davis Medical Center. Univariate analyses for their FIB-4, NFS, BARD, and APRI scores were conducted by using Chi-Square or Fisher’s Exact Test for categorical variables, two-sample t-test for continuous variables, and Wilcoxon-Mann-Whitney test for continuous non-parametric variables. Associations between ethnicity (Hispanic vs. Caucasian) and liver fibrosis severity calculated using the four cirrhosis predictor models were explored using backward selection multinomial logistic regression.

Summary of Results We observed that Hispanic women compared to Caucasian showed lower BMI (p<0.001), higher HDL (p<0.05), higher HbA1c (p<0.001), higher prevalence of stage 4 and 5 CKD (p = 0.005), lower prevalence of bariatric surgery (p=0.002), lower likelihood to smoke (p=0.002), and had lower APRI score (p=0.03). The percentage of advanced fibrosis in each models and ethnicity was as follows: Hispanic NFS: 35.1%, BARD: 82.6%, FIB-4: 3.5%, and APRI: 1.3%; in Caucasian NFS: 36.3%, BARD 83.2%, FIB-4: 7.7%, APRI: 6.3%. The odds of having a severe
fibrosis or cirrhosis was not significantly higher in those who are Hispanic compared to Caucasian in FIB-4, NFS, BARD, and APRI (p=0.16, 0.78, 0.97, and 0.18 respectively).

**Conclusions** Based on our preliminary results, we cannot conclude that ethnicity alone can be used to predict the severity of fibrosis or cirrhosis in women with NAFLD in the multiple predictor models.

**Abstracts**

### #247 EXAMINING CLINICAL PRESENTATION AND WORKUP OF VETERANS WITH IRRITABLE BOWEL SYNDROME IN A SINGLE MEDICAL CENTER

1PL Claassen*, 2M Riddle. 1Washington State University Elson S Floyd College of Medicine, Spokane, WA; 2VA Sierra Nevada Health Care System, Reno, NV

10.1136/jim-2022-WRMC.244

**Purpose of Study** Irritable Bowel Syndrome (IBS) is a disorder of gut-brain interaction (DGBI) that affects about 5% of the population with significant quality of life and economic impacts. Recent ACG guidelines have been established to improve diagnosis and management of IBS. Within the Veteran population, there is poor understanding of the epidemiology of DGBI despite significant occupation exposures, comorbid mental health problems and risk of deployment-associated diarrhea. Less is known about the management of DGBI in Veterans compared to other populations.

**Methods Used** As part of a quality improvement initiative, a case series analysis of Veterans seen at a new IBS Clinic was performed to better understand the patient population and their preceding diagnostic workup. Data from the VA Electronic Health Record and a standardized IBS Clinic intake form was retrieved. All charts were comprehensively reviewed to determine the dates of gastrointestinal (GI) symptom onset, first functional GI disorder diagnosis, first IBS diagnosis, as well as basic demographics, current comorbidities and medications. Descriptive analytic methods were applied.

**Summary of Results** A total of 11 Veterans referred to the IBS Clinic were included with a median age of 43 years, all male, majority Caucasian (81.8%) and predominantly from the Persian Gulf war (72.7%), post-Vietnam (18.2%) and Vietnam era (9.1%). A median of 4 (IQR 2.5–14.5) years delay between time of symptom onset to first functional GI disorder diagnosis was noted. Of those diagnosed with IBS, the phenotypes were diarrhea (57.1%), mixed (28.6%) and unclassified (14.3%). Patients had a median of 10 (IQR 6–15) comorbidities and took a median of 9 (IQR 5–10) medications (i.e. selective serotonin reuptake inhibitors) which could contribute to diarrhea. Mental health problems were very common, with 55% having post-traumatic stress disorder. Additionally, 60% of patients presented with at least one red flag symptom that warranted colonoscopy. With respect to ACG guidelines, less than 50% received testing for celiac disease and inflammatory bowel disease. Only 20% received unnecessary stool ova or parasite testing and 10% an unneeded colonoscopy.

**Conclusions** IBS is understudied in Veterans. Many of these patients have psychological comorbidities and take medications which may contribute to their symptoms. While some aspects of their workup aligned well with guidelines, others did not. Future efforts are needed to improve diagnosis and management of Veterans with DGBI.

### #248 RISK FACTORS ASSOCIATED WITH INCREASED FIBROSIS AND STEATOSIS IN NON- ALCOHOLIC FATTY LIVER DISEASE UTILIZING TRANSIENT ELASTOGRAPHY (FIBROSCAN) IN A COMMUNITY MULTI SPECIALTY PRACTICE

1B Stratford*, 2C Gonzales. 1Weber State University College of Science, Ogden, UT; 2Ogden Clinic, South Ogden, UT

10.1136/jim-2022-WRMC.245

**Purpose of Study** Non-alcoholic fatty liver disease (NAFLD) is associated with increased risk of fibrosis in patients with obesity, diabetes, and metabolic syndrome. Noninvasive liver assessment using FibroScan is an important part of the evaluation of these patients. The aim of this study is to assess the efficacy of the FibroScan in helping to risk stratify patients with NAFLD in order to improve patient outcomes.

**Methods Used** Over a 16 month period (3/2020 to 6/2021), 331 patients with suspected NAFLD underwent liver FibroScan assessment. A retrospective chart review was conducted for risk factors of metabolic syndrome (obesity, HTN, diabetes, sleep apnea, hyperlipidemia), FibroScan measurements of liver stiffness (MLS), steatosis, and liver biopsy outcomes.

**Summary of Results** A total of 331 patients were reviewed with a mean age of 49 years and mean BMI of 35 (range 18–58). Fifty-three percent of patient were obese (BMI > 30) and 20% were morbidly obese (BMI > 40). Metabolic risk factors were common; HTN (45%), hyperlipidemia (37%), sleep apnea (31%) and diabetes (23%). Greater fibrosis risk was seen with increased BMI (normal vs. morbidly obese BMI, mean MLS 5.5 kPa vs. 9.9 kPa, p < 0.001), metabolic risk factors (0 vs. 5 risk factors, mean MLS 5.7 kPa vs. 13.5 kPa, p < 0.0003) and presence of diabetes (no diabetes vs. diabetes, mean MLS 7.3 kPa vs. 10.3 kPa, p < 0.003). Increased steatosis scores were seen in increased BMI (normal vs. morbidly obese BMI, mean CAP 236 dB/m vs. 334 dB/m, P < 0.0001), metabolic risk factors (0 vs. 5 risk factors, mean CAP 263 dB/m vs. 340 dB/m, p < 0.0001) and presence of diabetes (no diabetes vs. diabetes, mean CAP 307 dB/m vs. 333 dB/m, p < 0.0005). Twenty-five patients underwent liver biopsy evaluation and all patients were diagnosed with non-alcoholic steatohepatitis (NASH). Stage II-III fibrosis was diagnosed in 12 patients (48%) and stage 4 bridging fibrosis/cirrhosis in 3 patients (12%).

**Conclusions** This cohort of patients undergoing FibroScan testing in a community practice were mostly obese and had >1 metabolic syndrome risk factors. Elevated liver stiffness and steatosis were significantly correlated with increasing BMI, presence of diabetes and increased risk factors for metabolic syndrome. These results will help guide clinician utilization of FibroScan testing in clinical practice.

### #249 ANTIPHOSPHOLIPID ANTIBODY SYNDROME IN A YOUNG MALE WITH BILIARY ADENOCARCINOMA

1H Ipalawatte*, 2S Sathian, 3K Radicic, 4S Mishra. 1Xern Medical Center, Bakersfield, CA; 2Ross University School of Medicine – Barbados Campus, Bridgetown, Barbados

10.1136/jim-2022-WRMC.246

**Case Report** Cholangiocarcinoma(ACC) is a rare cancer of the biliary system, common in elderly patients. Risk factors vary including alcoholic liver disease, hepatitis, HIV infection, or parasitic infection. Common associations include ulcerative
Ileal Endocrine Tumor Without Metastasis

V Rendica, J Raheesh, B Quraishi, S Pathapati. Texas Tech University Health Sciences Center School of Medicine, Amarillo, TX

Case Report A 26-year-old male with no known medication history of hypertension and recent substance abuse presented to the ED with 2 months of watery diarrhea consisting of 3–4 episodes a day. He was transferred from an outside facility for acute renal failure, severe metabolic acidosis, diarrhea, dehydration, and hypokalemia.

Initial episodes of watery diarrhea presented with fever, chills, and nausea. Watery diarrhea increased to about 10 episodes a day during this initial course. He also reported generalized weakness and recurrent falls due to syncope. He self-medicated Ivermectin in high doses meant to be used for horses that he was able to buy from his hometown veterinarian with no prescription meant to treat his diarrhea as well as prevent Covid-19 infections. He used this drug for 10 days and reported a 20 pounds weight loss over 2 months as well as worsening oliguria.

Follow this initial course, his diarrhea continued at a lesser rate of only 3–4 episodes a day. However, three days prior to presenting to the ED he noticed fresh blood in his stool, prompting him to seek medical attention. On admission he denied abdominal pain, flushing, dyspnea, or chest pain. On examination, he had mild diffuse abdominal tenderness. The patient denied any history of hepatitis, pancreatitis, peptic ulcer disease or GERD. He had never had a colonoscopy or endoscopy at this point.

On admission his labs showed acute renal failure with an SCr of 18, a BUN of 161, and bicarb of 6. After he was started on fluid resuscitation his kidney function improved and urinary output increased, and his acute kidney injury eventually resolved.

His CT abdomen was remarkable for ‘misty mesentery’ suggestive of pancreatitis. There was also an enhancing nodularity of the distal small bowel mesentery in the right mid abdomen with mild to moderate circumferential bowel wall thickening of the underlying small bowel loops. In addition to this there were scattered diverticula of the sigmoid colon with no diverticulitis. The carcinoid tumor was also able to be identified on CT.

Colonoscopy showed a large 3 cm submucosal polypoid nodular mass at 10 cm distance from ileocecal junction. Biopsy reported a well differentiated neuroendocrine tumor (carcinoid), spanning at least 2 mm involving the lamina propria and submucosa. Liver ultrasound showed only hepatic steatosis with no lesions. 24hr urine HIAA was 12.5, chromogranin A 1048ng/mL and serum serotonin level of 83.8ng/mL. TTE showed no valvular abnormalities.

Right hemicolectomy with small bowel mass was resected, surgical pathology reported well differentiated neuroendocrine tumor (carcinoid), spanning at least 2 mm involving the lamina propria and submucosa. Liver ultrasound showed only hepatic steatosis with no lesions. 24hr urine HIAA was 12.5, chromogranin A 1048ng/mL and serum serotonin level of 83.8ng/mL. TTE showed no valvular abnormalities.

Purpose of Study Indeterminate colitis (IC) is a form of Inflammatory Bowel Disease (IBD), when the diagnosis of Ulcerative colitis (UC) or Crohn’s disease cannot be made due to mixture of findings. The incidence of Clostridioides difficile (C. diff) infection is 6–9% in patients with IBD, which can lead to increased risk for colectomy or death. Here we discuss a case of newly diagnosed IC with superimposed C. diff infection.

Methods Used Retrospective chart review after IRB approval. Summary of Results A 26-year-old male with no known medical history presented to our facility with progressive one month history of abdominal cramping pain and bloating. This was associated with watery hematochezia up to 15 episodes
per day. Upon admission, he was found to have fever as high as 39.4°C, WBC 7.5, CRP 23.8 mg/dL, ESR 68 mm/hr, Hgb 12.7 g/dL. His work up also showed stool Calprotectin 3520 mcg/g, positive stool Lactoferrin, and atypical P-ANCA titer of 1:160, commonly elevated in UC. Stool studies were also positive for C.diff toxin B PCR and GDH antigen, and oral vancomycin was started. Colonoscopy with biopsies showed mixture of findings, from 2 pathology readings, for UC and CD with focal active colitis with focal ulcerations (consistent with UC), involvement of the crypts, lamina propria, and submucosa (consistent with CD). The diagnosis of IC was made and he was started on Methylprednisolone 60 mg TID and Mesalamine 1600 mg BID. Patient eventually improved with less frequency of hematochezia. Upon follow up outpatient clinic his was down to only 5 episodes of diarrhea without hematochezia. Follow up colonoscopy when C.diff infection is resolved is planned for making a definitive diagnosis between UC and CD.

Conclusions Diagnosis of Ulcerative Colitis vs Crohn’s Disease can be challenging in the setting of superimposed infection with Clostridiodes difficile. Management of this coexistence is difficult, and duration of treatment is also not well studied. Further work up and repeat biopsy might be needed for definitive diagnosis.

Session: genetics I
Concurrent session
8:00 AM
Friday, January 21, 2022

#252 DPH5: A NOVEL GENE CAUSING DIPHTHAMIDE BIOSYNTHESIS DISORDERS

SP Shankar*, K Grilmerud, L Lanoue, A Egenese, B Willis, P Shankar, J Horberg, L Alabed, K Mayer, K Otkar, M Monaghan, J Krie, R Stoier, N Schaffrath, K Alkuraya, U Brinkmann, L Erikson, K Lloyd, KA Raun. University of California Davis, Sacramento, CA; University of California Davis, Davis, CA; Goteborgs universitet, Goteborg, Sweden; King Faisal Specialist Hospital and Research Center, Riyadh, Saudi Arabia; Hoffmann-La Roche AG Research and Development Division, Penzberg, Germany; Genex, Gaithersburg, MA; Universitat Kassel Institut fur Biologie, Kassel, Germany; Boston Children’s Hospital, Boston, MA

Purpose of Study Neurodevelopmental disorders (NDDs) are genetically heterogeneous lifelong conditions with a known etiology in approximately 50% of individuals. Here, we report DPH5 (Diphthamide biosynthesis protein 5) as a novel cause of embryonic lethality, multisystem dysfunction and profound NDDs in three unrelated families. DPH5 is critical to the biosynthesis of diphthamide, a post translationally modified histidine on eEF2 (eukaryotic elongation factor 2), essential for ribosomal translation and protein synthesis in cells.

Methods Used Exome or genome sequencing, transgenic Dph5 phH260R knockin (C57BL/6Ncl-DphS<sup>emb<sup>1Mmucd</sup></sup> mouse model development, patient recruitment through GeneMatcher program, ADP-ribosylation assays in DPH5ko cells transfected with pN110S, pN174fs, pR207*, and p260R, site-directed mutagenesis in yeast cells for the missense variants and functional assays, computational modeling to evaluate effect of variants on DPH5 protein structure and assess its interaction with eEF2.

Summary of Results DPH5 variants, homozygous pH260R, compound heterozygous pN110S and pR207* and homozygous pN174fs were found in three unrelated families (F1, 2 & 3). Similar craniofacial features such as broad forehead, sparse eyebrows, epicanthal folds, short upturned nose, downturned corners of the mouth, profound NDDs, CNS anomalies such as enlarged cisterna magna, cardiac features such as tricuspid septal defect, pulmonary stenosis and pericardial effusion, and digital anomalies were noted. Dph5 pH260R targeted knockin mouse model generated only one homozygous (HOM) female exhibiting extremely low birthweight, craniofacial dysmorphism, polydactyly and abnormal behaviors with early death at 24-days of age. Embryonic studies of HOMs revealed abnormal head shape, exencephaly, eye anomalies, edema shortened frontronatal prominence, facial cleft and polydactyly. ADP ribosylation assays showed absent to decreased function in KO and yeast cells. Insilico modeling showed disruption of interaction of DPH5 with eEF2.

Conclusions We provide clinical and functional evidence that DPH5 variants in three unrelated families are likely pathogenic establishing DPH5 as a novel cause of embryonic lethality or profound NDDs with multisystem involvement expanding the Diphthamide biosynthesis disorders.

#253 MACROANGIOPATHY, NOT SO COMMON IN MELAS?

JP Ramos*, JA Morales, M Manning, Stanford Medicine, Stanford, CA

10.1136/jim-2022-WRMC.250

Purpose of Study MELAS (mitochondrial encephalopathy, lactic acidosis, and stroke-like episode) is a multisystem disorder caused by a known mitochondrial DNA mutation most commonly affecting the MT-TL1 gene. The majority of individuals develop symptoms between the ages of 2 to 40 years old.

When MELAS affected patients present with stroke-like symptoms, the expected etiology is the underlying mitochondrial disorder causing a stroke-like episode. Common clinical findings include stroke-like episodes, encephalopathy with seizures or altered mental status, muscle weakness, hearing impairment, peripheral neuropathy amongst others. MELAS is a condition with phenotypic variability due to multiple causes (heteroplasmia, tissue distribution and threshold effect).

There are approximately 13 reported cases of macroangiopathy in patients with MELAS. We report a previously undiagnosed cause of MELAS in an adult patient who presented with right middle cerebral artery (MCA) ischemic stroke and found to have severe intracranial atherosclerosis, a finding not typically reported in MELAS patients.

Methods Used Chart review, physical examination, and literature review

Summary of Results A 44 year-old female was evaluated for a possible mitochondrial condition given a past medical history of sensorineural hearing loss following physiologically demanding circumstances (pregnancy), type 1 diabetes and acute neurologic symptoms concerning for a stroke after removal of a cochlear implant. Molecular testing revealed a mtDNA pathogenic variant: 3243A>G (tRNA Leu gene) with 24.7% heteroplasmy, a finding highly indicative of MELAS. Head CT angiography found severe intracranial atherosclerosis involving large vessels requiring thrombectomy. Imaging noted a tapering
occlusion in a large segment of the right MCA, multifocal irregularity and moderate stenosis of the left internal carotid artery and proximal left middle cerebral artery, with diffusely narrowing caliber of the internal carotid arteries.

Conclusions Of the MELAS reported cases, intracranial large vessel involvement (macroangiopathy) is a rare finding in MELAS, described in very few individuals. Macroangiopathy can include stenosis, vasocostriction, atherosclerosis, and aneurysms, and theoretically could be a lesser-known finding in patients with chronic MELAS. Given the severity of clinical complications in our patient, we call attention to macroangiopathy as a possible complication in patients with MELAS to inform their clinical team when preparing for invasive procedures, especially those involving intracranial vascular system.

**#254 MT-ATP6-ASSOCIATED MITOCHONDRIAL DISEASE CAN PRESENT WITH LOW CITRULLINE BY NEWBORN SCREENING AND MORE: DEFINING THE BIOCHEMICAL PHENOTYPE**

1CG Tise*, 1C Lee, 1B Mendelsohn, 1,2J Woods, 1K Cusmano-Ozog, 2Stanford University, Stanford, CA; 2Kaiser Permanente, Oakland, CA; 3Valley Children’s Healthcare, Madera, CA

Purpose of Study MT-ATP6 encodes a subunit of mitochondrial complex V (ATP synthase). Pathogenic variants are associated with a form of Leigh syndrome exhibiting considerable phenotypic heterogeneity and maternal inheritance. We have identified seven children from five families with MT-ATP6-associated disease who presented with low citrulline by California newborn screening (NBS). Further evaluation revealed abnormalities on plasma amino acids (PAA), urine organic acids (UOA), and acylcarnitine profile (ACP), and in several cases, maternal relatives with a similar phenotype. We report these findings and characterize the biochemical phenotype associated with MT-ATP6-associated disease.

Methods Used Medical literature and chart review, physical examination, and laboratory testing.

Summary of Results Seven probands were found to have a maternally-inherited homozygous pathogenic variant in MT-ATP6 (m.8993 T>G, p.L156R) after presenting with abnormal NBS for low citrulline. Biochemical evaluation revealed PAA with low citrulline and high alanine, UOA with elevations of lactate and 3-hydroxyisovaleric acid without orotic aciduria, and ACP with elevations of propionylcarnitine (C3) and 3-hydroxyisovalerylarnitine (C5-OH). Overall findings were suggestive of a mitochondrial disorder, as opposed to the intended primary screening target of a proximal urea cycle disorder, warranting mitochondrial DNA studies. Some maternal relatives were found to have the same biochemical phenotype as the probands (n=3), as well as clinical features of this disorder including ataxia, neuropathy, myopathy, and/or retinitis pigmentosa; others are reportedly asymptomatic.

Conclusions MT-ATP6-associated disease can present with low citrulline by NBS and a unique biochemical phenotype consisting of the triad of low citrulline, elevated C3, and elevated C5-OH, in addition to evidence of mitochondrial dysfunction by PAA and UOA. For this reason, we recommend including ACP in the evaluation of individuals who screen positive by NBS for low citrulline; maternal history and laboratory studies should also be considered. Future studies include metabolomic profiling of NBS dried blood spots from individuals with MT-ATP6-associated disease and relevant negative and positive controls to further delineate the biochemical phenotype and aptly identify individuals with this mitochondrial disorder.

**#255 ZELLWEGGER SPECTRUM DISORDER IN CALIFORNIA’S CENTRAL VALLEY: EVIDENCE OF A FOUNDER MUTATION IN MIXTECO PATIENTS AFFECTED WITH A NOVEL PEX6 VARIANT**

1KR Wong*, 1,2J Woods, 1J Carmichael, 1C Galarreta Aima, 1Valley Children’s Hospital, Madera, CA; 2Stanford Medicine, Stanford, CA

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Purpose of Study Zellweger spectrum disorders (ZSD) are a group of autosomal recessive disorders caused by mutations in the PEX genes essential for peroxisomal biogenesis. Affected patients have defective lipid metabolism and cell detoxification, resulting in hypotonia, neurologic deficits, congenital malformations, adrenocortical dysfunction, and liver disease. Prevalence varies among different regions of the world and founder mutations in distinct populations have previously been reported. California’s Central Valley has a large Mixteco population compared to other parts of the United States as many of these families have emigrated from the southern Mexico states of Oaxaca, Guerrero, and Puebla. We have observed an increased frequency of ZSD at our center over the past few years in patients of Mixteco ethnicity. This study reports the spectrum of clinical and genotypic features of ZSD patients at our institution.

Methods Used We performed a retrospective chart review by searching for ZSD patients seen at our center between 2010 and 2020 using ICD-10 codes E71.50, E71.51, E71.511, E71.518, E71.53 and E71.54, and ICD-9 code 277.86. Patients with an alternative diagnosis were excluded.

Summary of Results Seven patients with ZSD were identified, all diagnosed over the past four years, presenting at birth and with generalized hypotonia and facial dysmorphisms such as large anterior fontanelles. Two had poor visual response to light, three had lagophthalmos, and four had failed hearing tests. Four patients are now deceased and three died prior to one year of age. Although two were lost to follow-up, they both presented with severe symptoms including worsening hypotonia and poor feeding by three months of age. One patient is currently alive at 12 months of age. Biochemical testing demonstrated characteristic elevations of very long chain fatty acids. Six patients were found to be homozygous for the PEX6 novel variant c.1409G>C (p.Gly470Ala) and were of Mixteco ethnicity. One was homozygous for PEX6 c.2095–21_2095del and was the only patient with parental consanguinity. This patient’s ethnicity is unknown.

Conclusions ZSD has a prevalence of 1:50,000 to 1:75,000 in the United States. In the Central Valley, we found an increased frequency of ZSD per zip code population in which these patients lived, ranging from 1:4,477 to 1:72,280, with a mean of 1:44,891. This over-representation of the novel PEX6 variant affecting ZSD patients of Mixteco ethnicity suggests a founder mutation within this patient population.
Purpose of Study CRB2-related syndrome was first described as a triad of cerebral ventriculomegaly, renal findings including nephrotic syndrome (NS), and greatly elevated alpha-fetoprotein levels. Additional clinical manifestations in CRB2-related syndrome have included congenital heart defects and retinitis pigmentosa (RP). The condition is rare and caused by biallelic, pathogenic variants in the CRB2 gene. Recent reports of CRB2-related syndrome have highlighted NS which can be associated with severe presentations. The objective of this project was to compile a list of the clinical and variant data for CRB2-related syndrome.

Methods Used We conducted a literature review of reported patients with biallelic pathogenic variants in CRB2 and identified additional unreported cases. We compared clinical features, survival, and variant location in CRB2 in patients with manifestations in different body systems. For those with NS, treatments and their effectiveness were recorded.

Summary of Results We ascertained 31 patients with biallelic, pathogenic variants in CRB2; detailed information was available for the majority. 18/29 (62%) were diagnosed during pregnancy and 9/29 (31%) from 0 to 10 years of age. Two patients were diagnosed at 46 and 51 years and had RP as the sole clinical finding. Of the 30 patients with recorded outcomes, 20 (66.7%) were alive, whereas 10 (33.3%) were deceased. 7 from termination of pregnancy. An analysis of clinical findings showed that renal involvement was the most frequent (22/31; 71%), with 11 (35%) patients having only renal manifestations with NS. 17/31 (55%) patients had manifestations in multiple body systems, and cardiac findings and abnormal retinas were each found in 3/31 (10%). Pathogenic variants were mostly located in exons 7, 8, 10, 12, and 13, with greatest representation of exon 10 in all patients, whereas exon 7 had the greatest representation in patients with only renal disease. Details of treatments and their effectiveness were inconsistently reported and prevented comparison, but one patient transiently responded to steroids and only one surviving patient required a renal transplant.

Conclusions Most patients with CRB2-related syndrome included in this review survived. The commonest finding in recently reported patients with pathogenic CRB2 variants was NS. Further information is needed to determine optimal treatment and patient care.

Purpose of Study Many regions in Asia including Taiwan hold a high thalassemia carrier rate. The diagnosis of thalassemia in these regions mainly consists of a screening algorithm relying on mean corpuscular volume (MCV) and mean corpuscular hemoglobin (MCH) values. However, individuals who are silent carriers and who do not phenotypically fit the screening algorithms tend to go undiagnosed, leading to future complications. While genetic screening panels may be useful in this context, a standardized regime is limited in this region. Thus, this study evaluates the clinical significance of implementing a standardized genetic screening panel for α-thalassemia. Overall, the study assesses the efficacy of this screening in identifying phenotypically asymptomatic patients who were otherwise undiagnosed, and the risk posed in passing down more severe carrier traits.

Methods Used This is a cross-sectional study utilizing next-generation sequencing (NGS) carrier screening panels to identify α-thalassemia silent carriers in a population within Taiwan. Long range PCR is followed to confirm patient’s genetic status.

Summary of Results A total of 22 patients were identified as thalassemia silent carriers. 21 out of 22 individuals (95.5%) were α-thalassemia silent carriers, with 2 of them being couples. Therefore, in this population set there is a 9.5% chance of passing down more severe alpha thalassemia carrier traits. Furthermore, only 1 out of 22 patients (4.5%) were found to be β-thalassemia carriers. These individuals were asymptomatic, and all held a MCV of >80. Of these individuals, only 4 were identified with a low hemoglobin level.

Conclusions Our data suggests the implementation of a standardized genetic screening for α-thalassemia will be beneficial in many regions within Asia, including Taiwan. Previous screening algorithms focused on measuring MCV and MCH levels. However, in this study we demonstrate that MCV and MCH levels alone may be insufficient in identifying carriers. Maintenance of this screening algorithm would thus result in false negatives within this patient population. Overall, the use of NGS-based carrier screening panels allowed for the diagnosis of individuals who were previously undiagnosed due to them not phenotypically conforming to the current regional screening algorithms. This study also allowed for the identification of individuals at an increased risk of passing down traits such alpha thalassemia carriers and hemoglobin H (HbH). The utilization of genetic screening for α-thalassemia would be cost-efficient and beneficial especially in these patients by allowing for earlier genetic counseling, education, and potential treatment to prevent costly complications later in life.

Purpose of Study Psychiatric illness is a pressing health concern and there is a greater need to understand genetic differences between influencing traits. This project analyzed genome-wide differences among three genome-wide association studies (GWAS), aiming to understand genetic differences between patients with depression and bipolar disorder. Being influenced by several genes and confounding factors complicates the genetic study of these polygenic illnesses. One tool available to help understand genome wide differences from GWAS is linkage disequilibrium score regression (LDSC) which allows

Abstracts

#256 CLINICAL FINDINGS IN CRB2-RELATED SYNDROME

K Mai, K Chen*. Western University of Health Sciences, Pomona, CA

10.1136/jim-2022-WRMC.256

#257 ASYMPTOMATIC THALASSEMIA: A REASON TO REFORM CURRENT SCREENING ALGORITHMS IN REGIONS OF ASIA

K Mai, K Chen*. Western University of Health Sciences, Pomona, CA

10.1136/jim-2022-WRMC.257

#258 LINKAGE DISEQUILIBRIUM SCORE REGRESSION ANALYSES TO DETERMINE THE GENETIC ARCHITECTURE OF THE DIFFERENCES BETWEEN DEPRESSION AND BIPOLAR DISORDER

S Asaf*. University of California Los Angeles, Los Angeles, CA

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for the analysis of differences among psychiatric patient cases who present with bipolar disorder (BP) or major depressive disorder (MDD) with different onsets.

Methods Used Three GWAS files were analyzed through LDSC: daner13 (case-only BP vs. MDD file), danerD (BP-onset-MDD vs. MDD file), and danerM (BP-onset-mania vs. MDD file). Unix and Python shell scripting were used to run a series of awk commands and the ldsc command-line tool MDD file). Unix and Python shell scripting were used to run a series of awk commands and the ldsc command-line tool MDD file), and danerM (BP-onset-mania vs. MDD file). Unix and Python shell scripting were used to run a series of awk commands and the ldsc command-line tool MDD file), and danerM (BP-onset-mania vs. MDD file).

Summary of Results The genetic correlation of 16 traits for the traits amongst two cohorts. The three GWAS were then compared and educational traits along with the heritability of traits a series of awk commands and the ldsc command-line tool MDD file). Unix and Python shell scripting were used to run a series of awk commands and the ldsc command-line tool MDD file), and danerM (BP-onset-mania vs. MDD file), and danerM (BP-onset-mania vs. MDD file).

Conclusions Typical intelligence, branchial sinus anomalies, and amastia/athelia in females appear to be distinct features of KMT2D-associated disorder. This syndrome presents with features overlapping with other multiple malformation syndromes including Kabuki, CHARGE, and brachiootooreal syndrome. These individuals exemplify the distinct phenotype of KMT2D-associated disorder and highlight the need for a natural history study of this condition to allow for better characterization and ultimately improved diagnosis and outcomes.

### Neonatology general III

#### Concurrent session

**Friday, January 21, 2022**

**#260** PREDICTIVE VALUE OF PATENT FORAMEN OVALE MEASURED AT BIRTH FOR DEVELOPMENT OF ATRIAL SEPTAL DEFECT IN EXTREMELY LOW BIRTHWEIGHT INFANTS

S Gaffar*, B Siasi, R Cayabyab, M Ebrahim, M Uzunyan, R Ramanathan. Los Angeles County University of Southern California Medical Center, Los Angeles, CA

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Purpose of Study The foramen ovale (FO) is a fetal channel that allows right-to-left (R-L) shunting throughout fetal development. After birth, some term infants have minor incompetence of the flap valve, allowing a small left to right (L-R) shunt which usually resolves by 18 days of postnatal life. The aim of this study is to evaluate the echocardiographic change in FO size with postnatal growth to determine if progression to atrial septal defect (ASD) can be predicted in extremely low birth weight infants (ELBW).

Methods Used Descriptive study of all inborn ELBW infants admitted to a neonatal intensive care unit between 2015 and 2021 who had echocardiograms (ECHO) performed within...
the first week of postnatal life and before discharge. Size of valve-incompetent FO was determined by measuring the width of L-R color doppler flow in coronal posterior and sagittal subcostal viewing windows. The largest measurement was taken as the diameter of the septal defect.

Summary of Results Fifty-two infants with median gestational age (GA) of 25 weeks (IQR: 24–26) and median birth weight (BW) of 733 grams (IQR: 633–835) were included in the study. First ECHO was obtained at a median postnatal day of life 2. Forty-eight infants had initial median FO diameter of 1.9 mm at birth that grew to 2.2 mm by median postmenstrual age (PMA) of 36 weeks. Majority of infants at birth (37/46, 80%) and at discharge (39/44, 89%) had L-R shunt. Only 1 infant (1.9%) at birth and 4 infants (7.7%) at discharge had a closed FO. In contrast, 4 ELBW infants born at median GA of 25 weeks (IQR: 24–26) and median BW of 700 grams (IQR: 615–748) had initial FO diameter of 3.3 mm that grew into a 6.2 mm septal defect by 38 weeks PMA. All 4 infants had L-R shunt at FO on discharge. Table 1 shows other echocardiographic parameters measured that were within normal limits.

Conclusions This study demonstrated that in ELBW infants, FO greater than 3 mm in diameter at birth enlarges with postnatal growth to possibly become a large atrial septal defect that needs echocardiographic follow up as outpatient. If these large defects remain as atrial septal defects on long term follow up, it will indicate a much higher incidence of ASD in ELBW infants compared to full term infants. Further study of a larger population of ELBW infants is needed to confirm this finding.

Abstract #261 SYSTEMIC AND END ORGAN HEMODYNAMIC CHANGES DURING BRADYCARDIC EVENTS IN INFANTS UNDERGOING THERAPEUTIC HYPOTHERMIA

D Cho*, T Wu. Children’s Hospital of Los Angeles, Los Angeles, CA; University of Southern California Keck School of Medicine, Los Angeles, CA

Purpose of Study Therapeutic hypothermia (TH) is the standard therapy for newborn hypoxic-ischemic encephalopathy (HIE). Lower temperature slows firing of sinoatrial node and can lead to sinus bradycardia. The heart rate threshold at which end organ perfusion may be compromised is unknown. We aim to investigate systemic and end-organ hemodynamic changes during bradycardic episodes.

Methods Used Continuous hemodynamic data, including heart rate (HR), mean arterial blood pressure (MBP), cardiac output (CO) by impedance cardiometry, and regional brain (CrSO2) and renal (RsSO2) oxygen saturation were collected prospectively and time-synchronized in infants with HIE undergoing TH (figure 1). Systemic vascular resistance (SVR) was derived using measured CO and MBP. Bradycardia was defined as sustained HR≤100 beats per minute (<1st percentile for age) for 3 minutes. For each bradycardic episode, we selected a corresponding 3-minute period when HR>100, within a 10-minute time frame of the bradycardic event. Hemodynamic parameters were compared between the two groups (HR≤100 vs. HR>100) and reported as medians with IQR (non-normally distributed). Mann-Whitney test was used for non-parametric comparisons.

Summary of Results Twenty infants (9 male, 11 female, mean GA 38 5/7 weeks) were included in the analysis. Cumulatively, 433 bradycardic episodes were noted and occurred in 13% (21.7 out of 160.7 hours) of the overall monitoring period.
DECREASING PERCENT OF INFANTS WITH VERY LOW 5-MINUTE APGAR SCORES AT A DENVER SAFETY-NET LEVEL III NICU

Abstract #261 Figure 1 Sample data collection for 1 patient to illustrate continuous monitoring of hemodynamic parameters (MAP: mean arterial blood pressure, SpO2: oxygen saturation, HR: heart rate, CO: cardiac output, Ch1rSO2: brain oxygen saturation, Ch2rSO2: renal oxygen saturation).

There was a significant decrease in MBP and CO, and a compensatory increase in SVR during bradycardic episodes (table 1). There was no significant difference in SaO2, CrSO2 or RrSO2.

Conclusions Although there were systemic hemodynamic alterations during bradycardia events, overall end organ perfusion and/or oxygenation remain unchanged.

Abstract #262 Table 1 Short- and long-term outcomes for infants with very low 5-minute Apgar scores

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Baseline (n=66)</th>
<th>Intervention (n=37)</th>
<th>Post (n=35)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hypothemia (%)</td>
<td>7 (11)</td>
<td>6 (16)</td>
<td>7 (20)</td>
<td>0.42</td>
</tr>
<tr>
<td>HIE (%)</td>
<td>16 (24)</td>
<td>11 (30)</td>
<td>7 (20)</td>
<td>0.16</td>
</tr>
<tr>
<td>HIE mild (%)</td>
<td>6 (9)</td>
<td>1 (3)</td>
<td>0</td>
<td>**</td>
</tr>
<tr>
<td>HIE moderate (%)</td>
<td>6 (9)</td>
<td>4 (11)</td>
<td>3 (9)</td>
<td>**</td>
</tr>
<tr>
<td>HIE severe (%)</td>
<td>4 (6)</td>
<td>5 (14)</td>
<td>4 (11)</td>
<td>**</td>
</tr>
<tr>
<td>HIE unknown (%)</td>
<td>4 (6)</td>
<td>5 (14)</td>
<td>4 (11)</td>
<td>**</td>
</tr>
<tr>
<td>Seizures (%)</td>
<td>4 (6)</td>
<td>5 (14)</td>
<td>7 (20)</td>
<td>0.10</td>
</tr>
<tr>
<td>Gastrostomy tube (%)</td>
<td>2 (3)</td>
<td>2 (5)</td>
<td>3 (9)</td>
<td>0.48</td>
</tr>
<tr>
<td>Conventional ventilation (%)</td>
<td>36 (55)</td>
<td>27 (73)</td>
<td>22 (63)</td>
<td>0.18</td>
</tr>
<tr>
<td>Long-term</td>
<td>Cerebral Palsy (%)</td>
<td>3 (5)</td>
<td>3 (8)</td>
<td>1 (3)</td>
</tr>
<tr>
<td>Developmental Delay (%)</td>
<td>11 (17)</td>
<td>10 (27)</td>
<td>14 (40)</td>
<td>**</td>
</tr>
<tr>
<td>Speech Delay (%)</td>
<td>17 (26)</td>
<td>12 (32)</td>
<td>10 (29)</td>
<td>**</td>
</tr>
</tbody>
</table>

** p-value not calculated due to missing data

Purpose of Study Low Apgar scores are a strong predictor of neonatal mortality and may be associated with poor neurologic outcomes. Our hospital’s percentage of infants born with very low Apgar scores at 5 minutes was higher than the Vermont Oxford Network national average (2.4%). We aimed to decrease the percentage of infants with Apgar scores <4 at 5 minutes of life from a mean of 5.2% (range 1–11%) to less than 2.5% and decrease the percentage of infants receiving chest compressions prior intubation from 21% to less than 5% by December 2017 and sustain through August 2021.

Methods Used Data was collected from January 2012 to August 2021. Four plan-do-study-act (PDSA) cycles were done from April 2015 through February 2017 including: formulated a multidisciplinary team (PDSA 1), provided 24-hour pediatric advanced life support (PALS) training to neonatal residents (PDSA 2), trained an APP-led hands-on compressions team (PDSA 3), and developed a Neonatal Resuscitation Program (PDSA 4). During the intervention period, protocol and compliance audit were performed to assess impact of the intervention and amount of protocol adherence.

Results A total of 1,228 live births were included in the study (667 baseline, 374 intervention and 187 post-intervention periods). There were no demographic differences between time periods. There was a significant decrease in MBP and CO, and a compensatory increase in SVR during bradycardic episodes (table 1). There was no significant difference in SaO2, CrSO2 or RrSO2.

Conclusions Our preliminary data show that as a result of interventions, significant decrease was achieved in the percentage of infants with very low 5-minute Apgar scores.
hour Advanced Practice Providers (APP) coverage and education campaign for providers to secure the airway prior to starting compressions (PDSA 2), initiated APP-led hands-on scenarios for rotating NICU residents (PDSA 3), and developed ‘Go Bags’ to be taken to deliveries to ensure adequate supplies. A statistical process control (SPC) p-chart was used to evaluate our primary outcome measure of 5-minute Apgar scores <4.

**Summary of Results** There were no demographic differences across the 3 time periods. The percentage of infants with Apgar scores <4 at 5 minutes decreased from 5.2% in the baseline and intervention period to 2.2% in the post-intervention period as special cause was detected in the SPC chart with 8 points below the center line. The percentage of infants receiving chest compressions prior to intubation decreased from 21% during the baseline period to 3% in the intervention and post-intervention periods (p<0.002), with a relative risk of 7.9 for infants born in the baseline period (95% CI: 1.9, 33.7).

**Conclusions** We were able to significantly decrease the percentage of infants with 5-minute Apgar scores <4 and percentage of infants receiving chest compressions prior to intubation which has been sustained for over 3 years. The addition of APPs as champions with a consistent presence in the NICU involved in all resuscitations and resident education has been critical to improving our care in the delivery room.

**#263 Vasoactive Therapy During Therapeutic Hypothermia Is Linked to Adverse Outcomes at Neonatal Intensive Care Unit Discharge**

1S Perugu*, 1M Morone, 2D Patel, 3J Cleary. Harbor-UCLA Medical Center Department of Pediatrics, Torrance, Ca; 2Children's Hospital of Orange County, Orange, CA; 3University of California Irvine, Irvine, CA.

10.1136/jim-2022-WRMC.260

**Purpose of Study** Evaluate if hemodynamic parameters and vasoactive medication requirements during therapeutic hypothermia predict outcomes such as duration of mechanical ventilator support, clinical parameters, and neurodiagnostics in neonates with moderate-severe Hypoxic Ischemic Encephalopathy (HIE).

**Methods Used** Retrospective cohort study of thirty four neonates with moderate-severe HIE from 2010 to 2013 at a quaternary neonatal intensive care unit (NICU) with neurodevelopmental assessments until 2016. Data points were extracted from electronic medical records and supplemented by chart review. The data set was analyzed to compare differences between two groups, with vasoactives (n=18) and without vasoactives (n=16) during therapeutic hypothermia. Primary outcomes (heart rate, blood pressures and cerebral NIRS) were analyzed at five time points. Secondary outcomes were duration of mechanical ventilation, clinical and biochemical parameters, brain MRI, EEG, and Bayley III scaled and composite scores between 6–24 months.

**Summary of Results** There were no statistically significant differences between heart rate, blood pressure or oxygen delivery as measured by cerebral NIRS [6,12, 24, 48 and 72 hours timepoints after birth] in babies with, and without vasoactive therapies. Neonates with vasoactive requirements had more severe hypoxemia, higher blood lactate, lower albumin and hemoglobin, and require prolonged ventilation (P=0.027). Additionally, they were 30% more likely to have abnormal background EEG with low voltage pattern during cooling (p <0.05). Almost 50% of the patients in the vasoactive group had moderate or severe brain injury on 10–12 day MRI. Despite these important differences at NICU discharge, newborns exposed to vasoactives did not have significantly worse neurodevelopmental delays at six months.

**Conclusions** Vasoactive medication requirement in babies with moderate-severe HIE predicts or perhaps contributes to adverse outcomes at NICU discharge. However, infants treated with vasoactives during cooling did not have significantly worse early neurodevelopmental delays. It is critical to comprehensively analyze if therapies are meeting their intended goal of oxygen delivery. Prognosticating risks for neurocognitive deficits in childhood thus remains as an important question for pediatric research.

**#264 Effects of Maternal Preeclampsia on Cardiac Structure and Function in Very Low-Birthweight Infants**

GA Kohbodi*, R Cayabyab, M Ebrahimi, R Ramanathan, B Siasi. University of Southern California Keck School of Medicine, Los Angeles, CA.

10.1136/jim-2022-WRMC.261

**Purpose of Study** Complications related to maternal preeclampsia (PE) are numerous, such as mortality, growth restriction, prematurity, respiratory distress, and morbidities related to neurodevelopment and cardiovascular system in later life. It has been shown in few small studies that exposure to PE can lead to cardiac remodeling in the fetus and term infants, but, this has not been studied in very low birth weight (VLBW) infants. The purpose of the study is to determine whether exposure to maternal PE is associated with hypertrophic cardiac changes and cardiac dysfunction in VLBW infants.

**Methods Used** We retrospectively reviewed the echocardiographic data on VLBW infants born to mothers with PE (cases) and compared them to infants not exposed to PE (controls) between January 2017 to December 2020. Cases were matched by birth weight (BW) and gestational-age (GA). Exclusion criteria were presence of congenital heart disease and no echocardiograms (ECHO) in first 7 days of life. Data was analyzed with Wilcoxon Sign Rank test and McNemar’s Test where appropriate.

**Summary of Results** Our data from 21 cases and controls showed that infants exposed to PE had higher lactate level at birth, significantly higher systolic blood pressure during the first three days of life and a lower rate of vasopressor use within the first 24 hours of life. On ECHO, infants exposed to PE had significantly lower ejection fraction (EF) and fractional shortening (FS) compared to controls. Patent ductus arteriosus (PDA) peak flows were lower in infants with PE compared to controls though not statistically significant (Table 1).

**Conclusions** Our preliminary data show that as a result of exposure to increased resistance to flow through placenta, VLBW infants born to preeclamptic mothers have higher systemic and pulmonary vascular resistance as evidenced by elevated systolic blood pressure after birth and lower peak flow velocity through PDA. Lower EF and SF may also be due to increase systemic vascular resistance. This information may be helpful for choice of inotropic agent for treatment of infants.
Abstracts

Abstract #264 Table 1  Demographics, clinical outcomes and echocardiographic parameters

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Exposure to Preeclampsia (Cases)</th>
<th>Non-Exposure to Maternal Preeclampsia (Controls)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Demographics</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gestational age (weeks)*</td>
<td>27.9 (26.3–30.9)</td>
<td>26.9 (25.4–29)</td>
<td>0.09</td>
</tr>
<tr>
<td>Birth weight (g)*</td>
<td>990 (665–1,255)</td>
<td>950 (710–1,240)</td>
<td>0.58</td>
</tr>
<tr>
<td>Small for Gestational Age, n (%)</td>
<td>10 (48)</td>
<td>3 (14)</td>
<td>0.04</td>
</tr>
<tr>
<td>Male sex, n (%)</td>
<td>12 (57)</td>
<td>12 (57)</td>
<td>1</td>
</tr>
<tr>
<td>Cesarean section, n (%)</td>
<td>19 (90)</td>
<td>15 (71)</td>
<td>0.29</td>
</tr>
<tr>
<td>Antenatal steroids, n (%)</td>
<td>18 (85.7)</td>
<td>21 (100)</td>
<td>0.25</td>
</tr>
<tr>
<td>Clinical outcomes</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiomegaly on admission Chest X-Ray, n (%)</td>
<td>6 (28.6)</td>
<td>1 (4.8)</td>
<td>0.13</td>
</tr>
<tr>
<td>Lactate Day 0 (mmol/L)*</td>
<td>3.3 (1.8–9.3)</td>
<td>1.8 (1.5–3.3)</td>
<td>0.03</td>
</tr>
<tr>
<td>Lactate Day 1 (mmol/L)*</td>
<td>2.9 (1.7–3.9)</td>
<td>1.7 (1.3–2.3)</td>
<td>0.11</td>
</tr>
<tr>
<td>Lactate Day 2 (mmol/L)*</td>
<td>1.6 (1.4–2.2)</td>
<td>1.65 (1.4–2.4)</td>
<td>0.47</td>
</tr>
<tr>
<td>Systolic blood Pressure Day 0*</td>
<td>49 (41–59)</td>
<td>42 (33–50)</td>
<td>0.04</td>
</tr>
<tr>
<td>Systolic blood Pressure Day 1*</td>
<td>45 (39–51)</td>
<td>41 (31–43)</td>
<td>0.03</td>
</tr>
<tr>
<td>Systolic blood Pressure Day 2*</td>
<td>49 (44–54)</td>
<td>43 (39–46)</td>
<td>0.02</td>
</tr>
<tr>
<td>Echocardiographic Data</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Fractional Shortening (%)</td>
<td>35.7 (31.8–41.9)</td>
<td>40.6 (36.8–44.6)</td>
<td>0.03</td>
</tr>
<tr>
<td>Ejection Fraction (%)*</td>
<td>69.1 (64.3–77.2)</td>
<td>76.1 (71.1–79.6)</td>
<td>0.04</td>
</tr>
<tr>
<td>Patent ductus arteriosus size (mm)*</td>
<td>2.2 (1.7–2.8)</td>
<td>1.57 (1.4–2.3)</td>
<td>0.21</td>
</tr>
<tr>
<td>Patent ductus arteriosus</td>
<td>1.3 (1–2)</td>
<td>1.95 (1.3–2.4)</td>
<td>0.17</td>
</tr>
</tbody>
</table>

*median (interquartile range).

Purpose of Study Bronchopulmonary dysplasia (BPD) primarily affects extremely preterm infants and has been associated with high mortality and morbidity. An online estimator for BPD severity was developed by NICHD that may guide interventions. This study aims to investigate the optimal use of the estimator for BPD prediction.

Abstract #265 Table 1

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Exploratory dataset for grid search</th>
<th>Postnatal day 1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensitivity</td>
<td>0.764</td>
<td>0.740</td>
</tr>
<tr>
<td>Specificity</td>
<td>0.741</td>
<td>0.750</td>
</tr>
<tr>
<td>Positive Predictive Value</td>
<td>0.750</td>
<td>0.720</td>
</tr>
<tr>
<td>Negative Predictive Value</td>
<td>0.755</td>
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<tr>
<td>Value</td>
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<td>0.759</td>
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<tr>
<td>Validation dataset</td>
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<td></td>
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<tr>
<td>Sensitivity</td>
<td>0.805</td>
<td>0.825</td>
</tr>
<tr>
<td>Specificity</td>
<td>0.641</td>
<td>0.675</td>
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<tr>
<td>Positive Predictive Value</td>
<td>0.702</td>
<td>0.717</td>
</tr>
<tr>
<td>Value</td>
<td>0.758</td>
<td>0.794</td>
</tr>
<tr>
<td>Negative Predictive Value</td>
<td>0.794</td>
<td>0.821</td>
</tr>
<tr>
<td>Value</td>
<td>0.848</td>
<td>0.848</td>
</tr>
<tr>
<td>Accuracy</td>
<td>0.725</td>
<td>0.750</td>
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</tbody>
</table>

Performance of the NICHD Neonatal BPD Outcome Estimator using combined scores from the severe and death categories with a cutoff number of 21, above which predictive of a severe (positive) disease outcome and below which predictive or a non-severe (negative) disease outcome.

Abstract #265 Figure 1

Abstract #265

COMBINING PROBABILITY SCORES TO OPTIMIZE CLINICAL USE OF THE ONLINE NEONATAL BRONCHOPULMONARY DYSPLASIA OUTCOME ESTIMATOR

Purpose of Study Bronchopulmonary dysplasia (BPD) primarily affects extremely preterm infants and has been associated with high mortality and morbidity. An online estimator for BPD severity was developed by NICHD that may guide with hemodynamic compromise born to mothers with PE. Data collection is ongoing to have a larger sample size to confirm these findings.

Abstract #266

Gestational Long Term Hypoxia and Phenotypic Transformation of Fetal Sheep Pulmonary Arteries

Purpose of Study Pulmonary hypertension is a serious medical condition that affects the heart and blood vessels of the lungs, which can lead to heart failure and other sequelae. Gestational long-term hypoxia predisposes severity category. A receiver operating characteristic curve and provides severity probability scores for five categories: no BPD, mild, moderate, severe, or death. Generalized additive model-

Abstract #303

Heart Failure and Other Pulmonary Hypertension in Pregnant Women

Abstract #304

Transcriptomic and Metabolomic Assessment of Pulmonary Hypertension in High Altitude Pregnant Women

Abstract #305

Mfsd2a Sex-Divergent Decreases in Serum and Lung Phosphatidylcholine (PC) and Phosphatidylcholine-DHA in Association with Pulmonary Hypertension in Pregnant Women.
Abstracts

Methods Used This retrospective single-center study included infants of 23–30 weeks gestation weighing ≤ 1,250 grams. Relevant variables were entered into the estimator which provides severity probability scores for five categories: no BPD, mild, moderate, severe, or death. Generalized additive modeling was used to develop probability score trajectories for each severity category. A receiver operating characteristic curve and the Youden’s J statistics, followed by a narrowed grid search, were used to assess predictability of severe BPD or death.

Summary of Results A total of 469 infants were included in the study. Direct use of the highest probability score for BPD prediction yielded 29% accuracy due to close proximity of probability scores between categories as assessed by probability score trajectories (figure 1A). Using combined severe and death scores as well as combined no BPD and mild BPD scores resulted in improved trajectory separations, which may convey improved predictability (figure 1B). After a grid search, we found a combined severe and death probability score with a cutoff value of 21% resulted in an optimal overall accuracy rate of 75% in predicting a composite outcome of severe BPD or death (table 1 and figure 1C).

Conclusions Combining probability scores of different categories may improve BPD outcome prediction.

Neonatology – perinatal biology I

Concurrent session

8:00 AM

Friday, January 21, 2022

GESTATIONAL LONG TERM HYPOXIA AND PHENOTYPIC TRANSFORMATION OF FETAL SHEEP PULMONARY ARTERIES

M Lee, C Leslie, C Guoqhe, L Zhang, M La Frano, S Wilson, Loma Linda University School of Medicine, Loma Linda, CA; University of New Mexico Health Sciences Center, Albuquerque, NM; California Polytechnic State University, San Luis Obispo, CA

Purpose of Study Pulmonary hypertension is a serious medical condition characterized by abnormally high blood pressures in the pulmonary circulation, which can ultimately lead to right-heart failure and other sequelae. Gestational long-term hypoxia (gLTH) is a known risk factor in the development of pulmonary arterial hypertension of the newborn. The pathophysiology of this process is not fully understood, though various processes that cause structural remodeling include enhanced cell growth and proliferation. The purpose of the current study was to test the hypothesis that gLTH would increase markers of cell growth and proliferation and decrease markers of smooth muscle differentiation.

Methods Used Pregnant sheep were divided into low- and high-altitude groups. Low-altitude sheep were transported to LLU (355 m) for study while high-altitude sheep were transported to the Barcroft Research Station at the University of California White Mountain Research Center (3801 m) and housed there for the latter ~110 days of gestation to induce gLTH, before being transported to LLU just prior to delivery. Fetal sheep were delivered by c-section near-term at 130–140 days of gestation at LLU. Fetal pulmonary arterial segments were isolated and frozen in LN2 and stored at -80C for subsequent analysis of metabolites as well as proteins using metabolomics and proteomics approaches along with miRNA in plasma exosomes through next-gen sequencing. Based on the results from the proteomics analysis, immunohistochemical analysis was performed as well.

Summary of Results Following gLTH there were increases in metabolic and proteomic markers of cell proliferation and phenotypic transformation that were associated with changes in exosomal miRNA. There was a decreased expression of a number of structural proteins including myosin heavy chain 11, several collagen isoforms, along with S100 proteins, histones, Caveolin 1, and markers of cellular inflammation. Metabolomic analyses indicate there were increases in oxidative stress, the pentose phosphate shunt, and arachidonic acid metabolism. There were select changes in exosomal miRNA, including upregulation of let-7a-5p, mir-221, mir-103, mir-21 and downregulation of let-7.

Conclusions Our data provide new evidence across multiple platforms regarding the mechanistic underpinnings associated with pulmonary arterial hypertension in newborns due to gLTH. Overall, the results indicate that following gLTH ovine fetal pulmonary arteries are taking on a synthetic phenotype, which likely promotes medial wall thickening along with increases in tissue inflammation and oxidative stress. We are in the process of examining the phenotypic transformation in more depth and working to elucidate the molecular underpinnings to the disease process.

IUGR DECREASES PHOSPHATIDYLCHOLINE-DHA AND LIPOPHOSPHATIDYLCHOLINE-DHA IN ASSOCIATION WITH INCREASED MFSD2A IN THE RAT LUNG


Purpose of Study Human neonates rely on adequate uptake of docosahexaenoic acid (DHA) for appropriate lung development. Low circulating neonatal DHA is associated with the development of bronchopulmonary dysplasia and is exacerbated by intrauterine growth restriction (IUGR). Lung uptake of DHA from the neonatal circulation is primarily in the form of phosphatidylcholine (PC) and lysophosphatidylcholine (LPC) DHA. Cellular uptake of PC-DHA and LPC-DHA is maximized by the omega-3 fatty acid transporter Mfsd2a, which in other tissue types increases expression under conditions of low DHA. We previously demonstrated, in a rat model, that IUGR decreases total circulating DHA in male but not female rat pups. However, the effect of IUGR on serum and lung PC-DHA and LPC-DHA, as well as on lung Mfsd2a mRNA levels is currently unknown. We hypothesize that IUGR causes sex-divergent decreases in serum and lung PC-DHA and LPC-DHA in association increased Mfsd2a in the rat.

Methods Used IUGR was induced via bilateral uterine artery ligation in pregnant Sprague Dawley rats. Control dams received anesthesia only. Rat pups were euthanized at term c-section delivery, and serum and lung tissue collected. PC-DHA and LPC-DHA were measured using GC/MS. Mfsd2a mRNA was measured using real-time RT PCR. Two-way ANOVA was used to assess interaction effects, t-test was used to assess group differences.
Summary of Results Results are IUGR as% of control±SD, *P<.05. A significant IUGR-sex interaction effect occurred for serum PC-DHA whereby IUGR decreased serum PC-DHA in male rat pups (64±7%*), with no change in female rat pups. No other interaction effects were observed. IUGR decreased serum LPC-DHA (65±30%*). Within the lung, IUGR decreased PC-DHA and LPC DHA (85±8%* and 79±23%* respectively). IUGR also increased lung Mfsd2a mRNA (159±76%).

Conclusions IUGR decreases lung PC-DHA and LPC-DHA in rat pups in association with increased Mfsd2a mRNA expression of both sexes. In contrast, the IUGR-induced decreased serum PC-DHA is restricted to male rat pups. We speculate that upregulation of lung Mfsd2a may be a compensatory mechanism to maximize lung DHA uptake in IUGR.

Purpose of Study Pregnancy complications such as gestational diabetes and maternal obesity, which can result in elevated insulin-like growth factor-1 (IGF-1) concentrations and fetal overgrowth, are associated with abnormal pancreatic islet development and beta cell failure later in life. We have demonstrated that a 1-week IGF-1 LR3 infusion into fetal sheep results in reduced in vivo and in vitro insulin secretion. However, acute exposure to IGF-1 enhances insulin secretion in adult rodent islets. Therefore, we hypothesized that a 90-minute IGF-1 LR3 infusion into fetal sheep would potentiate in vivo and in vitro glucose-stimulated insulin secretion (GSIS).

Methods Used Late gestation fetal sheep (n=10) were infused for 90 minutes at 6.6 μg/kg/hr with either IGF-1 LR3 (IGF-1) or vehicle control (CON). Each animal received both infusions spaced 2–3 days apart. Fetal arterial blood gases and plasma insulin, glucose, and lactate were measured. At minute 90 of infusion, and with infusates still running, GSIS was measured using a hyperglycemic clamp with frequent glucose and insulin measurements. At the end of the second infusion (IGF-1, n=5; CON, n=5), fetal islets were isolated and incubated with 1.1, 2.7, or 11 mM/L glucose or 30 mM/L KCl and then pelleted. Insulin concentrations of the media and cell pellet were measured by ELISA. Insulin secretion was calculated as the fraction of total islet insulin secreted into the media.

Summary of Results Plasma insulin concentrations decreased during IGF-1 infusion and were 48% lower at the end of infusion compared to CON (*P<0.05). Fetal arterial blood gases and plasma glucose and lactate were not different between IGF-1 and CON at the end of infusion. During the GSIS study, glucose concentrations were similar, but insulin concentrations were 66% lower with IGF-1 infusion compared to CON (*P<0.0001). Insulin secretion in isolated fetal islets was not different based on infusion just prior to necropsy.

Conclusions IGF-1 LR3 infusion for 90 minutes into late gestation fetal sheep lowers plasma insulin concentrations and attenuates fetal GSIS. However, reduced insulin secretion does not persist in isolated fetal islets exposed to IGF-1 in vitro at the time of necropsy. Therefore, while acute increases in fetal IGF-1 may directly suppress insulin secretion, the fetal beta cell in vitro retains the ability to recover GSIS. With more prolonged elevations in fetal IGF-1, beta cells become programmed to secrete less insulin in response to glucose. We speculate that chronically, but not acutely, elevated IGF-1 concentrations during the fetal period in pregnancies complicated by gestational diabetes or maternal obesity may underlie the risk of beta cell failure and diabetes later in life.
Abstracts

#270  Elevated triglyceride concentrations in umbilical cord plasma from human intrauterine growth restricted pregnancies correlate with ultrasound markers of poor fetal wellbeing

SS Chassen*, K Zemski-Berry, S Raymond-Whish, C Driver-Rigdon, J Hobbins, T Powell. University of Colorado, Aurora, CO

10.1136/jim-2022-WRMC.267

Purpose of Study Intrauterine growth restriction (IUGR) is associated with decreased subcutaneous fat depots in the fetus and health consequences including neurodevelopmental impairment for the child. Intrauterine brain development depends on long chain polyunsaturated fatty acids (LCPUFA), but their metabolism in IUGR pregnancies is poorly understood. Circulating fetal LCPUFA levels and their relation to ultrasound (US) markers of fetal growth and wellbeing (estimated fetal weight [EFW], umbilical artery [UA] and middle cerebral artery [MCA] Doppler) in human pregnancies have also not been elucidated. We hypothesized that LCPUFA concentrations are decreased in IUGR cord plasma and correlate with US markers of severe IUGR.

Methods Used Pregnant women with an EFW of <10th% for gestation in 2nd or 3rd trimester were included. Prenatal US was obtained every 1–4 weeks from enrollment, with UA and MCA pulsatility index (PI) assessed by Doppler. Birth weight (BW) was used to classify the infants as pathologically small (BW <3rd%, IUGR), small for gestational age (BW 5–10th%, SGA), or appropriate for gestational age (BW >10th%, AGA). Cord plasma samples (umbilical vein) were collected at delivery and targeted lipidomic analyses performed by liquid chromatography mass spectrometry (LCMS) following extraction of fatty acids in triglycerides (TG), phosphatidylcholine (PC), and lysophosphatidylcholine (LPC) lipid classes. Statistical differences and correlations were assessed using student’s t test and Pearson’s coefficient.

Summary of Results Birth weights were 25% smaller (p=0.004) and UA PI 122% higher (p=0.02) in the IUGR group (n=8) compared to combined AGA/SGA group (n=12). Concentrations of all LCPUFA in the TG fraction were greater in IUGR cord plasma compared to combined AGA/SGA group (p<0.05), and negatively correlated with EFW (avg r = -0.577, p<0.05), BW (avg r=-0.592, p=0.01), and MCA PI (r = -0.469, p<0.05). LCPUFA concentrations in the PC fraction were similar between groups, though LPC-docosahexaenoic acid (DHA) was decreased in IUGR (p<0.01).

Conclusions We found increased TG-LCPUFA in IUGR cord plasma that correlated with US markers of compromised fetal growth. LPC-DHA (vital for brain development) is believed to be the lipid form for transport across the blood brain barrier, but our data show that LCPUFA packaging into TG (a storage form) may occur at the expense of LPC formation. We speculate that the increased stored TG-LCPUFA represents the fetal response to an adverse in utero environment, such as oxidative stress, and may have detrimental consequences for ultimate LCPUFA transport to the brain for development.

#271  Effects of chronic high altitude hypoxia during gestation on mitochondrial oxygen consumption rates in pulmonary arteries from fetal and newborn sheep

SF Hanson*, ST Murray, S Wilson, L Zhang, A Blood. Loma Linda University School of Medicine, Loma Linda, CA

10.1136/jim-2022-WRMC.268

Purpose of Study Chronic hypoxia during gestation has been associated with neonatal pulmonary hypertension in newborns of both humans and sheep. MicroRNA-210 levels have been found to be upregulated in response to hypoxia via HIF-1a in nearly every mammalian tissue studied. This, together with evidence that miR210 orchestrates many cellular responses to hypoxia, has led to use of the term ‘universal hypoxamir.’ One effect of miR210 is downregulation of the expression of iron sulfur cluster assembly proteins that are critical for normal mitochondrial function. We hypothesized that chronic hypoxia during gestation results in decreased mitochondrial oxygen consumption capacity in pulmonary arteries of near-term fetal and 2-week-old newborn lambs.

Methods Used Pregnant ewes were quartered at either low altitude (330m) or high altitude (3801m) for the last 100 days of gestation (term = 150 days). Lungs were harvested from fetal lambs following c-section within 12 days before term gestation, and from newborn lambs at 10 to 14 days after birth. Pulmonary arteries were isolated from the 4th to 6th branching generation, segments of these arteries were studied with a Seahorse XFe 24 analyzer to evaluate oxygen consumption rate (OCR) associated with basal, ATP-linked, maximal, proton leak, and non-mitochondrial respiration. OCR was normalized to vessel weight, and comparisons between normoxic/hypoxic and fetal/newborn groups were made using 2-way ANOVA with post hoc analysis (significance at p<0.05).

Summary of Results Compared to sea level controls, chronic hypoxia resulted in a significant decrease in basal oxygen consumption in both fetal (from 241.23 ± 32.89 to 176.74 ± 31.91 ml/mg protein) and newborn (from 281.74 ± 26.56 to 168.03 ± 43.25 ml/mg protein) pulmonary arteries. However, contrary to our hypothesis, maximal oxygen consumption rates were not affected by chronic hypoxia. Likewise, hypoxia had no effect on ATP-linked or non-mitochondrial OCR. Unexpectedly, hypoxia resulted in a decrease in oxygen consumption due to mitochondrial proton leak in both fetal (from 53.69 ± 9.34 to 13.76 ± 6.37 ml/mg protein) and newborn (from 71.25 ± 12.16 to 37.62 ± 4.98 ml/mg protein) hypoxic arteries. The magnitude of decrease in OCR due to proton leak nearly accounted for the overall decrease in baseline OCR, suggesting that the decrease in baseline OCR is due largely to decreased proton leak.

Conclusions These differences in proton leak when exposed to hypoxic conditions might be explained by downstream effects in the regulation of mitochondrial respiration involving uncoupling proteins and ROS signaling in complex with fatty acid oxidative metabolism and a glycolytic shift.
Surgery III
Concurrent session
8:00 AM
Friday, January 21, 2022

#272 URINE SODIUM TO URINE CREATININE RATIO AS A MARKER OF TOTAL BODY SODIUM IN INFANTS WITH INTESTINAL FAILURE

Choi S1, L Casey1, Albersheim1, Van Oerle K4, Irvine A2, Piper HG1. The University of British Columbia Faculty of Medicine, Vancouver, BC, Canada; BC Children’s Hospital, Vancouver, BC, Canada; The University of British Columbia Women’s Hospital and Health Centre, Vancouver, BC, Canada; BC Centre for Disease Control, Vancouver, BC, Canada; Simon Fraser University, Burnaby, BC, Canada

Purpose of Study Urine sodium (UNa) is a measure of total body sodium in infants with intestinal failure (IF) but can be misleading as it does not reflect volume status. Urine sodium to urine creatinine ratio (UNa:UCr) may offer a more accurate measure, but is not routinely used. This study compares UNa:UCr to UNa as a marker of sodium status in infants with IF.

Methods Used A retrospective review of infants with IF, from a single centre, from 2018–2020 was conducted (REB H20–00816). IF etiology, intestinal anatomy, nutritional intake, urine electrolytes and anthropometrics were collected. Linear mixed effects models adjusting for repeated measures were used to associate UNa and UNa:UCr with weight gain and sodium intake.

Summary of Results Twenty-two infants with a median gestational age of 31 weeks were included. IF etiology included gastrostomy (41%), necrotizing enterocolitis (23%), and intestinal perforation (14%). Infants had an average of 3 paired UNa and UNa:UCr measures for a total of 74 paired measurements. UNa:UCr more strongly correlated with sodium intake compared to UNa (R=0.25, p=0.032 vs. R=0.10, p=0.38). Overall, neither UNa (p=0.21) nor UNa:UCr (p=0.16) were significantly correlated with weight gain. However, for infants receiving ≤50% nutrition enterally, weight gain correlated with UNa (p=0.01) and UNa:UCr (p=0.01). UNa:UCr >35 predicted adequate growth regardless of enteral intake (92% sensitivity, 59% specificity).

Conclusions UNa:UCr is a measure of total body sodium that correlates with sodium intake in infants with IF. Our study indicates UNa:UCr >35 is associated with adequate growth and can be used to guide further validation studies.

#273 IDENTIFYING NONADHERENCE IN LIVER TRANSPLANT CANDIDATES USING SELF-REPORTED QUALITY OF LIFE

Nichols J1, Vutien Z2, Biggins S2, McCandlish K2, Bambaia J3, Rees J3, Perkins D3. University of Washington School of Medicine, Seattle, WA; University of Washington School of Social Work, Seattle, WA; University of Washington School of Social Work, Seattle, WA; University of Washington Medical Center, Seattle, WA; Seattle Children’s Hospital, Seattle, WA

Purpose of Study Nonadherence remains an intractable barrier to undergoing liver transplantation (LT) for some transplant candidates and can result in candidate delisting and poor survival. Currently, there is no objective tool to evaluate pre-transplant nonadherence risk. We investigated if the Liver Disease Health-Related Quality of Life Short Form (SF-LDQOL) could help proactively identify LT candidates at risk of nonadherence.

Methods Used We conducted a retrospective cohort study of 720 adults (≥18 years old) for LT at the University of Washington Medical Center from 9/1/2012 to 8/30/2017, including those who completed the SF-LDQOL prior to listing. Univariable and multivariable competing risk analysis was performed to estimate the risk of delisting due to nonadherence. Results were statistically significant if P < 0.05.

Summary of Results 358 (49.7%) LT candidates responded to the SF-LDQOL prior to being listed. Ultimately, 23 (6.4%) were delisted due to nonadherence, 205 (57.3%) underwent transplantation, 79 (22.1%) died on the wait list or became too sick for transplantation, 19 (5.3%) were delisted due to cancer progression beyond the Milan Criteria, 19 (5.3%) were delisted for clinical improvement, and 13 (3.6%) remained actively listed at the end of follow up. In the multivariable competing risk analysis, SF-LDQOL responses indicating ‘poor memory’ (SHR: 3.48; 95% CI: 1.44–8.42; P=0.006) and ‘poor future outlook’ (SHR: 2.96; 95% CI: 1.01–8.62; P=0.047) and being listed for repeat LT (SHR: 16.80; 95% CI: 1.76–160.30; P=0.01) were associated with a higher risk of delisting due to nonadherence. Female sex (SHR: 0.30; 95% CI: 0.10–0.94; P=0.04) and having a history of previous abdominal surgery (SHR: 0.28; 95% CI: 0.10–0.80; P=0.02) were associated with a lower risk of delisting due to nonadherence.

Conclusions Responses in the SF-LDQOL indicating ‘poor memory’ and ‘poor future outlook’ may suggest an increased risk of nonadherence in LT candidates. Tools that assess for these specific measures may help identify those candidates in need of additional support to avoid delisting.

#274 LAMINECTOMY OR FUSION FOR INTRADURAL EXTREMEDULLARY TUMORS?

Mo K1, Al Farii H2, Lee S3. Case Western Reserve University School of Medicine, Cleveland, OH; Western University School of Medicine, Pomona, CA; Johns Hopkins University, Baltimore, MD

Purpose of Study Laminectomy and laminectomy with fusion have both been demonstrated as surgical techniques that can treat intradural extramedullary tumors (IDEM). However, we are not aware of any studies comparing outcomes between laminectomy versus laminectomy with fusion. The purpose of this study was to compare the rate of 30-day complications following laminectomy with or without fusion for IDEM.

Methods Used Adult patients undergoing laminectomy for IDEM from 2012 to 2018 were identified in the National Surgical Quality Improvement Program database. Patients undergoing laminectomy for IDEM were stratified into 2 cohorts: those who received fusion and those who did not. In this analysis, pre-operative patient characteristics and demographic variables such as Age, BMI, Gender, ASA Class, Smoking, Functional Status, Diabetes, Chronic Obstructive Pulmonary Disease, Congestive Heart Failure, Hypertension, Renal Failure, Dialysis, Weight Loss, Steroid Use, Bleeding Disorder, Dyspnea at Rest, Dyspnea at Moderate Exertion, and...
Abstracts

#274 Table 1  Fusion with laminectomy by location of intradural extramedullary tumor

<table>
<thead>
<tr>
<th></th>
<th>Odds Ratio</th>
<th>P-Value</th>
<th>Lower CI</th>
<th>Upper CI</th>
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</thead>
<tbody>
<tr>
<td>Length of Stay &gt; 5 days</td>
<td>2.73</td>
<td>&lt;0.001</td>
<td>1.95</td>
<td>3.820</td>
</tr>
<tr>
<td>Transfusion</td>
<td>3.15</td>
<td>&lt;0.011</td>
<td>1.78</td>
<td>5.57</td>
</tr>
<tr>
<td>Myocardial Infarction</td>
<td>5.62</td>
<td>0.241</td>
<td>0.31</td>
<td>101.00</td>
</tr>
</tbody>
</table>

General Anesthesia were assessed. 30-day wound, sepsis, cardiac, pulmonary, renal, and thromboembolic complications, as well as mortality, postoperative transfusions, extended length of stay, and reoperation were assessed. Bivariate analyses, including chi-squared and t-test, and multivariable logistical regression were performed.

Summary of Results Of 2,027 total patients undergoing laminectomy for IDEM, 181 (10%) also had fusion. There were 72/373 (24%) laminectomies with fusion in the cervical region, 67/801 (8%) laminectomies with fusion in the thoracic region, and 42/776 (6%) laminectomies with fusion in the lumbar region. Following adjustment, patients who received laminectomy with fusion were more likely to have increased length of stay (OR 2.73, P<0.001) and increased rate of postoperative transfusion (OR 3.15, P<0.001). Patients undergoing laminectomy in the cervical spine for IDEM had a tendency to receive additional fusion (P<0.001). Laminectomy with fusion for IDEM is associated with higher anesthesia class (P=0.046), hypertension (P=0.035), dialysis (P=0.004), and steroid use (P=0.03).

Conclusions Increased length of stay and rate of post-operative transfusion were associated with laminectomy with fusion for IDEM. Patients undergoing laminectomy in the cervical spine for IDEM had a tendency to receive additional fusion. Laminectomy with fusion was also associated increased anesthesia class, hypertension, dialysis, and steroid use.

#275 TIME EFFICIENCY AND PERFORMANCE OF DISPOSABLE VERSUS REUSABLE CYSTOSCOPY: A PROSPECTIVE, RANDOMIZED BENCHTOP COMPARISON

1C Baz*, 2R Chen, 1DR Perervi, 1IC Hartman, 2A Amanapali, 2OD Belle, 2E Baldwin, 2D Baldwin. 1Loma Linda University School of Medicine, Loma Linda, CA; 2Loma Linda University Medical Center, Loma Linda, CA

Purpose of Study While reusable cystoscopes are commonly used for urologic procedures, bulky equipment and lengthy sterilization times can impact time efficiency. Disposable cystoscopes may offer advantages including increased availability, greater portability, and faster setup times. The aim of this study is to compare procedure time, cystoscope specifications, and physician satisfaction between disposable and reusable cystoscopes.

Methods Used Ten urologists (5 attendings and 5 residents) performed timed, simulated bedside cystoscopies with target identification using a prospective, randomized, crossover study design. Each subject used both a new disposable Ambu aScope 4 Cysto and a reusable Olympus CYF-5 flexible cystoscope. Afterward, participants completed a satisfaction survey. Time required for supply-gathering, setup, cystoscopy, and cleanup were compared. Image definition, field of view, deflection angle, force required for deflection, irrigation rate, weight, and working length were also compared.

Summary of Results The disposable cystoscope required less time for supply-gathering (187.5 vs. 289.4s, p<0.05), setup (203.3 vs. 327.5s, p<0.01), and cleanup (183.7 vs 356.2s, p<0.05) compared to the reusable, while cystoscopy times were similar (230.4 vs. 274.1s; p=.575).

Optical testing showed higher image definition for the disposable cystoscope (6.30 vs 2.00 line pairs/mm, p<0.001), but a smaller field of view (66.54° vs 107.91°, p<0.001) and a lack of user-adjustable optical settings. The disposable cystoscope also had increased deflection (214° up/182° down vs 198° up/109° down, p<0.001), yet required more force to deflect 180° up (6.86 vs 4.46N, p<0.001) and 90° down (4.66 vs 3.55 N, p<.01). The reusable cystoscope had a greater mass (325.11 vs 159.03 g, p<.001), shorter working scope length (37.45 cm vs 38.97 cm, p<.001), and a faster irrigation rate at 200 cm H2O (494.10 vs 387.40 mL/min, p<.001). Post-testing, deflection was reduced for two of the ten disposable cystoscopes.

Survey results showed higher ratings in time-efficiency (9.5 vs 6.2/10, p=.000) and overall satisfaction (9.3 vs 7.9/10, p=.004) for the disposable cystoscope. There was no difference in ratings for maneuverability, image quality, confidence in sterility, or cystoscopy time (p>.05 for all).

Conclusions While the disposable cystoscope had better image quality, greater deflection, and was faster to assemble and disassemble, the reusable cystoscope had greater durability, more optical settings, wider field of view, faster irrigation rates, and required less force for deflection. Knowledge of the strengths and weaknesses of each device could assist surgeons in optimizing cystoscope utilization in specific clinical scenarios. Further studies are warranted and should include relative cost-effectiveness and infection risks.

#276 SELF-DETERMINATION OF APTITUDE IN SURGICAL SPECIALITIES

M Dea*. Rocky Vista University College of Osteopathic Medicine, Parker, CO

Purpose of Study The field of surgery remains a highly coveted and competitive specialty. There are currently no standardized assessments used to assess a student’s psychomotor skills, innate aptitude and other technical surgical skills. Laparoscopic surgery involves specific skills such as a strong appreciation of depth perception, manual/bi-manual dexterity, reduced tactile feedback all with operating with increased hand tremor. This study investigated two main objectives. Firstly, to evaluate the relationship between motor skills, such as drawing, playing videogames or musical instruments and surgical simulator performance. The second objective was to evaluate how a student’s self-perception of their motor skills before and after a simulation assessment related to their actual performance.

Methods Used A cohort of unconditioned medical students defined as having no prior experience with surgical simulators completed a pre-task survey, which included demographic information, previous exposure to playing videogames, painting/drawing, playing a musical instrument, organized sports and knitting/sewing. The students also provided information on their medical specialty of interest and their own predictive
LEADERSHIP AND COMMUNICATION IMPROVEMENTS IN ORTHOPEDIC SURGERY OPERATING ROOM TIME-OUT HUDDLES

Purpose of Study: Time-out huddles, in which all information is confirmed, have significantly reduced adverse outcomes, near misses, and never events. During this critical portion of the procedure, the group leader takes charge and reviews necessary information with the team. The procedure does not continue until each person agrees on all information. This creates an interesting dynamic where everyone shares responsibility for a high-stakes event though most communication and leadership are in the hands of a sole spokesperson. We aim to uncover whether or not members of the team feel valued in the decision-making process as well as suggest means of improvement.

Methods Used: We sent out surveys across the United States to active members operating inside the orthopedic surgery operating room and asked them to verify their position. We then asked them to express how valued they feel in terms of the time-out huddles on a scale from 1 to 10 with 1 being not valued at all and 10 being extremely valued. 30 orthopedic surgeons, 41 anesthesiologists, 66 nurses, 59 medical students, and 79 technicians were interviewed. The response rate remained at 78.57%.

Summary of Results: Each team member’s average response in alphabetic order is presented. Anesthesiologists reported an average response of 8.44. Medical students reported an average response of 5.56. Nurses reported an average response of 8.52. Orthopedic surgeons reported an average response of 9.87. Technicians reported an average response of 8.80.

Conclusions: Our study shows significant differences in the reported level of feeling of value in the time-out huddles in orthopedic surgery operating rooms. There are three significantly different groups at a confidence level of 99%. The high level among orthopedic surgeons can be explained by the fact that these are typically the ones conducting the discussions and therefore they have a lot of control. The middle group includes anesthesiologists, nurses, and technicians. The group with the lowest level was medical students. This is concerning because keeping future health care professionals engaged and responsible is crucial in part of a sustainable future. Many reported that they were passive observers rather than participants. They also reported fear of poor evaluations if they spoke out in any way. It remains true that this is the only group that could be graded.
Conclusions The skull shape differences seen between the two surgical cohorts right after surgery were expected, given the immediate expansion that occurs with PVR. By 2-year follow-up, both surgical cohorts had morphometric results that were comparable, and both cohorts’ cranial vaults had normalized to match control CT scans. Our chart review supports differences in anesthesia exposure, blood loss, and follow-up duration. When choosing a surgical approach for SS, a holistic evaluation that considers other factors beyond the morphometric outcome may be beneficial.

Behavior and development I
Concurrent session
10:15 AM
Friday, January 21, 2022

BEHAVIORAL OUTCOMES IN INDIVIDUALS WITH FRAGILE X SYNDROME AND VARYING AUTISM SEVERITY

Y Tak, H Biag, M Salcedo-Arellano, R Ashworiya, K Kim, D Hess, RJ Hagemeier. University of California Davis MIND Institute, Sacramento, CA; University of California Davis School of Medicine, Sacramento, CA

Purpose of Study Fragile X Syndrome (FXS) individuals have developmental delay, learning disabilities and social/behavioral deficits. Approximately 60% of males and 20% of females with FXS have Autism Spectrum Disorder (ASD). Individuals with both FXS and ASD show more severe behavioral, cognitive and language problems than individuals without ASD; however, the behavioral profile varies depending on ASD severity. We aim to evaluate the behavioral profile of individuals with FXS and varying degrees of Autism.

Methods Used Outcome measure data was collected from baseline behavioral questionnaires/assessments of patients (n=41, ages 6 to 25) enrolled in the UC Davis MIND Metformin Clinical Trial. Outcomes included five different self-reported questionnaires [Pediatric Quality of Life (PedsQL), Anxiety, Depression and Mood Screening (ADAMS), Aberrant Behavior Checklist (ABC), Child Sleep Habits Questionnaire (CSHQ), Swanson, Nolan and Pelham-IV (SNAP-IV)] and 2 standardized assessments (Leiter and Vineland). Autism severity was measured by the Autism Diagnostic Observation Scale (ADOS-2), which indicated ASD severity: no/mild ASD (n=8), moderate ASD (n=15), severe ASD (n=18). One-way ANOVA compared mean differences in behavioral outcomes among three ASD severity groups followed by Tukey’s post-hoc tests. Significance of differences was determined at p-value < 0.05.

Summary of Results The Leiter (non-verbal IQ) score of the no/mild ASD group significantly differed from the moderate ASD (p=0.0348) and severe ASD (p=0.0045) groups. The Vineland adaptive behavior composite score was significantly different between the no/mild ASD and severe ASD groups (p=0.0109) (Table 1). Of the five different self-reported questionnaires, one sub-score from the Peds Quality of Life showed a significant difference between the moderate ASD and severe ASD groups (School Functioning, p=0.005). The remaining PedsQL scores (Physical Functioning, Emotional Functioning and Social Functioning), Child Sleep Health total, SNAP-IV scores, Aberrant Behavior Checklist total, Anxiety, Depression and Mood Screening scores did not show significant differences between the groups.

Conclusions Our results identified significant differences between individuals with varying autism severity in non-verbal IQ and adaptive behavior. There were no significant differences in other parent-reported behavioral measures and overall pediatric quality of life except for that related to school functioning. Increased sample size may lead to emergence of further significant differences in the other behavioral measures. Nonetheless, facilitating further school accommodations/inclusion for children with ASD can help to improve their school-related quality of life.

INVESTIGATING RELATIONS BETWEEN MATERNAL MOOD DISORDERS AND RATES OF DEVELOPMENTAL CARE PRACTICES IN THE NEONATAL INTENSIVE CARE UNIT

M Chan Morales, E Brignori-Pérez, VA Marchman, E Armer, R Shaw, KE Travis, M Scala. Stanford University, Stanford, CA

Purpose of Study Developmental care practices in the NICU, such as positive touch and skin-to-skin holding, have led to substantial improvements in clinical and neurodevelopmental outcomes in children born preterm, and represent the largest source of NICU parenting behaviors. Parents of children born preterm are at risk for experiencing mood disorders, such as depression, anxiety, and post-traumatic stress disorder (PTSD). Mood disorders have been shown to have negative impacts on parenting behaviors for healthy term infants. Developmental care activities have been used at other institutions as part of treatment interventions for NICU parents with mood disorders. However, few studies have explored relations between perinatal parental mood disorders and rates of participation in developmental care activities. The purpose of this study is to examine the relationship between maternal mood disorders and rates of mother-delivered developmental care activities.

Methods Used Mothers of infants born preterm (n=59) completed screensers for mood disorders when their infant was approximately 14 days old, deriving rates of maternal mood disorders, specifically Depression, Anxiety, and PTSD. Rates of mother-delivered developmental care (minutes of
developmental care/days of hospitalization) were documented daily in the electronic medical record throughout the infants’ hospital stay and were extracted for analyses.

Summary of Results No significant correlations were observed between mood scores and developmental care rates normalized for the total length of stay (Depression r = 0.02, p = 0.90; Anxiety r = 0.00, p = 0.99; PTSD r = 0.02 p = 0.87). Rates of developmental care were not significantly different in mothers who were classified as qualifying for referral to mental health services (N=19) compared to those who did not (N=40) (Depression t = 0.48, p = 0.63, Anxiety t = -0.34, p = 0.74; PTSD t = -0.20 p = 0.85 ). Results were similar after controlling for factors that may have affected rates of developmental care, including SES, gestational age, and infant health.

Conclusions While perinatal maternal mood disorders remain an important issue in the NICU, perinatal mood may not be an important factor affecting engagement in developmental care. Findings suggest that maternal engagement in developmental care activities may not serve as an appropriate indicator of mental health issues. As such, perinatal mood screeners should be implemented in infant healthcare environments, to gauge maternal mental health. Further research should be conducted to determine the extent to which perinatal screening for mental health can inform ways to improve maternal engagement in developmental care and other parenting activities both during and after infant hospitalization.

Summary of Results A significant positive correlation was found for BSID-III language scale and maternal frequency of maintaining behavior (R = 0.510, p = 0.001). A significant negative correlation was found for BSID-III language scale and frequency of maternal redirecting behavior (R = -0.383, p = 0.018) and duration of redirecting (R = -0.388, p = 0.018). Using partial correlations to control for maternal education level, significant positive associations between BSID-III language score at 18 mo persisted for frequency and duration of maintaining behavior (R = 0.446, p = 0.004 and R = 0.522, p = 0.001 respectively). In addition, a significant positive correlation between the PSI parent distress scale and duration for redirecting behavior was found (R=0.373, p=0.042) when controlling for maternal education.

Conclusions We demonstrate that early child language development was positively impacted by maternal maintaining behavior and negatively impacted by maternal redirecting behavior during play. Increased maternal stress levels resulted in increased redirecting behavior. Overall, these results suggest higher parental stress may have a negative impact on early child language development.

THE INFLUENCE OF MATERNAL AND CHILD INTERACTIVE BEHAVIORS ON EARLY CHILD DEVELOPMENT

1KN Baille*, 1T Emery, 1L Lowe, 1S Markoe. 1University of New Mexico School of Medicine, Albuquerque, NM; 2University of New Mexico Health Sciences Center, Albuquerque, NM

Purpose of Study The quality of interactions between children and their caregivers significantly impacts a child’s development. Positive maternal behaviors during play are associated with better self-regulation and fewer behavioral problems, while poor parental well-being is associated with increased developmental difficulties. We explored the relationship between positive and negative maternal interactive behaviors during play, parental well-being and cognitive and language development in children aged 18 months old (mo). We hypothesized positive maternal behaviors during play will positively correlate with higher Bayley Scales of Infant Development III (BSID-III) cognitive and language scores at 18 mo and poor parental well-being will lead to more negative directing behaviors and child disengagement.

Methods Used Healthy term infants were recruited in Albuquerque, NM between 2010–2012 as part of a longitudinal study of parenting and child development (n=40). BSID-III was used as an objective measure of cognitive and language development. Parental well-being was measured using a parental stress index (PSI) questionnaire in addition to socioeconomic status (SES). The Maternal Attention Directing Manual (S. Landry 2000) was used to code six minute free-play videos of mother-child dyads, focusing on positive maternal directing behaviors, such as ‘maintain’ and ‘introduce’ and negative behaviors, such as ‘redirect’ or ‘zaps’. Intercoder reliability was greater than 80%.

Purpose of Study Temperament of very young children with sex chromosome trisomy (SCT) has yet to be described in the literature. Young children with developmental differences may present with variation in temperament profiles. Temperament consists of a child’s activity level, rhythmicity, approach or withdrawal tendencies, adaptability, threshold of responsiveness, intensity, mood, distractibility, and persistence; all of which may affect a child’s development and have diagnostic and treatment implications. This project aimed to evaluate temperament profiles in infants with prenatally identified SCTs using standardized measures.

Methods Used The eXtraordinarY Babies study is a longitudinal natural history study examining developmental, medical, and psychosocial factors in children with prenatally identified SCT (XXX, XXY, or XXX). Data from the Carey Temperament Scale (CTS) completed by parents at the 12-month-visit were used for this analysis. The CTS is a validated tool designed to assess innate behavioral patterns in children. Caregivers rated the frequency of behaviors on a 6-point scale, yielding z-scores in 9 categories of temperament characteristics, as well as a rating of overall manageability of the child’s behavior. Descriptive statistics were used to describe the sample, one-sample t-tests were used to compare the CTS sample to the norming sample, and a one-way ANOVA was used to compare SCT groups.

Summary of Results In young children with prenatally identified sex chromosome trisomies (SCTs) are at an increased risk for neurocognitive and behavioral conditions, hypoplastic left heart syndrome, twin to twin transfusion syndrome, and SCAs. The literature. Young children with developmental differences may present with variation in temperament profiles. Temperament consists of a child’s activity level, rhythmicity, approach or withdrawal tendencies, adaptability, threshold of responsiveness, intensity, mood, distractibility, and persistence; all of which may affect a child’s development and have diagnostic and treatment implications. This project aimed to evaluate temperament profiles in infants with prenatally identified SCTs using standardized measures.
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(\(p<.001\)), and high threshold of responsiveness (\(p=.006\)). There were no statistically significant differences between the SCT conditions. A significant majority (82%) of parents rated their child’s overall manageability as ‘easy’ or ‘very easy’.

Conclusions Results indicate that 12-month old infants with SCTs have temperamental differences from the general population. These data confirm parent-report of child behavior, and previous studies which suggest similar temperamental differences in older children with SCTs. The temperament profile found reflects what specific characteristics make these children easier to manage in the first year of life and defines a new constellation of behavioral patterns that comprise what is commonly referred to as an ‘easy baby’ in SCTs. Future analyses will further examine temperamental patterns beyond 12 months of age, the relationship of temperament to development and parent-child relationships, and inform genetic counseling, assessment, and intervention recommendations.

#284 PERINATAL DISTRESS ASSOCIATED WITH NEONATAL CONDITIONS: ANXIETY AND DEPRESSION

D. Cooke*, H. Nasier, J. Kelleher, J. Dempsey, AG. Dempsey. University of Colorado Denver School of Medicine, Aurora, CO

10.1136/jim-2022-WRMC.280

Purpose of Study The purpose of this study was to examine self-reported perinatal depression and anxiety symptomology associated with discrete fetal medical complexities seen in a Fetal Care Center (FCC) in a Children’s Hospital.

Methods Used The Edinburgh Postnatal Depression Scale (EPDS) and General Anxiety Disorder (GAD-7) were administered to all patients at a major FCC as part of a quality improvement project for detection of PMADs in the center. Questionnaires were given to all women presenting for both initial and return appointments, however only data from the initially administered questionnaires were included. The EPDS is a self-report measure of depressive symptoms in antenatal populations, with a cutoff of 11.5 suggested for women with high-risk pregnancies. The GAD-7 is a standardized self-report measure of anxiety symptoms, with a cutoff of 10. Higher scores on both indicate greater symptomatology. Scores and information regarding fetal diagnoses were obtained via retrospective chart review from 141 women and examined descriptively.

Summary of Results The pattern of depressive and anxious symptomatology at first administration varied by medical diagnosis (figure 1). Within the sample 37% reported an EPDS score of 10 or greater, 21% reported an EPDS score of 12 or greater, and 20% reported a GAD-7 score of 10 or greater. Of the observed medical conditions, myelomeningocele, gastrointestinal conditions, and omphalocele were associated with the highest average rates of distress (the average EPDS scores were greater than 10 and GAD-7 scores 9 or greater). Neurologic conditions, hypoplastic left heart syndrome, twin to twin transfusion syndrome, and twins were associated with the second highest average rates of distress (average EPDS scores greater than 8 and GAD-7 scores were greater than 5.75).

Conclusions Certain fetal medical conditions appear to be associated with greater levels of distress in perinatal women. More research is needed to determine if fetal medical conditions increase risk for the development of PMADs. Women with these diagnoses may benefit from increased psychosocial monitoring and support.

#285 EDUCATIONAL NEEDS OF CHILDREN WITH SEX CHROMOSOME ANEUPLOIDIES

1,2TG Thompson*, 1,2J Janusz, 1,2S Davis, 1,2N Tartaglia; 1University of Colorado Denver School of Medicine, Aurora, CO; 2Children’s Hospital Colorado, Aurora, CO

10.1136/jim-2022-WRMC.281

Purpose of Study Children with sex chromosome aneuploidies (SCAs) are at an increased risk for neurocognitive and behavioral disorders that may interfere with success in school and ultimately with quality of life, including early developmental delays, learning disabilities, executive function problems, and social communication deficits. The current study aimed to update and extend our understanding of educational supports and outcomes for students with these increasingly common genetic diagnoses.

Methods Used Parents of children with a diagnosed SCA, birth to 21 years, living in the U.S. (N=248), consented to participate in an electronic survey. Descriptive statistics were used to quantify rates of school support plans, academic accommodations, educational therapies, and school completion. Logistic regression was used to calculate group differences. An inductive qualitative thematic analysis was applied to free-text survey responses.

Summary of Results Results revealed high rates of delayed kindergarten (20%), grade retention (17%), and individualized educational programs (IEPs; 71%). Despite a clear profile of educational need, a majority of respondents with children over age 18 (N=41) reported their child successfully completed high school, and nearly half pursued post-secondary education opportunities. A majority of parents described their child’s teachers as having little to no knowledge of how SCAs can impact a child’s learning. Analysis of open-ended parent responses emphasized that challenges with reading, executive function, reduced stamina, social skills deficits, and emotional dysregulation acted as barriers to learning, and were frequently triggered by busy or noisy classroom environments. Further, skills hovering in the borderline range are common to the SCA phenotype, and are not often well served by
special education systems challenged by limited resources and strict cut-offs for qualification. Parents reported the need to advocate strongly for their child to receive adequate school support services.

Conclusions Pediatricians should be aware of the frequent need for accommodations and individualized support plans in children with SCAs, so they can educate families and advocate for early and comprehensive evaluations and intervention plans. Our findings justify a need to train teachers and policy makers in the unique educational needs of children and adolescents with SCAs. We recommend increased collaboration between medical and educational teams and acknowledgement of the significant role the genetic condition plays in the educational experiences of students with SCAs.

#286 LOW SENSITIVITY OF THE AGES & STAGES QUESTIONNAIRE IN INFANTS WITH SEX CHROMOSOME TRISOMY AT 6 AND 12 MONTHS OF AGE

J. Newar, T.G. Thompson, L. Pyle, J. Ross, N. Tartaglia. Children’s Hospital Colorado, Aurora, CO; University of Colorado Denver School of Medicine, Aurora, CO; Alfred I DuPont Hospital for Children, Wilmington, DE

Purpose of Study Sex chromosome trisomy (SCT) conditions are a group of genetic disorders characterized by an extra X or Y chromosome, including XXY/Klinefelter syndrome and XYY syndrome in males, and Trisomy X syndrome in females. In early childhood, the most common features of SCT include developmental delays, with previous studies showing that 50–70% of young children have delays that require early intervention therapies such as speech therapy and/or physical therapy. The diagnosis of SCT in the prenatal period has drastically increased over the past decade, leading to the need for developmental screening guidelines for the large group of SCT infants at risk for delays. The aim of this project is to evaluate a commonly-used standardized developmental screening measure called the Ages and Stages Questionnaire (ASQ) to determine if it is valid and effective in identifying young children with SCT with developmental delays.

Methods Used The eXtraordinarY Babies Study is a longitudinal natural history study examining developmental, medical, and psychosocial factors in children with prenatally identified SCT (XXY, XYY, or XXX). Data from the 6 and 12 month visits were used for this analysis. Parents (n=121) completed the 6 and 12 month ASQ, and results were compared to direct developmental assessment using the Bayley Scales of Infant and Child Development-3 (BSID-III). Sensitivity, specificity, positive and negative predictive values were calculated for domains of communication, fine motor, gross motor, and problem solving, using both a strict ASQ cutoff (≥2SD below the mean) and an expanded cutoff (≥1.5SD below the mean).

Summary of Results The sensitivity of all analyses ranged from 0.1–0.6, all well below the standard accepted sensitivity of 0.7–0.8, except for the 12-month gross motor analysis. An expanded cut-off improved sensitivity to 0.2–0.9, but results remained below acceptable levels for a reliable screening tool. Specificity rates were high at a range of 0.7–0.10 for all analyses and fell within the standard accepted specificity of 0.7–0.9.

Conclusions Based on findings of low sensitivities and high false negative rates of the ASQ, we conclude that the ASQ is not a good measure to use in the SCT population at 6 or 12 months of age. Future analyses will compare the ASQ to other screening measures to determine if others provide acceptable statistical properties for developmental screening. Based on these results, direct developmental assessment during the first year of life should be part of the care recommendations for all children with SCT until a more rigorous screening measure is identified or developed for this at-risk population.

Diversity, equity and inclusion II

Concurrent session

10:15 AM

Friday, January 21, 2022

#287 DISPARITIES IN PARENT-DOCTOR COMMUNICATION DURING THE HOSPITALIZATION OF CHILDREN WITH MEDICAL COMPLEXITY

S. Hussain, D. Chin, H. Nguyen-Tang, R. Afghani, UC Irvine School of Medicine, Orange, CA; Children’s Hospital of Orange County, Orange, CA

Purpose of Study While children with medical complexity (CMC) represent less than 1% of the pediatric population, they account for one-third of pediatric health care spending. Unfortunately, there is a lack of clinical initiatives and research efforts geared towards improving their care, let alone assessing their health literacy. This population experiences frequent hospitalizations and is often inundated with medical jargon. Clear communication from providers is a key component of health literacy and empowers families to make medical decisions. The objective of this study is to assess communication among physicians and parents prior to discharge.

Methods Used A survey via REDCap was distributed to parents of CMC on the day of discharge from the hospital. The survey was based on the CAHPS Health Literacy survey, a validated tool used to examine the communication barriers that providers can target to improve patient and caretaker’s understanding. The survey was modified for the pediatric inpatient population.

Summary of Results Twenty surveys were collected. 20% said they speak English not well or none at all. About 66% of the parents had completed high school and some college and the remainder had a college or postgraduate degree. A majority of respondents (72%) identified themselves as belonging to an
Abstracts

#288 A CALL TO CREATE EVIDENCE-BASED, UPSTREAM PREVENTION PROGRAMS TARGETING YOUTH MENTAL HEALTH THAT ARE EQUIVALENT ACROSS ETHNIC/RACIAL SUBGROUPS: ADVOCATES FOR ALL YOUTH (ALLY)

1J Kaar*, 2AE Bowen, 3M Pangelinan, 4A Dademathews, 5S Simon, 6L Shomaker, 7University of Colorado Anschutz Medical Campus, Aurora, CO; 2Children’s Hospital Aurora, Aurora, CO; 3Auburn University, Auburn, AL; 4Colorado State University, Fort Collins, CO

Purpose of Study Youth from disadvantaged racial/ethnic groups have higher rates of anxiety and depression compared to non-Hispanic White youth; however, are less likely to receive mental healthcare. Factors that may underlie these disparities include acculturation pressures and stressors, trauma, peer victimization and microaggressions, and discrimination. With mental health distress exacerbated during and in the wake of the COVID-19 pandemic, there is a need for evidence-based school-based programs that have the potential to reach youth who otherwise may not have access to mental healthcare.

Methods Used ALLY was created based on a review of existing school-based, mental health programs and with input from key stakeholders.

Summary of Results Prevention programs that target the improvement of individual-level protective factors exist, however, tend not to be equitable across ethnic/racial subgroups, especially among low-income Black and Hispanic communities. Schools are viewed as an attractive platform for such programs because they can alter the environment and education of a large number of youths simultaneously. Further, when implemented successfully into the school's structure, have the potential to be integrated and sustained long-term. Despite these advantages, few school-based efforts focused on improving protective factors, such as self-efficacy and resilience, to reduce symptoms of anxiety and depression have been conducted.

Conclusions With continued increases in mental health distress, there is a need for school-based programs that have the potential to reach youth from disadvantaged backgrounds. ALLY is a school-based positive psychology program aimed to increase youth's well-being and reduce symptoms of depression and anxiety. Facilitators are selected from the community in which the school is located and receive training in supportive conversations and cultural competency & humility as well as how to reach youth with varying cultural, gender identity, or neurodiversity differences that may exist. ALLY is examining mental health outcomes among diverse school settings in terms of program participation, efficacy, and implementation. Our findings will inform to what extent the program is efficacious among underrepresented communities.

#289 SOCIOECONOMIC DIFFERENCES ASSOCIATED WITH CONSUMPTION OF A PLANT-BASED DIET: RESULTS FROM THE NATIONAL HEALTH AND NUTRITION EXAMINATION SURVEY

M Gonzalgo*, S Nackeeran, A Mouzannar, R Blachman. University of Miami School of Medicine, Miami, FL

Purpose of Study A plant-based diet (PBD) has been associated with potential health benefits, but factors that may affect access to and consumption of a PBD are not well defined. The purpose of this study was to determine the association between plant-based dietary content, racial, ethnic, and socioeconomic status among participants enrolled in the National Health and Nutrition Examination Survey (NHANES).

Methods Used This was a cross-sectional study using data obtained from the NHANES database. The following covariates were assessed: age, sex, race/ethnicity, educational level, marital status, smoking status, physical activity, alcohol use, history of diabetes, and hypertension. Socioeconomic status was categorized according to poverty-income ratio (PIR). Food frequency questionnaires were used to calculate plant-based diet index (PDI) and healthful plant-based diet index (hPDI). Multivariable-adjusted logistic regression was performed to determine the association between PIR, clinical, demographic, and plant-based diet indices.

Summary of Results A total of 5,037 participants were included in the final cohort. Median age of participants was 51 ± 18.5 years. Overall PDI and hPDI were 50 (46 – 54) and 52 (47 – 57), respectively. Median PDI consumption was significantly different among PIR groups (PDI, p = 0.018; hPDI, p < 0.001). On multivariable analysis, participants in the poorest socioeconomic group (PIR ≤ 130%) were more likely to have lower consumption of a healthful PBD (hPDI).

Conclusions Lower socioeconomic status (PIR ≤ 130%) was associated with decreased consumption of a healthful plant-based diet. These data suggest that socioeconomic disparities...
may limit consumption of healthier food and contribute to the high prevalence of adverse health conditions that exist in certain population groups.

#290 BILLINGS, MONTANA LGBTQ+ HEALTHCARE ALLIES TRAINING
M Turner*, T Keys. University of Washington, Seattle, WA
10.1136/jim-2022-WRMC.286

Purpose of Study The aim of the project was to improve LGBTQ+ care in Billings (pop. 125,000 and Montana’s largest city) by offering a training to healthcare providers and support staff that educates on how to provide affirming healthcare and reduce barriers to access for the LGBTQ+ population.

Methods Used Bridgercare is a family planning clinic located in Bozeman that offers educational workshops such as the LGBTQ+ Healthcare Allies Training. The 3-hour training educates healthcare providers and support staff and seeks to provide a safe environment for participants to engage with facilitators and a 20-page booklet to follow along with. The evidenced-based booklets are kept by participants and encouraged to be shared after training. The training is led by three facilitators, who either: have experience providing affirming care in a medical setting, belong to an LGBTQ+ community, and/or are allies of LGBTQ+ communities. Bridgercare offered to come to Billings, provide a training, and partner to establish a long-term education class for the Billings area. The training was held at a local museum. Recruitment for attending the training was done through word of mouth and social media. Post-training surveys were given following training.

Summary of Results 30 healthcare professionals attended the first training. Individuals came from 9 different organizations. The training consisted of administration leadership, physicians, PAs, NPs, RDs, nurses, counselors, professors, pharmacists, and care managers. Surveys indicated that 100 percent of individuals felt the training was highly beneficial and learned something. 7 people who were at the training signed up to receive info on how to become a facilitator.

Conclusions Given the positive feedback, next steps will be to partner with another community partner and to train 10 facilitators by December 2021. Trainings with facilitators from Billings will start January 2022. During Spring 2022, training will start to be assessed for how training is impacting LGBTQ+ care. Additional metrics to see whether training is an effective intervention may include measuring: the amount of healthcare workers with formal LGBTQ+ education, healthcare worker confidence in treating LGBTQ+ patients, and the amount of healthcare workers who believe their clinic or department adequately addresses the needs of their LGBTQ+ patients.

#291 PRESENTATION OF ERYTHEMA IN ABSCESSES FOUND IN SKIN OF COLOR
1S Bodapati*, 2C Wong, 3S Weis. 1Western University of Health Sciences, Pomona, CA; 2University of North Texas Health Science Center, Fort Worth, TX; 3Medical City Fort Worth, Fort Worth, TX
10.1136/jim-2022-WRMC.287

Purpose of Study Erythema, one of the cardinal signs of inflammation, is one of the hallmark descriptors of a skin abscess. In individuals with lighter skin, such as Caucasians, erythema presents markedly differently from unaffected skin tissue. In individuals with darker skin tone, abscesses may not be as strikingly apparent when discerned based on appearance of erythema alone. The masking of erythema by darker skin tones is a result of differences in skin biology and the overall composition of skin. It highlights the need for dermatologists and medical professionals to rely on other signs of inflammation when diagnosing skin abscesses.

Methods Used Two cases of patients with skin abscesses were selected from University of North Texas Health Science Center and John Peter Smith Hospital. One male and one female case was selected. Photographs were captured in clinic and image analysis of erythema was performed visually.

Summary of Results Variance in the degree of skin pigmentation results from differences in skin composition, and this changes the appearance of erythema. Differences include denser stratum corneum, a greater ratio of eumelanin to pheomelanin, and more oval-shaped melanocytes in darker skinned individuals. Erythema is a result of pathologic increase in inflammation and red blood cells. Hemoglobin is the primary chromophore in red blood cells, and reflects visible red light, causing the typical red appearance. However, as demonstrated in these presentations, appreciating erythema visually as redness is unreliable in darker skinned individuals, because the red hue of oxyhemoglobin often presents as violaceous, brown, or black. While erythema is commonly one of the first described features of an abscess, relying on indicators such as swelling may prevent delays in diagnosis, particularly in people of color.

Conclusions More documentation of the presentation of inflammatory conditions in individuals with skin of color will help build knowledge in diagnosis of dermatologic conditions. Focusing on color-independent markers of inflammation may provide a more equitable approach to patient care.

#292 DISPARITIES IN USE OF SYRINGE SERVICES PROGRAMS IN KING COUNTY, WA
1KR Salawu*, 2S Glick. 1University of Washington School of Medicine, Seattle, WA; 2University of Washington Department of Medicine, Seattle, WA
10.1136/jim-2022-WRMC.288

Purpose of Study Syringe service programs (SSPs) are an effective harm reduction resource for people who use drugs (PWUD). Recently published cross-sectional studies found that SSPs in two East Coast cities (Miami and New York) were not reaching Black communities. Additionally, qualitative literature showed that women who use drugs experience greater stigma than men and they may feel less inclined to seek harm reduction services. No papers have been published regarding disparities among SSPs in West coast cities, which have different drug use trends and demographics. We used data from the Seattle area to identify disparities in SSP use, and hypothesized that Black PWUD, woman, and gender minorities would be underrepresented in a large Seattle SSP.

Methods Used We compared data from two cross-sectional surveys. One survey was conducted in 2019 at three sites that are part of Public Health – Seattle & King County’s SSP, and evaluated the demographics of 432 SSP clients. The
comparison survey, the 2018 National HIV Behavioral Surveillance Injection Drug Use (NHBS-IDU) Survey, utilized respondent-driven sampling to create a representative sample of people who inject drugs (PWID) in the region; 555 PWID in the Seattle area completed this survey.

The primary variables of interest were race and gender. Exploratory covariates included age, men who have sex with men (MSM), and housing status. We used chi-squared tests to evaluate differences between people using the SSP and PWID in the Seattle area.

**Summary of Results** The majority of SSP clients identified as white, which was similar to the demographics of the NHBS-IDU survey participants (p=0.086). American Indian/Alaska Native and Black PWID were underrepresented in the SSP client survey (p<0.001 for both). Latinx/Hispanic and Native Hawaiian/islander PWID were also underrepresented (p=0.014 and p=0.039 respectively). Approximately, two-thirds of SSP clients were men and one-third were women, with <1% of clients reporting another gender identity and <1% reporting being transgender. The gender of SSP clients largely reflects the same distribution seen in a community sample of PWID (p=0.378). Participants in the SSP survey tended to be younger and less likely to be experiencing homelessness than the NHBS-IDU participants (p<0.001 for both). The distribution of MSM participants in the two surveys were similar (p=0.409).

**Conclusions** This study provides further evidence that Black PWID are underrepresented in SSPs, consistent with studies in other large US cities. Both nationally and in Seattle, overdose deaths have been increasing among Black PWID, and harm reduction strategies are vital to reversing this trend. SSPs should explore and test ways to be more accessible to underrepresented populations.

### Abstract #292 Table 1 Comparison of SSP users and PWID in King County, WA

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>SSP Survey Participants 2019 N=432 N (%)</th>
<th>NHBS-IDU Survey Participants 2018 N=555 N (%)</th>
<th>χ² test p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>18–29</td>
<td>101 (23.4)</td>
<td>100 (18.0)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>30–39</td>
<td>163 (37.8)</td>
<td>164 (29.6)</td>
<td></td>
</tr>
<tr>
<td>40–49</td>
<td>94 (21.8)</td>
<td>127 (22.9)</td>
<td></td>
</tr>
<tr>
<td>50+</td>
<td>73 (16.9)</td>
<td>164 (29.6)</td>
<td></td>
</tr>
<tr>
<td>Gendera</td>
<td></td>
<td></td>
<td>0.378</td>
</tr>
<tr>
<td>Women</td>
<td>147 (34.0)</td>
<td>211 (38.0)</td>
<td></td>
</tr>
<tr>
<td>Men</td>
<td>281 (65.1)</td>
<td>337 (60.7)</td>
<td></td>
</tr>
<tr>
<td>Transgender</td>
<td>3 (0.7)</td>
<td>7 (1.3)</td>
<td></td>
</tr>
<tr>
<td>Other Gender Identity</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>American Race</td>
<td></td>
<td></td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>American</td>
<td>53 (12.3)</td>
<td>125 (22.5)</td>
<td></td>
</tr>
<tr>
<td>Native</td>
<td>330 (76.4)</td>
<td>397 (71.5)</td>
<td>0.086</td>
</tr>
<tr>
<td>Asian</td>
<td>21 (4.9)</td>
<td>14 (2.5)</td>
<td>0.049</td>
</tr>
<tr>
<td>Asian/South</td>
<td>26 (6.0)</td>
<td>110 (19.8)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Black/African American</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Native</td>
<td>9 (2.1)</td>
<td>25 (4.5)</td>
<td>0.039</td>
</tr>
<tr>
<td>Latinx/Hispanic</td>
<td>33 (7.6)</td>
<td>69 (12.4)</td>
<td>0.014</td>
</tr>
<tr>
<td>Hawaiian/Pacific Islander</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>White</td>
<td>330 (76.4)</td>
<td>397 (71.5)</td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>12 (2.8)</td>
<td>–</td>
<td></td>
</tr>
<tr>
<td>Missing</td>
<td>0 (0.0)</td>
<td>10 (1.8)</td>
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</tr>
<tr>
<td>MSM</td>
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<td></td>
<td>0.409</td>
</tr>
<tr>
<td>No MSM</td>
<td>236 (84.0)</td>
<td>297 (86.3)</td>
<td></td>
</tr>
<tr>
<td>Yes MSM</td>
<td>45 (16.0)</td>
<td>47 (13.7)</td>
<td></td>
</tr>
<tr>
<td>Housing Status</td>
<td></td>
<td></td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Currently homeless</td>
<td>198 (45.8)</td>
<td>338 (60.9)</td>
<td></td>
</tr>
<tr>
<td>Other housing status</td>
<td>234 (54.2)</td>
<td>217 (39.1)</td>
<td></td>
</tr>
</tbody>
</table>

a) SSP participants could select more than one gender, while NHBS-IDU participants could not. b) Both SSP and NHBS-IDU participants could select more than one race.
Purpose of Study
People living in rural and frontier areas experience higher morbidity and mortality in part due to limited access to health care. Whether rurality affects care and outcomes for very low birthweight (VLBW) infants is not well understood. This study compares prenatal factors, care patterns, and health outcomes between VLBW infants born to mothers living in rural versus urban zip codes in California.

Methods Used
We used data from the California Perinatal Quality Care Collaborative (CPQCC) registry to conduct a population-based retrospective cohort study of VLBW infants born between 2011 and 2018. This descriptive analysis examined associations between the degree of maternal residential rurality and maternal, infant, hospital, and community/county level characteristics based on Rural Urban Commuting Area (RUCA) codes. The associations between all factors and maternal RUCA codes were stratified and compared using Chi-squared and ANOVA tests.

Summary of Results
Our study included 38,614 VLBW infants. Of those, 36,936 were born to mothers residing in an urban area, 1,113 in a large rural area, and 561 within a small rural or isolated area. Mothers of VLBW infants who live in large rural and small rural/isolated areas were more likely to be younger, publicly insured, have education limited to high school, smoke during pregnancy, and have delayed or no prenatal care.

Infants born to mothers living in large rural and small rural/isolated areas were more likely to have their first hospital within the CPQCC network as a level 4 NICU with greater than 110 annual VLBW admissions but with less than 3500 annual deliveries compared to urban infants.

Birth weight, gestational age, and infant sex were similar across geographic groups. Survival without major morbidities decreased across the rural-urban continuum, with urban maternal residence having the highest percentage of VLBW infants surviving without major morbidities.

Conclusions
VLBW infant survival without major morbidity decreased with increasing rurality. This descriptive study highlights various risk and protective factors which may modulate this association, and which suggest the possibility for health policy intervention.

#294
RISK FACTORS FOR MORBIDITY AND MORTALITY IN VERY LOW BIRTH WEIGHT INFANTS FROM RURAL VERSUS URBAN COMMUNITIES

10.1136/jim-2022-WRMC.290

Purpose of Study
Surveying California family physicians and Los Angeles County pharmacists: pill prescribing and dispensing practices.

Methods Used
After we obtained a list of all pharmacies in each LA County zip code using Yellow Pages, we organized them by the 8 LA County Service Planning Areas (SPAs). We chose how many pharmacies to contact in each SPA based on the percentage of women living in each SPA. We contacted a proportional number of national and local pharmacies in each SPA based on the distribution of pharmacies in that area. A standardized secret shopper script was used by all researchers when calling pharmacies. We recorded if the pharmacy agreed to dispense 13 packs and if not, how many packs they would dispense and why, among other parameters. Pharmacies were classified as unable to determine if they stated they would need us to come into the pharmacy to determine prescription and insurance eligibility.

Conclusions
While the regulations requiring pharmacies to dispense a one-year supply of oral contraceptive pills to Family PACT patients who have an appropriate prescription was introduced in January 2017, the lack of knowledge of this insurance program and its requirements by pharmacies in this area, 1,113 in a large rural area, and 561 within a small rural/isolated area were more likely to be younger, publicly insured, have education limited to high school, smoke during pregnancy, and have delayed or no prenatal care.

Infants born to mothers living in large rural and small rural/isolated areas were more likely to have their first hospital within the CPQCC network as a level 4 NICU with greater than 110 annual VLBW admissions but with less than 3500 annual deliveries compared to urban infants.

Birth weight, gestational age, and infant sex were similar across geographic groups. Survival without major morbidities decreased across the rural-urban continuum, with urban maternal residence having the highest percentage of VLBW infants surviving without major morbidities.

Conclusions
VLBW infant survival without major morbidity decreased with increasing rurality. This descriptive study highlights various risk and protective factors which may modulate this association, and which suggest the possibility for health policy intervention.

Healthcare delivery research II
Concurrent session
10:15 AM
Friday, January 21, 2022

#295
SURVEYS OF CALIFORNIA FAMILY PHYSICIANS AND LOS ANGELES COUNTY PHARMACISTS: PILL PRESCRIBING AND DISPENSING PRACTICES

I. Gedestad*, A. Matlack, M. Munnangi, A. Nelson. Western University of Health Sciences, Pomona, CA

10.1136/jim-2022-WRMC.291

Purpose of Study
The California Family PACT insurance program provides family planning and reproductive health services at no cost to at least half a million women ages 15–44 who fall below 200% of the Federal Poverty Line and who are at risk for unintended pregnancy. As shown in previous studies, women who received 13 packs (a 1 year supply) of oral contraceptives were less likely to become pregnant, conceive an unplanned pregnancy, or receive an abortion when compared to women who received one or three cycles of pills. The purpose of this study was to determine whether women insured by the Family PACT program are able to receive 13 packs of oral contraceptive pills at once from their pharmacies as now required by CA law.

Methods Used
After we obtained a list of all pharmacies in each LA County zip code using Yellow Pages, we organized them by the 8 LA County Service Planning Areas (SPAs). We chose how many pharmacies to contact in each SPA based on the percentage of women living in each SPA. We contacted a proportional number of national and local pharmacies in each SPA based on the distribution of pharmacies in that area. A standardized secret shopper script was used by all researchers when calling pharmacies. We recorded if the pharmacy agreed to dispense 13 packs and if not, how many packs they would dispense and why, among other parameters. Pharmacies were classified as unable to determine if they stated they would need us to come into the pharmacy to determine prescription and insurance eligibility.

Conclusions
While the regulations requiring pharmacies to dispense a one-year supply of oral contraceptive pills to Family PACT patients who have an appropriate prescription was introduced in January 2017, the lack of knowledge of this insurance program and its requirements by pharmacies in this area, 1,113 in a large rural area, and 561 within a small rural/isolated area were more likely to be younger, publicly insured, have education limited to high school, smoke during pregnancy, and have delayed or no prenatal care.

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Birth weight, gestational age, and infant sex were similar across geographic groups. Survival without major morbidities decreased across the rural-urban continuum, with urban maternal residence having the highest percentage of VLBW infants surviving without major morbidities.

Conclusions
VLBW infant survival without major morbidity decreased with increasing rurality. This descriptive study highlights various risk and protective factors which may modulate this association, and which suggest the possibility for health policy intervention.
Abstracts

CHARACTERIZING OPIOID USER TYPES AND IDENTIFYING PRESCRIBING FACTORS CONTRIBUTING TO LONG TERM USE

R Xu, J Bone, R Coutremanche, L Yetef, M Bueno, M Simmonds, E Cartoni, G Lauder, D Coutremanche. The University of British Columbia Faculty of Medicine, Vancouver, BC, Canada; Children’s Hospital, Vancouver, BC, Canada; University of Toronto Temerty Faculty of Medicine, Toronto, ON, Canada

Purpose of Study Prescription opioid use is associated with subsequent risk of opioid use disorder and opioid-related death. Variability of prescription practice may further escalate this risk. This is paramount in British Columbia (BC); BC had the highest 2020 Canadian provincial opioid overdose related deaths per capita. Our recent study described BC outpatient opioid prescription patterns (2013–2017). Opioid user types (single, short, intermediate, and long-term) are characterized here with the same dataset to identify prescription factors that increase long-term use.

Methods Used Anonymized data of all BC outpatient opioid prescriptions (2013–2017): patient age and sex, prescriber type, generic drug, dose, and total days of opioid supply, was used. Incident episodes that began within the study timeline were included; those on opioid agonist therapy were excluded. Opioid episode durations: time between first and last day of prescription with none in the following 180 days. Opioid user types: single (one prescription); short-term (<90-day); intermediate (≥90-day: <10 prescriptions & <120-day supply); and long-term (≥90-day: ≥10 prescriptions over >90 days or ≥120-day supply).

Opioid user types were described by episode number, median total daily supply, median total MME (morphine milligram equivalents) prescribed, first prescribing specialist and first opioid prescribed.

Incidence/prevalence of user types, and prescription trends were described. Hazard ratios for prescription factors associated with long-term use were estimated with cox regression models.

Summary of Results 1,920,073 incident episodes (8,635,831 prescriptions) over the 5 years were classified as single (70%), short (17%), intermediate (9%), and long-term (4%). Median episode length was 5, 13, 15, and 202 days, respectively. Median total MME prescribed was 4.5, 11.3, 13.5, and 79 MME, respectively. General practitioners were the first prescribers for >50% of episodes across all user types apart from single (45%). First opioid prescribed across all user types was codeine (50%-70%), tramadol/tapentadol (19%), and hydromorphone (5%-15%).

Incidence of long-term users decreased over the 5 years (664 to 537 per 100,000) while prevalence remained similar (3000 per 100,000). Factors associated with longer episodes included initial prescription of oxycodone (HR 0.82) or hydromorphone (HR 0.85) relative to codeine; first prescriber specialty: psychiatry (HR 0.79) or diagnostic services (0.89) relative to general practice; and increasing age (HR 0.97 by 5-year increments).

Conclusions This population-level data provides a better understanding of the role of prescriber types/prescriptions and factors that contribute to increased likelihood of long-term use.

BARRIERS TO ACCESSING HEALTHCARE BY THE FEMALE HOMELESS POPULATION: A SURVEY OF SERVICE PROVIDERS

A Akcam*, AL Nelson, M Fraix. Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA

Purpose of Study This study surveyed service providers to assess perceived barriers that the growing population of homeless, reproductive-age women face accessing health care from the view of those who work with them in San Diego County.

Methods Used Between October 2020 and February 2021, 25 organizations working with unsheltered women were sent links to an anonymous, voluntary survey to distribute to English-speaking employees over age 18. It assessed perceived barriers; subgroup analysis was done by participant role in the organization. IRB approval was obtained.

Summary of Results Eighteen of the contacted agencies forwarded the survey to staff; 49 members participated; 38 completed it entirely. Participating organizations included clinics (70.2%), housing programs (15%), and outreach and/or school programs (11%). Of respondents, 42% are nurses, 17% are clinicians, 15% were members of the management team, and 8.5% were nursing students.

The top ‘major barriers’ endorsed by service providers to homeless women accessing care were lack of transportation (73.8%), cost of care (71.4%), and unfamiliarity with resources (64.3%).

Regarding menstrual care, 38 participants reported that women used pads (32.5%), tampons (28%) and other products (tie shirts, socks, etc.) (25%); 60% provided menstrual supplies. Cost/unfamiliarity with places for low-cost/free products (60%), lack of privacy for use (37.5%), and lack of storage (35%) were endorsed as the top ‘major barriers’ to care by service providers.

43 participants estimated the number of clients who were sexually active; 21% estimated 25–50%; 40% estimated 50–75%, and 18.4% estimated ≥ 75%. Nearly 2/3’s said their organization provided some contraceptive supplies to homeless women; 41% provided barrier methods, 18% provided both injectable and oral birth control. Major barriers included cost (76.3%) and unfamiliarity with free contraception sites (50%).

Conclusions Unsheltered women face numerous financial and knowledge barriers to care. Based on our results, the best options to increase access to care for homeless women include decreasing costs, and educating women on locations of available products and services. Additionally, organizations like mobile clinics may benefit women more as they eliminate the transportation barrier.

BARRIERS AND FACILITATORS TO PRIMARY CARE ACCESS FOR REFUGEES IN SPOKANE, WASHINGTON

K Glover*, A White, AM Cole. University of Washington School of Medicine, Seattle, WA

Purpose of Study Refugees in the United States lack access and quality of care, and barriers contribute to their poor health. To identify barriers and facilitators to primary care access, a mixed-methods survey was distributed in a refugee population in Spokane, Washington.

Methods A mixed-methods survey was distributed to refugees in Spokane, Washington, and data was collected from September 8, 2021 to November 1, 2021. The survey included a combination of open-ended and closed-ended questions on barriers and facilitators to primary care access.

Summary of Results The survey was completed by 48 refugees, who identified several barriers to primary care access, including language barriers, transportation barriers, lack of insurance, and lack of awareness of available services. Additionally, refugees identified several facilitators, including the availability of interpreters, the use of mobile clinics, and the reception of cultural and linguistic considerations.

Conclusions The findings of this study highlight the need for greater access and quality of care for refugees in the United States. Programs and practices that address language barriers, transportation barriers, lack of insurance, and lack of awareness of available services are essential to improving the health status of refugees.

Supported by the National Institutes of Health (NIH) grant T32HD090868.
Purpose of Study The United States has resettled over three million refugees and Spokane, WA is home to a community of over 10,000 refugees. There is a paucity of research on barriers and facilitators for refugee primary care access (PCA). This study sought to identify barriers and facilitators to PCA amongst resettled refugees in Spokane, WA.

Methods Used This study emphasizes community participation and was developed with local healthcare and refugee assistance organizations (CBO’s). Snowball sampling generated key informants from initial referrals from CBOs. Eligible participants were: 1) refugees resettled in Spokane through a formal resettlement agency; 2) healthcare workers who care for refugees; 3) resettled refugees who now provide refugee services. A semi-structured interview guide was developed for each group assessing known barriers and facilitators to PCA, as well as questions developed collaboratively with CBO’s to explore refugees’ experience of the COVID-19 pandemic. Thematic analysis was performed on the transcribed interview text using an established coding schema of supply- and demand-side barriers and facilitators. Demand-side barriers are defined as individual, household or community characteristics that influence the demand for services, and supply-side as ‘characteristics of the health system that exist beyond the control of potential health service users, such as health facilities, drugs, equipment, finances, human resources, geographic distance, etc.

Summary of Results We interview 24 individuals including refugees (N=9), care providers (N=9), and refugees who have become providers (N=6). Thematic analysis (ongoing at this time) revealed extant supply-side barriers including the complexity of the US healthcare system and inadequate interpretation. Novel supply-side barriers specific to the COVID-19 pandemic emerged such as loss of in-person interpretation services, as well as demand-side barriers including the spread of COVID misinformation. Novel supply-side facilitators include use of communication technologies to share health information. This is one of the first studies of barriers and facilitators to refugee PCA to directly engage refugees resettled on the West Coast. It is also the first such study done in Eastern Washington.

Conclusions These data contribute to the literature on refugee healthcare access in the US in general, as well as in Spokane, WA. This is among the first studies of barriers and facilitators to refugee PCA to assess the impact of the COVID-19 pandemic. These findings offer evidence for CBO’s and local and state governments to implement policies that remove barriers and leverage facilitators to refugee PCA.

Abstract #299 Table 1

<table>
<thead>
<tr>
<th>Chief Complaint</th>
<th>Percent</th>
<th>Total (n=359)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wound</td>
<td>38.2%</td>
<td>137</td>
</tr>
<tr>
<td>Medication-related</td>
<td>20.3%</td>
<td>73</td>
</tr>
<tr>
<td>Pain</td>
<td>18.4%</td>
<td>66</td>
</tr>
<tr>
<td>Follow-up</td>
<td>14.5%</td>
<td>52</td>
</tr>
<tr>
<td>Cardiovascular</td>
<td>12.8%</td>
<td>46</td>
</tr>
<tr>
<td>Infectious</td>
<td>11.7%</td>
<td>42</td>
</tr>
<tr>
<td>Respiratory</td>
<td>10.9%</td>
<td>39</td>
</tr>
<tr>
<td>Other</td>
<td>9.7%</td>
<td>35</td>
</tr>
<tr>
<td>Trauma</td>
<td>8.4%</td>
<td>30</td>
</tr>
<tr>
<td>Mental Health</td>
<td>8.1%</td>
<td>29</td>
</tr>
</tbody>
</table>

#299 AN ANALYSIS OF STREET MEDICINE CHIEF COMPLAINTS AND TREATMENTS FOR THE HOMELESS POPULATION OF SPOKANE, WASHINGTON

AS Hoppe*, T Kagele, B Hall, BD Messner, A Nichols. Washington State University, Spokane, WA

Purpose of Study The purpose of this study is to evaluate the chief complaints treated by the Spokane Street Medicine team for purposes of resource requirements and allocation.

Methods Used In this analysis, a retrospective chart review of the RANGE Spokane Street Medicine Team was completed on encounters seen between January 2021 and March 2021. To

begin this process, 359 paper charts were transcribed into a Qualtrics format for analysis. Categories included chief complaints, type of chief complaint (acute or chronic), treatment received, and demographic information. These categories were confirmed in meetings with RANGE Street Medicine team members. A qualtrics analysis of basic chief complaint and treatment frequencies was then performed. This is an ongoing analysis.

Summary of Results Primary analysis revealed that the majority of street medicine chief complaints were acute conditions. Of these, the top 5 complaints were Wounds (38.2%), Medication-Related (20.3%), Pain (18.4%), Follow-up (14.5%), and Cardiovascular (12.8%) (see table 1.1). Mental health was the 10th most common complaint. In terms of treatment, follow-ups were scheduled with local organizations for ⅓ of patients (33.93%). Medications were prescribed to 122 patients (39.97%). No resuscitations were performed or Narcan used.

Conclusions Wound care, medication-related complaints, and pain accounted for the majority of complaints from the Spokane, WA Street Medicine team. Follow-up proved to be another commonly addressed problem, suggesting the difficulties of continuity of care within the homeless population. As such, team resources should be directed towards wound care equipment and assisting with health coordination within the Spokane community. While there is still need for other types of acute care, the team may be best served by pursuing clinic and pharmacy connections for efficient treatment and follow-up.

#300 TEXT MESSAGING INTERVENTION TO PROMOTE CARDIAC HEALTH AND COMBAT LONELINESS DURING SOCIAL DISTANCING

1K Uhls*; 2,3K Watson; 2,3T Horwich; 2,3M Callon-Press. 1University of California Los Angeles, Los Angeles, CA; 2Ronald Reagan UCLA Medical Center, Los Angeles, CA; 3University of California Los Angeles David Geffen School of Medicine, Los Angeles, CA

Purpose of Study The COVID-19 pandemic, with its physical distancing, can lead to loneliness, isolation, and unhealthy behaviors. We aimed to develop an intervention that could address these known cardiovascular risks and hypothesized
that a health text-based message intervention (HTMI) could mitigate unhealthy behaviors and increase healthy behaviors.

Methods Used We prospectively recruited 101 adults in Southern California. After informed consent, participants received HTMIs for 31 consecutive days. Self-reported health behaviors were assessed by the Health Promoting Lifestyle Profile (HPLP). Subjective loneliness was measured by the UCLA Loneliness Scale 3 (LL), and isolation was measured objectively by the Social Network Index (SNI). Surveys were analyzed via paired T test and regression analysis. HPLP and LL were scored on a 4-point scale per question.

Summary of Results Participants’ median HPLP score increased significantly 2.67 to 2.82 signaling increased health behaviors, and the median LL score decreased signaling decreased loneliness 2.08 or 2.01. SNI showed no significant change in social contacts. 75% of participants reported increased knowledge of heart health and 81% reported increased health awareness.

Conclusions HTMIs demonstrated rapid and efficient delivery to a population affected by the COVID-19 pandemic. HTMIs were associated with increased health behaviors and decreased self-reported loneliness.

#301 RELATIONSHIP BETWEEN PHYSICAL ACTIVITY VITAL SIGN AND HEALTHCARE SYSTEM-utilization

Purpose of Study Insufficient physical activity (PA) is a leading risk factor for major chronic diseases and all-cause mortality. Engaging in regular PA is associated with improved health, lower risks of chronic disease, and lower all-cause mortality. Insufficient PA has been associated with substantial costs to healthcare systems globally, although reports on these relationships are few, and the association between insufficient PA and healthcare utilization is less known. This study examined the association between self-reported PA with healthcare system utilization for adults in a single academic healthcare system in the United States.

Methods Used The Physical Activity Vital Sign (PAVS) is a standardized and validated tool for collecting self-reported PA from patients into electronic health records. PAVS data from 23,914 patients were collected at outpatient visits between 2018–2020. Association between 6 months of healthcare utilization and the concurrent and prior 6 months (12 months total) of PAVS was evaluated every 6 months for each patient. Generalized estimating equations were used to assess relationships between PAVS and hospital inpatient admissions, emergency department visits, urgent care visits, and primary care outpatient visits respectively.

Summary of Results Sufficient aerobic PA per public health guidelines of ≥ 150 mins/wk was associated with fewer hospital admissions (p<0.001), emergency department visits (p<0.001), and primary care visits (p<0.001) after adjusting for age, sex, body mass index (BMI), and race. PAVS did not show a statistically significant relationship with urgent care visits. Older patients had a larger association between PAVS and fewer hospital admissions and primary care visits (p=0.002). Sex, BMI, and race were not found to modify the association between PAVS and healthcare utilization.

Conclusions We found that higher PAVS is associated with fewer hospital admissions, emergency department visits, and primary care visits. In the public health setting, these findings suggest that PA may be an important modifiable risk factor for health outcomes and healthcare system utilization. Clinically, the results highlight the importance of medical providers assessing PAVS routinely with patients. Implementation of PAVS in the outpatient setting can help identify patients who need support to become more active for their health. Methods and challenges of implementing PAVS as well as increasing PA should be explored. Finally, further research is needed to understand the potential contribution of increasing PA to reductions in healthcare system costs.

#302 THE IMPACT OF DIRECT-TO-HOME TELEMEDICINE VISITS ON PARENT, PHYSICIAN, AND THERAPIST EXPERIENCE FOR CHILDREN WITH SPECIAL HEALTHCARE NEEDS

Purpose of Study Pediatric physiatrists provide necessary specialty care to children with physical disabilities. However, pediatric physiatrists are scarce and are concentrated at academic medical centers. To expand access to care for children with special healthcare needs, UC Davis created the School-Based Tele-Physiatry Assistance of Rehabilitative and Therapeutic Services (STARS) Program, a novel telemedicine program that connects pediatric physiatrists to patients receiving in-person care from a therapist at a local medical therapy unit (MTU). As a result of the COVID-19 pandemic, this model of care was disrupted, as MTUs moved to direct-to-home video visits with patients. The purpose of this qualitative study was to examine the experience of parents, therapists, and physiatrists on direct-to-home tele-physiatry visits with the goal of informing future use of telemedicine for children with special healthcare needs.

Methods Used Parents were recruited from a list of randomly generated parents of patients who had participated in the STARS program. All therapists and physicians providing care at the participating MTUs were invited to participate. We used a semi-structured guide based on the domains of the Practical Robust Implementation Sustainability Model (PRISM). Following principles of qualitative content analysis, two researchers separately coded each transcript for themes and subthemes using line by line coding in the Dedoose software program. Themes and sub-themes were developed using the constant comparative method and were refined by the larger study team.

Summary of Results Six focus groups (4 parent groups, 1 therapist group, and 1 physician group) were conducted before thematic saturation was reached. Direct-to-home telemedicine visits were more effective when physicians had previously established a therapeutic alliance with patients. Although in-person visits were perceived as higher quality, physicians, therapists, and parents identified aspects of care that improved over telemedicine, including using tools within the patient’s home for therapy, understanding the child’s environment, and practicing daily tasks. Parents saw direct-to-home video visits as a good option when their child’s condition is static, and
when travel isn’t possible due to weather, childcare or work. Effective direct-to-home telemedicine visits are possible when parents are engaged and capable of positioning the child. Using pre-made cards or a doll to show positions to parents helped to improve the quality of visits.

**Conclusions** Despite inherent limitations, direct-to-home telemedicine visits could benefit children when used appropriately in conjunction with in-person visits. Our findings suggest that there are specific practices that facilitate effective direct-to-home telemedicine visits.

**Purpose of Study** This study was part of an ongoing effort at our institution to identify characteristics in our patient population associated with long term (defined as visit between 4–12 weeks) postpartum visit (PPV) attendance and specifically examines whether initiation of telemedicine during the COVID-19 pandemic was a predictor of PPV compliance.

**Methods** A retrospective chart review was performed of patients who delivered between April 10, 2019 - August 31, 2019 and during the same time period in 2020. The main outcome of interest was percentage of deliveries with PPV follow-up in 2020 compared to 2019. Demographics including age, race/ethnicity, parity, total prenatal visits, delivery route, gestational age, birthweight, and length of newborn hospital stay were also reviewed. Statistical analysis was performed using two sample T-test or Mann-Whitney/Wilcoxon Two-Sample Test, or Chi-Squared test as appropriate. Logistic regression was used to identify independent predictors of PPV.

**Summary of Results** In total, 587 patients were identified in 2019 and 552 in 2020. 13 duplicate charts were removed for a total of 1126 charts in the final analysis. The 2 cohorts were determined to not be significantly different except in terms of parity which was higher in 2019. In 2020, 79% of patients were compliant with PPV, in 2019 71% were compliant. This difference was determined to be statistically significant with p-value = 0.001. 2019 PPV were conducted in-office only while 2020 PPV were offered as telemedicine or in-person. Approximately 53% of visits in 2020 were telemedicine.

Other variables predictive of postpartum follow-up included higher age, increased parity, greater total prenatal visits. Cesarean delivery was positively associated with follow-up but it was not significant. Gestational age, race/ethnicity, birthweight, and newborn length of stay were not identified as predictive of follow-up. In the logistic regression delivery year, age, prenatal visits, and parity remained significant and independent predictors of follow-up.

**Conclusions** This study reviewed 2 cohorts of patients during time periods before and after initiation of telemedicine postpartum visits during the COVID-19 pandemic. Previous research at our institution indicated that telemedicine follow-up was non-inferior to office follow-up for short term visits in patients with hypertensive disorders. This study expands upon this work by considering long term follow-up in all postpartum patients. Additionally, while demographic characteristics predictive of postpartum attendance have been considered elsewhere in the literature there are fewer studies examining access to telemedicine as a variable. These findings suggest telemedicine may be effective for increasing compliance with postpartum visits.

### Hematology and Oncology II

**Concurrent session**

**10:15 AM**

**Friday, January 21, 2022**

**#304 COMPREHENSIVE CARE MODEL FOR SICKLE CELL DISEASE CARE AT KERN MEDICAL**

*R Sharma*, L Moosavi, E Cabos. Kern Medical Center, Bakersfield, CA

**Purpose of Study** Sickle cell disease is the most common genetic hemoglobinopathy in the US. It affects over 100,000 individuals. Sickle cell disease is a blood disorder causing a mutation in the hemoglobin beta chain. It is characterized by acute pain episodes, emergency department visits, hospitalizations, anemia and early mortality. This disease requires a programmatic, multidisciplinary and team-based approach. Currently there are three potential models of care for sickle cell disease patients: classic comprehensive model, embedded care model and specialized medical home model. The model adapted at our institution is a blend of all the three potential models incorporating as per our institutional considerations. This study presents the results following the multidisciplinary care implementation at our institution over the last year.

**Methods** This study was approved by the Institutional Review Board of Kern Medical. A retrospective review of post initiation of a multidisciplinary care intervention at a public academic medical center was reviewed.

**Summary of Results** This intervention was implemented on July 13th 2020. The multidisciplinary team includes hematologist/oncologist, pharmacists, primary care physician, pain specialist, sickle cell specialist, behavior health worker, clinical health worker, medical assistant and registered nurse. In the last six months a total of 170 patients were scheduled to be seen at the sickle cell clinic. 14 patients were admitted inpatient in the last six months which is a decrease in number compared to the data prior to implementation. 23 additional patients were newly referred from outside institutions. Our findings thus far highlight the importance of gaining patients’ trust, providing social support and providing patients with appropriate resources. In the last month we have also established a virtual reality program for pain management for patients. In December of 2020 at the state level, we also gained recognition for standardizing care for sickle cell patients.

**Conclusions** In conclusion, a multidisciplinary comprehensive care for sickle cell patients is a necessity. At present there are only ten sickle cell clinics in the state of California. Our academic center became one of the founding institutions of the upcoming Networking California for Sickle Cell Model. The model of care for implementation of this disease varies highly upon location, population served, financial and institutional needs.
Purpose of Study
Approximately 20% of newly diagnosed cancer patients are between the typical parenting ages of 20 and 54, and so many of these patients are also the primary caregivers of children. Qualitative studies focusing on this demographic indicate that patients who are parents struggle to balance their own care needs with those of their children, but to date, no research has formally evaluated the need for on-site childcare services at cancer centers. This study aims to explore the need for childcare support for cancer patients from the perspective of healthcare professionals providing care at a major Canadian cancer center.

Methods
Used
Between May and April 2021, healthcare providers at one major Canadian cancer center were invited to partake in semi-structured interviews. The interview guide was developed through consultation with a multidisciplinary team and aimed to elicit the perspectives of healthcare providers on the importance and potential benefits of childcare services for their patients. Specific questions were also designed to explore what might constitute optimal childcare solutions for patients, and how cancer centers and healthcare providers could help to address this need. Interview transcript data was interpreted using thematic analysis.

Summary of Results
Semi-structured telephone interviews were conducted with 28 healthcare professionals providing care at a major Canadian cancer center between April and May 2021, including medical, surgical, and radiation oncologists, psychiatrists, general practitioners, registered nurses, radiation therapists, and social workers. Managing childcare responsibilities was described as challenging for patients, and the introduction of childcare services for patients on-site at cancer centers was seen as a way to reduce emotional and financial stress. Other identified benefits of introducing childcare support services for patients included increased system efficiency, improved treatment compliance, increased trust in providers and cancer centers, and additional emotional support for the children of patients.

Conclusions
These results indicate that childcare issues are broadly impactful for parents battling cancer, and that providing childcare support for these patients could be highly beneficial from both a medical and social perspective. As such, these findings suggest that cancer centers could consider the implementation of childcare support services to provide a more patient-centered approach to care.

#306 OUTCOME OF PLANNED CHECKPOINT INHIBITOR DISCONTINUATION IN METASTATIC MELANOMA PATIENTS IN A COMMUNITY ONCOLOGY PRACTICE: A CASE SERIES

L Perez*, 1W Samlowski, 1RD Lopez-Flores, 1University of Nevada Reno School of Medicine, Reno, NV; 2University of Nevada Las Vegas, Las Vegas, NV

Purpose of Study
Check-point inhibitor (CKI) therapy for melanoma has dramatically changed the prognosis for patients with metastatic melanoma. Twenty to 40% of patients appear to achieve long-term remissions. It is not clear whether patients require ongoing therapy to maintain remissions or whether treatment can be safely discontinued. We evaluated a potential strategy for elective treatment discontinuation.

Methods
Used
A retrospective chart review of patients with metastatic melanoma treated with CKI based treatment was performed. Patients who underwent elective treatment discontinuation after 2 negative scans 3 months apart or had biopsies confirming complete remission (CR) were identified for analysis.

Summary of Results
Of 139 checkpoint inhibitor treated patients, 53 individuals achieved a CR. The progression free survival was 100% at 1 year, 95% at 3 years after treatment discontinuation, with a median duration of follow up of 31 months. Four of 53 individuals with complete remissions (7.5%) eventually relapsed. The median overall survival of the entire cohort was not reached and was 100% at 1 year, and was 95% at 3 years.

Conclusions
In this retrospective analysis, we demonstrate an elective treatment discontinuation strategy that is generalizable to a variety of CKI ± targeted therapy regimens. We have found a high complete response duration after elective treatment discontinuation without a requirement for a specific period of therapy. We hypothesize that this approach can potentially decrease treatment related financial burden and toxicities faced by responding patients with improved quality of life.
Models were trained to predict the ordinal NMRI scores using a regression objective function, to take advantage of the ordinal scores. To assess the models as a clinical screening tool, performance was evaluated as a binary classification task where predicted scores were mapped to low- and high-severity labels. Regression training improved performance compared with binary classification training.

The reports were split into train (70%), validation (15%), and test (15%) sets. Model hyperparameters were tuned using the training and validation sets to maximize area under the receiver operating characteristic curve (AUC), and final performance was evaluated on the withheld test set.

Summary of Results Results from each approach are reported in table 1. Although DistillBERT is a more sophisticated architecture, both models performed similarly.

While neither model completely generalized to the test set, both are shown to have predictive value. Since NMRI scores were annotated by many individuals over 9 years, scores may vary between raters and over time, limiting model performance.

Conclusions This work presents two candidate architectures for a severity screening tool for incident reports in radiation oncology. We show AUC scores up to 0.77, demonstrating that these architectures have the potential to identify safety-critical reports and reduce latency of interventions. Future work includes incorporating radiation oncology-specific text and experimenting with BERT variants that are pretrained on clinical text.

### Abstract #307 Table 1 Performance metrics on test set

<table>
<thead>
<tr>
<th></th>
<th>AUC</th>
<th>F1</th>
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<tr>
<td>LSTM-A</td>
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<tr>
<td>DistillBERT</td>
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**#308 INVESTIGATING BARRIERS TO RADIATION ONCOLOGY TREATMENT FOR AMERICAN INDIAN AND ALASKA NATIVE PATIENTS**

D Marashi*, M Greer, L Moriguchi Halasz. University of Washington School of Medicine, Seattle, WA

10.1136/jim-2022-WRMC.304

**Purpose of Study** American Indian/Alaska Native (AI/AN) citizens comprise 2% of the population, both in the United States and Washington State. Among this population, cancer is the leading cause of death for females and the second for males. Radiation therapy (RT) is recommended for about 50% of cancer patients. Unfortunately, few data are available regarding the barriers to RT treatment for AI/AN populations. Our objective is to characterize the barriers AI/AN patients experience while accessing cancer care at the University of Washington Medical Center, and their affiliated hospitals, and to determine which barriers (if any) are captured and addressed in the electronic health record.

**Methods Used** Levesque et al. describe a ‘5A’ theoretical framework – approachability, acceptability, availability, affordability, and appropriateness – that considers supply-side factors (e.g., health systems, institutions, organizations, providers) alongside demand-side factors (e.g., individual, household, community, population), as a framework to describe the complex process patients experience utilizing healthcare. We conducted an analysis based on this framework using 36 AI/AN patient charts, focusing on their radiation oncology and social work notes from the last five years. All data were double coded. Quotes describing each barrier were used to support the coding process, and these were referred to to resolve any discrepancies between coders.

**Summary of Results** When looking specifically at the supply-side dimensions of access to oncologic services, availability/ accommodation (25.0%) and affordability (16.6%) appeared as the most common barriers to care. In contrast, a very small percentage of patient charts mentioned acceptability as a barrier to care (1.0%). Most chart descriptions did not feature demand-side abilities of patients or populations to access services (5.0%). Illustrative vignettes were also organized to represent the different types of barriers discovered in patient charts.

**Conclusions** AI/AN communities find it challenging to access health services (either the physical space or healthcare providers) in a timely manner, and potentially lack the economic resources to pay for healthcare services without catastrophic expenditures. Our review also uncovered the supply-side dominance in terms of dimensions of health care access. In other words, despite evidence suggesting individual-level interventions are more effective at improving access for patients, there was still an emphasis on services provided rather than self-management and health literacy approaches. It is not known whether this reflects that providers do not address demand-side dimensions of access or simply do not record it in the chart. Regardless, the findings suggest efforts to address all dimensions of access may be important to diminishing health care disparities.

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**#309 CONCURRENCE OF COCCIDIOIDOMYCOSIS AND MALIGNANCY**

J Slaton, R Sharma*, M Mistry, N Dhillon, A Heidari, R Johnson, R Kuran, R Polineni, E Cobos. Kern Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.305

**Purpose of Study** Coccidioidomycosis (CM) is endemic to Southwestern USA with over 150000 cases per year. Concurrency of CM and malignancy has been reported sporadically but the interaction and association is yet to be studied. This study is designed to identify and describe cases with CM and malignancy from a single center located in highly endemic area of Coccidioides in California.

**Methods Used** A retrospective chart review following IRB approval at our institution was conducted. The inclusion criteria was patients greater than or equal to 18 years of age, positive CM serology and positive diagnosis of malignancy on biopsy. Pediatric age group, inmates, pregnant females, human immunodeficiency virus patients and patients with incomplete medical records were excluded from this study.

**Summary of Results** Twelve total patients were included in this study. Characteristics of cancer by location of CM are displayed in table 1. 5/12 patients presented with associated diabetes mellitus. 11/12 patients underwent therapeutic treatment with fluconazole and one patient didn’t receive antifungal therapy. 11 patients underwent surgical resection for cancer, one patient had zero-reactivation of CM post fluconazole.
failure and 1 patient succumbed to life. Common cancer types noted with concurrence CM infection in this study are colorectal, prostate, gynecologic, renal cell, gastric and skin. Onset of cancer prior to CM infection and vice versa was noted in almost equal number of patients. Only one case demonstrated simultaneous onset of cancer and CM. Extrapulmonary dissemination of CM was noted only among renal cell carcinoma and melanoma patients.

Conclusions Coexistence of coccidioidomycosis and cancer is not common but when it occurs, the interaction between the two conditions is not well understood and management is formidable. A close collaboration between oncology and infectious diseases teams is paramount.

### #310

**OUTCOME OF PLANNED CHECKPOINT INHIBITOR DISCONTINUATION IN METASTATIC NON-MELANOMA SOLID TUMOR PATIENTS IN A COMMUNITY ONCOLOGY PRACTICE: A CASE SERIES**

1.64% of patients with other tumor types can achieve durable remissions. The duration of treatment or whether treatment can be safely stopped to maintain a complete response is controversial. Based on melanoma-derived data, we tested whether CKI treatment could be safely discontinued in patients with other solid tumors.

Methods Used A retrospective analysis was performed in adults with metastatic non-melanoma solid tumors treated with CKI-based therapy. Patients with solid tumors who achieved a complete remission on 2 sequential scans at least 3 months apart were identified from our computerized patient database. Patient data was analyzed for patient characteristics, as well as progression-free, and overall survival.

Summary of Results Sixty-nine non-melanoma solid tumor patients were treated with CKI-based regimens in our clinic and 15 achieved a complete remission (21.7%). Five patients were female (35.7%) and the remaining nine were male (64.3%). A 100% PFS was reported for these patients who achieved CR with over 20 months of median follow up after elective treatment discontinuation. Median OS was not reached. CKI toxicity primarily included acute toxicity with rare, chronic unresolved toxicity over 6-month duration in two patients.

Conclusions In this retrospective case series, all patients remained in complete remission after treatment discontinuation. We hypothesized that appropriate selection of patients for early treatment discontinuation may decrease their economic burden related to ongoing treatment, limit potential toxicity, and improve quality of life.

### #311

**NOVEL FLOW CYTOMETRY ANTIGENS TO DIFFERENTIATE FOLLICULAR LYMPHOMA AND DIFFUSE LARGE B-CELL LYMPHOMA**

L. Schwarz*, I. Fromm. University of Washington School of Medicine, Seattle, WA

10.1136/jim-2022-WRMC.307

Purpose of Study Flow cytometry can aid in the identification of blood cancers, including Non-Hodgkin's Lymphomas. Its ability to quantify the antigen expression of cells is often used to differentiate between similar neoplasms. However, there is currently a dearth of antigens which differentiate Diffuse Large B-cell Lymphoma (DLBCL) and Follicular Lymphoma (FL). We evaluated the ability of novel antigens including CD32, CD38, CD40, CD44, CD71, and CD75 to discriminate between the two lymphomas.

Methods Used Tissue samples had been previously collected, immunohistochemically and morphologically categorized, and cryopreserved in long-term liquid N2 storage. DLBCL and FL samples were identified using an internal database, thawed, stained with a 10-color antigen panel: CD40/CD32/CD3/CD44/CD38/CD75/CD71/CD19/CD5 and run on a BD LSR II Flow Cytometer. Benign follicular hyperplasia (FH) samples were also processed and run as a negative control. Populations were isolated and analyzed using WoodList, an internal software program. Median fluorescence intensity (MFI) for each antigen, forward scatter (FSC), and side scatter (SSC) were collected for neoplastic cells in DLBCL and FL samples and for all B-cells in FH. Values were exported to Excel and used to create two-tailed boxplots using the maximum, minimum, median, and first (Q1) and third (Q3) quartile values. Antigens were considered ‘successful’ in differentiating the two lymphomas if there was no overlap between Q3 of one neoplasm and Q1 of another.

Summary of Results While there were apparent differences between FL and DLBCL (both germinal and non-germinal center) in the expression levels of CD32, CD38, CD40, CD44, and CD75, there was significant overlap in the plots for each of these antigens. For CD71 however, FL was adequately differentiated from both germinal and non-germinal center DLBCL. Both forward scatter and side scatter differentiated low-grade but not intermediate-grade FL from DLBCL. Incidentally, CD44 and CD32 showed the ability to separate non-germinal center from germinal center DLBCL. However, that data is limited by the low sample size of non-germinal DLBCL (n=5).
Conclusions CD71 was the sole antigen which successfully discriminated FL from both germinal and non-germinal center DLBCL in isolation. However, combining parameters, such as FSC and CD71, may better separate these two lymphomas. Further research with a more robust sample size is needed to confirm the ability of CD44 and CD32 to differentiate between germinal and non-germinal center DLBCL.

Abstracts

FREQUENCY OF ACUTE LEUKEMIA IN PEDIATRIC PATIENTS WITH BELL PALSY

#312

1CM Tanji*, 2,3LG Yamamoto. 1Hawaii Pacific Health, Honolulu, HI; 2University of Hawai‘i at Manoa, Honolulu, HI; 3Kapi‘olani Medical Center for Women and Children, Honolulu, HI

10.1136/jim-2022-WRMC.308

Purpose of Study A recent study identified a 0.6% rate of acute leukemia in children with facial palsy. In response, the primary objective of our study was to confirm if there is a relationship between Bell Palsy and acute leukemia in children younger than 19. If found, obtaining a Complete Blood Count (CBC) screening upon Bell Palsy diagnosis would be in order and reconsideration of corticosteroid treatment, which has been shown to interfere with leukemia diagnosis, if the CBC is suspicious. If not found, this would cast serious doubt on whether a true relationship exists.

Methods Used A retrospective chart review was conducted (n=513) of patients < 19 years old with Bell Palsy between April 7, 1995 to June 4, 2021. Data was collected using the hospital billing system and an electronic medical record from 4 community hospitals in Hawai‘i. Statistical analyses were conducted to describe the patient demographic.

Summary of Results 53% of the patients with Bell Palsy were female and the remaining 47% were males, with around 67% of patients in the 6–12 and 13–19 years age groups. Of the 513 pediatric patients with Bell Palsy, one patient developed leukemia, leading to an association of 0.19%. However, the patient’s leukemia diagnosis was 3.5 years after their initial Bell Palsy encounter, suggesting these two events might not be medically related.

Conclusions Our data is unable to confirm an association between Bell Palsy and acute leukemia.

Abstract #312 Figure 1 Among the 513 patients, only one patient with Bell Palsy was diagnosed with acute leukemia.

Infectious diseases II
Concurrent session
10:15 AM
Friday, January 21, 2022

#313

3-AMINOIMIDAZO[1,2-A] PYRIDINE DERIVATIVES WITH ANTIBACTERIAL ACTIVITY AGAINST STAPHYLOCOCCUS AUREUS

1L Johari*, 2E Bentley, 3D Vosburg, 4H Szurmant. 1Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA; 2Harvey Mudd College, Claremont, CA

10.1136/jim-2022-WRMC.309

Purpose of Study According to the World Health Organization (WHO), antimicrobial resistance is a serious public health issue on a global scale. Nationally, the US Centers for Disease Control and Prevention (CDC) estimate that 3 million infections are caused by antibiotic-resistant bacteria, resulting in approximately 35,900 deaths annually. Antimicrobial resistances occur when bacteria are forced to evolve due to widespread exposure to antimicrobial drugs. One way to address antimicrobial resistance is to identify novel antimicrobial compounds that can target essential microbial cellular processes. Among a diverse library of novel heterocycles produced in one of our laboratories, we previously identified one, a 3-Aminoimidazo [1,2-a]pyridine with moderate antimicrobial activity against methicillin resistant Staphylococcus aureus (MRSA). To identify the substituents of the compound necessary for killing activity and to potentially improve the initial hit compound, a medicinal chemistry approach was pursued.

Methods Used 3-aminoimidazo [1,2-a]-heterocycles were synthesized using a green, microwave-assisted Groehke-Blackburn-Bienaymé multicomponent reaction. Compounds were screened for their ability to inhibit growth of MRSA. For screening purposes, we utilized the Kirby-Bauer disk diffusion assay, according to Clinical Laboratory Standards Institute protocols. To this end, bacterial suspensions with a density equivalent to a 0.5 McFarland standard were evenly spread over fresh agar plates. Paper disks containing 100 µg of the respective compounds, vancomycin as a positive control, or no compound as a negative control were prepared. Compound and control disks were evenly distributed on the agar plates and plates were incubated for 12h at 37°C to allow growth of the bacteria. In vitro testing showed potent activity against a variety of Gram-positive and Gram-negative bacteria. However, there were no clear dose-dependent effects by the absorbance reader.

Summary of Results A series of 3-aminoimidazo[1,2-a]-heterocycles were synthesized and subjected to antimicrobial screening. In addition to the previously identified active compound, a second compound was found to exhibit similar moderate antimicrobial activity against MRSA. The active compounds share 4-chlorophenyl and tert-butyl substituents and varied core heterocyclic structures. Equipped with this knowledge, further medicinal chemistry efforts are underway to improve the activity of these compounds.

Conclusions Compounds similar to the two candidates with antimicrobial activities have also been identified to have anti-inflammatory, anticancer, and anti-fibrosis effects. Thus 3-Ami-noimidazo[1,2-a]pyridines and related structures have
demonstrated a wide range of biological activities, including antimicrobial effects. Further efforts are underway to select for improved killing activity and move towards cytotoxicity studies.

**#314** THE ANTIBIOTIC SPECTRUM OF NOVEL METHIONYL-TRNA SYNTHETASE INHIBITORS

1CE Liston*, 1N Molasky, 1A Mushkaq, 1JR Gillespie, 1F Buckner, 1University of Washington, Seattle, WA; 2University of Washington School of Medicine, Seattle, WA

Purpose of Study As antibiotic resistance continues to increase worldwide, patient care and outcomes will decline. Therefore, it is necessary to investigate alternative biochemical pathways in bacterial lifecycles and select new drug targets for novel inhibitory compounds. One target of interest is the Methionyl-tRNA Synthetase (MetRS) enzyme which is critical in protein synthesis. Typically, Gram-positive bacteria express type-1 MetRS and Gram-negative express type-2 MetRS. The purpose of this study is to explore the efficacy of novel MetRS inhibitors on type-1 MetRS enzymes compared to control antibiotics in Gram-positive and Gram-negative bacteria.

Methods Used All preparations for bacterial growth and cultivation followed the standardized guidelines outlined by the Clinical Laboratory Standards Institute. Minimum Inhibitory Concentration (MIC) assays were performed in triplicate and repeated at least twice. The selected Gram-positive strains were non-anthracs *Bacillus ssp.*, *Listeria monocytogenes*, *Staphylococcus epidermidis*, and the Gram-negative bacterial strains were *Serratia marcescens*, *Pseudomonas aeruginosa*, and *Burkholderia cepacia*. Each strain was streaked from frozen stock to strain-appropriate agar and incubated at 37°C for 24 hours. Single colony inocula were collected from the plate and cultured to the exponential growth phase in strain-specific broth. The culture was adjusted to MacFarland 0.5 standard and quantified through the absorbance reader (OD 600). The microbroth dilution was completed in 96-well plates in a 2-fold dilution using novel MetRS inhibitory compounds ChemIDs: 2541, 1986, 1717, 2536, 2144, 2093, 2596 and control antibiotics. The MacFarland adjusted bacteria were added to the plates. Negative and positive controls were established in each plate and after an additional 18–24 hours of incubation at 37°C, the MIC90 was calculated from the data generated by the absorbance reader.

Summary of Results The novel MetRS inhibitors demonstrated variable yet potent inhibitory effect against type-1 MetRS enzymes in the Gram-positive strains. The MICs ranged from <0.0312–0.25 μg/mL. In most cases these MICs were more efficacious than the control antibiotics. As predicted, the compounds showed little activity in the type-2 MetRS enzymes in Gram-negative bacteria.

Conclusions In vitro testing showed potent activity against a variety of Gram-positive bacteria species. Parallel studies not reported here demonstrated in vivo activity in mouse models of MRSA infection. The compounds have excellent potential for development as selective agents for treating Gram-positive infections.
#316  CHARACTERISTICS AND OUTCOMES OF HOSPITALIZED COVID-19 PATIENTS AND DNR

1JY Hwang, C Lee, T Nguyen, T Buck, K Mun, T Vanteru, D Baral, D Tirschwell, A Kim. Washington State University Elson S Floyd College of Medicine, Spokane, WA; University of Washington School of Medicine, Seattle, WA

Purpose of Study Do-not-resuscitate (DNR) orders allow patients, per their wishes, to decline cardiopulmonary resuscitation (CPR) in a cardiac or respiratory arrest event. DNR status has been linked with severe illness, advanced age, poor disease prognosis, and deteriorating health status with impending death. Conversely, DNR status has not been associated with a reduction in priority of care in the intensive care units. In the current COVID-19 pandemic, patients and families are often faced with a need to make a DNR decision with limited information. Some hospital systems had to impose a unilateral DNR policy, resulting in unprecedented, unresolved ethical dilemmas. This study further investigates the characteristics and outcomes of hospitalized COVID-19 patients in association with DNR status.

Methods Used An observational study of hospitalized patients with COVID-19 was conducted in the greater Seattle area from January 2020 to December 2020. Data was collected through review of the electronic medical record as part of the national American Heart Association COVID-19 registry. Inclusion criteria for DNR included patients with any documentation of a DNR order, partial DNR order, or advanced directive related to restricting attempts to resuscitate prior to or after arrival to the hospital. Physician Orders for Life-Sustaining Treatment form, Do Not Attempt Resuscitation form, and the Medical Orders for Scope of Treatment form were all considered as documentation of DNR. We performed a chi-square test of independence and one-way ANOVA analysis.

Summary of Results Of the total 396 hospitalized COVID-19 patients in our registry, DNR status was available for 368 patients. Of the 368 patients, 114 (31%) were deemed to be DNR. The average age of patients who were deemed DNR was 72 years, while the average age for those who were not DNR was 55 years (F(1,366) = 73.37, p < .001). Patients with comorbidities were more likely to be DNR (34%) compared to patients with no past medical history (18%) (χ² = 5.802, df = 1, p = 0.019). With regard to mortality, 69 out of the 114 patients with a DNR order expired (61%), while 11 out of the 254 patients without a DNR order expired (4%) (χ² = 142.768, df = 1, p < .001). Of the total 80 expired patients, 86% were DNR. There was no statistically significant difference in the average length of hospitalization between the two groups (F(1,366) = 1.44, p = 0.231).

Conclusions Our data suggest that DNR is associated with increased mortality, age, and comorbidities. However, there were no statistically significant associations observed between sex and length of hospitalization. These results potentially reflect the overwhelming burden our healthcare system has faced during the COVID-19 pandemic when resources have been substantially limited.
community educational program will help with the complexities of understanding vaccines. This study aims to minimize vaccine hesitancy, increase the uptake of COVID-19 vaccines, and promote herd immunity.

**Methods** A COVID-19 vaccine educational program was created for a diverse audience to enhance the knowledge in underserved communities on the benefits of vaccines to prevent diseases. The training module materials presented in the program provided information on the background and significance of vaccines to the COVID-19 pandemic. Following the training, the participants completed demographics, test review questions, and a one-page survey feedback questionnaire. SPSS statistical software was used for descriptive One-Way ANOVA analysis for the data calculations.

**Summary of Results** After examining the results from 103 participants, an exemplary average score of 98.4% was observed for the review section. The mean average response on the Likert-scale survey questions was ‘Strongly Agree’ for comprehending new information about COVID-19 vaccines. One-Way ANOVA produced statistically significant p values ≤ 0.05 by SPSS. Additionally, 84 participants agreed to educate 10 additional individuals about the benefits of vaccines, expanding the population of the study to be inclusive of 840 participants.

**Conclusions** The community engagement educational training program demonstrated a decrease in vaccine hesitancy and promoted the uptake of COVID-19 vaccines in underserved communities, fundamentally assisting the elimination and prevention of the COVID-19 pandemic by contributing to herd immunity.

**#319 EFFECTS OF GEOGRAPHY, SOCIODEMOGRAPHICS AND POLITICAL LEANING ON COVID-19 CASE-FATALITY RATE IN WEST COAST STATES**

JS Gerken*, D Xuivinen, D Zapata, I Zapata. Rocky Vista University College of Osteopathic Medicine, Parker, CO

Purpose of Study The COVID-19 pandemic, caused by the SARS-Cov-2 virus, has impacted nearly every aspect of scientific research. Many factors such as comorbidities, sociodemographic descriptors along with political associations have been associated to fatality rates. One specific factor that has continuously suspected to have a potential impact on COVID-19 fatality rates is Vitamin D deficiency. Few studies have evaluated the effect of environmental, geographic, or sociodemographic variables together on COVID-19 case-fatality rate. Our study evaluates COVID-19 case-fatality rate on a per-county basis of the Pacific Coast states using comorbidity, sociodemographic and political leaning data in conjunction with geographic variables such as average precipitation, air pollution, average temperature, latitude and elevation which have been shown to correlate to Vitamin D deficiency.

**Methods** This analysis evaluates the effect sizes of each particular factor described using Generalized Additive Models (GAM). This type of model was selected to accommodate for non-parametric evaluations of continuous predictors. Estimates obtained through these models then are standardized and sorted by their effect size and direction to allow for fair interpretation with a standardized context.

**Summary of Results** Our analysis showed that geographic and environmental variables associated with Vitamin D levels did not affect COVID-19 case-fatality rates; however, factors such as Medicare flu vaccination rate, voting Democrat during the 2020 Presidential Election, and being uninsured were shown to be the most important factors in reducing COVID-19 case fatality rate. Comorbidities, disproportionate higher females in a county, mental health, and voting Republican in the 2020 Presidential Election significantly increased COVID-19 case fatality rates.

**Conclusions** The effect of Vitamin D deficiency, indirectly measured through geographic variables, was not supported by our results. However, there were other clinically significant factors such as comorbidities, socioeconomic, and political leaning factors that influenced COVID-19 case fatality rates.

This work was presented in part at Rocky Vista University Appreciation Day (virtual) on October 15, 2021

**#320 THERAPEUTIC ANTICOAGULATION IN HOSPITALIZED PATIENTS WITH COVID-19**

1C Lee*, 1T Nguyen, 1JY Hwang, 2T Buck, 2K Mun, 2T Vanteru, 2D Baral, 2D Tirschwell, 1A Kim, 1Washington State University Elson S Floyd College of Medicine, Spokane, WA; 2University of Washington School of Medicine, Seattle, WA

Purpose of Study Throughout the COVID-19 pandemic, providers observed increased rates of thrombus and inflammation among COVID-19 patients which affected multiple organs and led to complications such as lung failure, myocardial infarction, and stroke. Furthermore, studies have shown increased D-dimer levels to be associated with worse outcomes in COVID-19 patients. A spectrum of anticoagulants is currently being used in COVID-19 patient management as needed to combat this hypercoagulable state. Current treatment guidelines and literature investigating the effects of different anticoagulants in COVID-19 patients are continuing to evolve. Here, we describe a preliminary study that investigated the effects of therapeutic anticoagulation on the outcomes of hospitalized COVID-19 patients.

**Methods** From a national registry developed by the American Heart Association, we performed a retrospective observational cohort study of hospitalized COVID-19 patients who were admitted to one of four hospitals in the Greater Seattle area from January 2020 to December 2020. Diagnosis of COVID-19 was made either clinically using hospital specific criteria, or by polymerase chain reaction testing. Medications given for therapeutic anticoagulation include parenteral unfractionated heparin, subcutaneous low molecular weight heparin full therapeutic dose, argatroban, bivalirudin, direct oral anticoagulant, and warfarin. Mortality was quantified by the number of patients who died during hospitalization or were discharged to hospice care.

**Summary of Results** Out of 396 hospitalized COVID-19 patients, 57 patients received one or more anticoagulants for therapeutic use. For 89% of those patients (51 out of 57), therapeutic anticoagulation was indicated due to pre-hospitalization anticoagulant use or development of myocardial infarction, pulmonary embolism, or deep vein thrombosis during hospitalization. Of those who received therapeutic anticoagulation, the survival rate was 67% (38 out of 57). On the contrary, out of the 339 patients who did not receive any type of therapeutic anticoagulation, the survival rate was 79% (267 out of 339).
out of 339). This difference of higher survival rate and lower mortality rate in patients who did not receive therapeutic anticoagulation was not statistically significant (p=0.066).

Conclusions Our preliminary findings indicate that therapeutic anticoagulation does not affect mortality in hospitalized COVID-19 patients. However, therapeutic anticoagulation may be necessary in high-risk patients who are more susceptible to a COVID-19 induced hypercoagulable state to reduce complications or death resulting from thromboembolic events. Further investigation is necessary with additional analyses performed to control for confounding by indication and other prognostic factors.

**Abstract #321**

**SHORT-COURSE ANTIBIOTICS FOR TREATMENT OF UNCOMPLICATED COMMUNITY ACQUIRED PNEUMONIA IN PEDIATRIC PATIENTS**

1YN Manely*, 1M Arunachalam, 1T Hoang, 1E Kang, 1H Lee, 1J Parikh, 1C Palomino, 1B Afghani. 1UC Irvine School of Medicine, Irvine, CA; 2Children’s Hospital of Orange County, Orange, CA

10.1136/jim-2022-WRMC.317

**Purpose of Study** Several studies of adults have shown that a 5-day course of antibiotics is as effective as a 10-day course in treatment of community acquired pneumonia (CAP). However, the effectiveness of short-course therapy in pediatric patients with CAP has not been well documented. The main goal of this study was to investigate the literature for studies that compare the effectiveness of short-course vs long course antibiotics in treatment of pediatric CAP.

**Methods Used** We conducted a literature search using search engines such as Google Scholar, SciHub, and PubMed. Key words used included: 'pediatric' 'children' 'pneumonia' 'short-course' 'long-course' and 'treatment'. Only studies of pediatric patients with uncomplicated community acquired pneumonia (CAP) that included a control group and used high-dose amoxicillin as part of the treatment were included in our review.

**Summary of Results** We found 4 studies that satisfied our inclusion criteria. All four studies excluded patients diagnosed with severe or complicated pneumonia and those with underlying conditions. Please see table 1 for summary of the results. The age of children in the studies ranged from 2 months to 18 years. The investigators measured cure rates and/or relapse rates during the follow-up period of 14 to 30 days. The studies showed that a short-course of 5–7 days was not inferior to a 10–14 days of treatment. The design was different in various studies as 3 of the studies required chest radiograph as well as clinical evidence of pneumonia while one study was based on clinical diagnosis of pneumonia.

**Conclusions** The results from our literature review show that a 5–7 day antibiotic course is non-inferior to a longer course of 10 to 14 days in treatment of previously healthy children with uncomplicated community acquired pneumonia. Larger studies are warranted to determine whether these results are reproducible across different populations or non-trail conditions.

**Morphogenesis and malformations**

**Concurrent session**

**10:15 AM**

**Friday, January 21, 2022**

**Abstract #322**

**SIN3A IS REQUIRED FOR EPIGENETIC REGULATION OF DIAPHRAGM AND LUNG DEVELOPMENT**

G Stokes, D McCulley*. UCSD, University of California San Diego, La Jolla, CA, La Jolla, CA

10.1136/jim-2022-WRMC.318

**Purpose of Study** One of the most common and severe congenital malformations is congenital diaphragmatic hernia (CDH) which occurs in 1 out of 3500 live births with a mortality rate of 20–50%. Although abnormal diaphragm formation is the hallmark of the disease, the high mortality rate is due to abnormal lung and pulmonary vascular development causing lung hypoplasia and pulmonary hypertension. Despite the frequency and severity of CDH, little is known about the underlying developmental and genetic mechanisms responsible for the disease. Our hypothesis is that a core group of genes are responsible for failure of diaphragm formation as well as defects in lung and pulmonary vascular development.

**Conclusions** These data demonstrate the importance of genetic mechanisms responsible for abnormal lung and pulmonary vascular development.
patients. To investigate this hypothesis, we are studying the role that genetic mutations identified in patients with CDH play in diaphragm, lung, and pulmonary vascular development. In this study we investigated the role played by SIN3A, a gene containing pathogenic sequence variants identified by whole genome sequencing in patients with CDH, in diaphragm, lung, and pulmonary vascular development.

Methods Used To investigate the role of SIN3A, we used tissue and cell-specific deletion in diaphragm and lung mesenchymal progenitor cells in mice. The diaphragm and lung phenotypes associated with loss of SIN3A were characterized using whole mount, brightfield, and immunofluorescence imaging. Pulmonary vascular defects associated with loss of SIN3A were characterized by fluorescence imaging as well as echocardiography and right heart catheterization. The genetic mechanisms responsible for abnormal lung and pulmonary vascular development were studied using gene expression analysis with RNA collected from whole lung and sorted lung mesenchymal cells.

Summary of Results Loss of SIN3A in somatic mesoderm derived skeletal muscle and lateral plate mesoderm derived mesothelium resulted in abnormal diaphragm development and CDH in mice. In the developing lungs, loss of SIN3A resulted in failure of alveologenesis and pulmonary hypertension. Lung defects in SIN3A deficient mice were evident at late embryonic stages with failure of mesenchymal cell differentiation, decreased cell cycling, and increased DNA damage. Loss of SIN3A was associated with an imbalance of histone acetylation that was restored by embryonic inhibition of histone acetyltransferase.

Conclusions These data demonstrate the importance of genetic evaluation in structural malformations such as CDH. In the case of SIN3A loss of function, impaired epigenetic regulation of gene expression in the diaphragm and lungs resulted in failure of diaphragm formation as well as lung and pulmonary vascular defects common among patients with CDH. These defects were reversed by restoring the balance of histone acetylation.

Abstract #324 BIALLELIC VARIANTS IN GPX4 ARE ASSOCIATED WITH A SPECTRUM OF SKELETAL ANOMALIES, HYPOTONIA, AND NEURODEGENERATION AND MAY INCREASE SUSCEPTIBILITY TO FERROPTOSIS

1K. Wigby*, 3H. Saneto, 1,2J. Friedman, 1,4H. Liu, 5B. Stockwell, 1University of California San Diego, La Jolla, CA; 2Rady Children’s Institute for Genomic Medicine, San Diego, CA; 3Seattle Children’s Hospital, Seattle, WA; 4University of Washington, Seattle, WA; 5Columbia University, New York, NY

10.1136/jim-2022-WRMC.320

Case Report Biallelic loss of function variants in GPX4 are associated an exceedingly rare pattern of malformation of Sedaghatian-type Spondylometaphyseal Skeletal Dysplasia (SSMD), brain malformations, hypotonia, and neonatal lethality. We report three long-term survivors in two unrelated families identified by whole exome sequencing to harbor a homozygous variant in GPX4 (c.455 G>A, p. R152H, based on NP_001354761.1, the mature short form of GPX4) and who extend the phenotype to include other skeletal anomalies and neurodegeneration.

Proband 1 (family 1) presented with IUGR, microcephaly (~4 SD), hypotonia, hypoplastic optic nerves, and bilateral auditory neuropathy. Skeletal survey demonstrated metaphyseal cupping and irregularity of all long bones. Spine series showed cervical platyspondyly. Serial brain MRIs revealed cerebellar atrophy and supratentorial white matter atrophy. Probands 2 and 3 (family 2) presented with infantile hypotonia and gross motor delays. Skeletal surveys revealed brachycephaly, proximal tibial metaphyseal flaring, and coxa valga. Now age 13 years, proband 2 has intractable epilepsy and has diffuse
supratentorial and infratentorial volume loss on serial brain MRIs. Proband 3 has bilateral auditory neuropathy.

GXP4 encodes for a phospholipid glutathione peroxidase that is critical for eliminating peroxidized phospholipids in membranes. It thus prevents ferroptosis, a type of regulated cell death driven by iron-dependent lipid peroxidation. Animal knockouts of Gpx4 are embryonic lethal; adult conditional Gpx4 knockout mice develop seizures, ataxia and neuronal loss [Yoo et al, 2012]. Computational and cellular modeling from patient-derived fibroblasts revealed a 40% reduction in activity via destabilization of secondary protein structure which in turn disrupts the active catalytic site. This data suggests that the p.R152H variant is a hypomorphic allele that produces sufficient enzymatic function to likely allow for survival beyond infancy but may result in increased susceptibility to ferroptosis culminating in cell death, tissue damage, and neurodegeneration.

#325 EXPANDING THE PHENOTYPE OF PIK3CA RELATED SYNDROME: REPORT OF TWO SIBLINGS WITH NOVEL FEATURES AND GENOTYPE

C Galareta Aina*, 1OR Blair, 1A Slavotinek. 1Valley Children’s Hospital, Madera, CA; 2University of California San Francisco, San Francisco, CA

Case Report Tiosano et al. (2019) identified a novel syndrome characterized by short stature, coarse facial features, ocular and skeletal abnormalities in five individuals with homozygous loss-of-function mutations in PIK3CA. This report describes a pair of siblings of Hispanic ethnicity with novel PIK3CA truncating variants and a phenotype consistent with original report.

The siblings, a 5-year-old boy and 2-year-old girl, share a strikingly similar facial gestalt with coarse appearance, sparse eyebrows, prominent epicanthal folds, broad nasal bridge, prominent nasal tip and full lips. Both have sparse hair, two hair whorls, broad distal phalanges, 5th finger clinodactyly, persistence of fetal pads and flat feet. The boy is short (height Z score -2.5), while the sister has normal height (Z score -1.3). Both have developed gibbus deformity, more severe in the boy who has hypoplasia of L2 vertebral body. He has significant ocular abnormalities, including bilateral congenital cataracts, nystagmus, esotropia and high myopia, and mild to moderate, right-sided hearing impairment that has been presumed to be sensorineural loss. The sister had an uncomplicated perinatal course, whereas the boy had significant respiratory depression at birth and developed persistent pulmonary hypertension in the neonatal period. Both children have dental decay and global developmental delay. The sister had GBS sepsis at 4-weeks-old and was found to have proteinuria which persisted at 2 years of age with UPcr/CR ratio range 3.4-6.5. Plasma albumin and renal ultrasound were normal. Exome sequencing analysis of both children and their mother revealed two novel PIK3CA variants in the siblings, c.4381delC (p.Arg1461Glufs*31) and c.1555C>T (p.Arg519Ter).

Previously reported features present in these patients include short stature, congenital cataract, hearing loss, coarse facial features and scoliosis. In contrast to brother and prior patients, the sister had normal stature and she also exhibits persistent proteinuria which has not been reported. This report brings to 7 the number of patients reported with PIK3CA-related syndrome, confirms core clinical features, establishes intra-familial variability and expands the phenotype to include proteinuria.

#326 ROHHAD MIMICKED BY A TUBB2B NOVEL VARIANT


Purpose of Study Although many individuals have been diagnosed with ROHHAD [rapid-onset obesity, hypothalamic, autonomic dysregulation, neuroendocrine tumor] since this condition was first defined in 2007, the genetic basis remains unknown. Here we describe individuals with ROHHAD-like presentation that underwent medical genetics evaluations. We review clinical characteristics of tubulopathies and query whether they have overlapping features with ROHHAD.

Methods Used Two individuals referred to the genetics clinic for ROHHAD had trio whole exome sequencing performed. One of the individuals was found to have a novel variant in TUBB2B gene. A literature search from PubMed on TUBB2B resulted in 66 articles. 16 were excluded based on a lack of clinical information. In total, 50 articles were reviewed for clinical features that have been described for TUBB2B. Prior studies on ROHHAD were reviewed.

Summary of Results Trio exome sequencing reported a de novo missense variant in TUBB2B c.421G>A (p.Gly141Ser) at a highly conserved amino acid residue in the patient with early-onset obesity (BMI 48 by 1 year life) and additional features. A second unrelated patient had non-diagnostic exome. The individual with TUBB2B variant is a 2-year-old female with severe obesity, developmental delay, hypotonia, obstructive sleep apnea (OSA), hypertension, ocular motor apraxia, hypothyroidism, colitis, clostridium difficile infection, and anemia. The hypothalamic-pituitary axis was tested by ACTH level but in general, the hypothalamic-pituitary axis was tested by ACTH level and stimulation testing and was not diagnostic. Variants in TUBB2B are typically associated with brain malformations and characterized by a wide spectrum of features. It is possible that early onset obesity may be a rare presentation of TUBB2B tubulopathies since the sequencing of 74 obesity genes did not yield an additional diagnostic variant.

Conclusions A novel TUBB2B variant was identified in a patient with brain MRI findings and severe obesity. TUBB2B has been reported to associated with developmental delay in 92% (45/49), motor impairment in 89% (32/36), microcephaly in 86% (44/51), impaired muscular tone in 87% (21/24), ocular features in 74% (17/23), seizures in 68% (34/50), behavior abnormalities 61% (8/13), and craniofacial dysmorphology in 71% (5/7) of the cases. OSA was reported in one case. An association of tubulopathies with rapid-onset obesity would be worth considering. There are numerous examples of genetic conditions with both neurodevelopmental and obesity phenotypes. Possible overlap in ROHHAD-like phenotypes and tubulopathies could involve the central nervous system, but in general, the TUBB2B condition can be discerned by genetic testing and imaging, distinguishable from classical ROHHAD, which would need tumor screening. Further studies may help elucidate the full spectrum of tubulopathies and their clinical presentations.
**Purpose of Study** Barth syndrome is an X-linked disorder that presents with dilated cardiomyopathy and/or left ventricular non-compaction, 3-methylglutacetic aciduria (3-MGA), neutropenia, and characteristic facial features in young males with hemizygous pathogenic variants in TAFazzin (OMIM: 300394). The finding of 3-MGA in patients with cardiomyopathy may be suggestive of Barth syndrome but is not specific to this disorder, and in some cases may misdirect the diagnostic work-up in patients with 3-MGA secondary to other etiologies. We present our institutional experience with the diagnosis and management of Barth syndrome, as well as the role of 3-MGA in the diagnostic evaluation of patients with cardiomyopathy.

**Methods Used** Retrospective chart review of probands with Barth syndrome evaluated at Lucile Packard Children’s Hospital from January 1, 2003 to the present.

**Summary of Results** Seven male patients were diagnosed with Barth syndrome via molecular genetic testing. Cardiac findings at presentation (median age: 2 months, range: neonate to 15 years) included dilated cardiomyopathy with left ventricular non-compaction (n=5) and cardiac arrest (n=2). Urine organic acid analysis noted 3-MGA in 6 of 7 patients, though most elevations were mild. Molecular diagnosis of Barth syndrome occurred via gene panel (n=3), single gene or familial variant testing (n=2), or unspecified genetic testing (n=2), with a median time to molecular diagnosis of 4 months (range: 2 weeks to 9 years). In 4 patients, cardiac function improved over time; 2 patients required cardiac transplant; and 2 patients (including 1 transplanted patient) died from cardiac causes. Median follow-up was 9 years (range: 3 weeks to 24 years).

**Conclusions** Among the 7 individuals with Barth syndrome, the most common features included dilated cardiomyopathy, left ventricular non-compaction, and 3-MGA. Although 3-MGA is a classically described finding in patients with Barth syndrome, elevated 3-MGA levels are not always present. Furthermore, as 3-MGA can also be seen secondary to acute clinical states and/or mitochondrial dysfunction, reliance on 3-MGA in boys with cardiomyopathy may promote premature narrowing of the differential diagnosis and ultimately delay diagnosis of other conditions. Further investigation into the prevalence of 3-MGA in other metabolic and genetic disorders, especially those with cardiac involvement, is warranted.

**Purpose of Study** Osteogenesis imperfecta (OI) is a skeletal disorder typically associated with defects in collagen proteins, the majority due to autosomal dominant pathogenic variants in COL1A1 and COL1A2. However, OI is increasingly found to be genetically heterogeneous with other inheritance patterns observed. Recently, biallelic variants in SPARC, encoding for a secreted protein, acidic, cysteine-rich (SPARC), have been reported in rare cases of autosomal recessive OI (type XVII). Literature review identified 6 cases. Here, we describe an individual who was found to have an earlier reported SPARC homozygous missense variant c.787G>A (p.Glu263Lys).

**Methods Used** Case report with chart and literature review.

**Summary of Results** The proband is a 6yo male born to consanguineous parents. Early on, he had global hypotonia with gross and fine motor delays and remains non-ambulatory. He had bilateral conductive hearing loss but normal speech development. MRI of the brain showed periventricular white matter volume loss. He did not have dentinogenesis imperfecta though did have multiple caries. At 6 months he sustained a femur fracture and subsequently developed >10 fractures requiring multiple surgeries. He developed severe thoracolumbar kyphoscoliosis with multilevel compression fractures. He also had thoracic cage deformity with bony fusion of several ribs, and generalized osteopenia. Molecular testing identified a homozygous variant in SPARC, and his brother with history of fracture after minor trauma was found to be heterozygous for the same variant.

**Conclusions** Our patient shares features common in the previously described cases such as early neuromuscular concerns, scoliosis, long bone and vertebral compression fractures, and delayed motor milestones, suggesting these are consistent across SPARC-related OI. Interestingly, our patient displayed white matter changes on MRI, also found in two other cases. However, our case experienced fractures at a younger age with a more severe course compared to most reports. He also had bony fusion of several ribs and hearing loss, which have not been reported in SPARC-related OI. His family history raises the possibility of mild manifestations for heterozygous carriers. Overall, our patient provides further insight into the variation of SPARC-related OI and expands on the phenotype of individuals with pathogenic variants in SPARC.
Whole genome sequencing (WGS) from the clinical laboratory identified a deep intronic \textit{PTCH1} variant (c.3+314G>T), present also in her healthy father and sibling, which was considered to be likely benign. However, when we reanalyzed the WGS data using two structural variant callers, RUFUS and Smoove, we detected signals suggesting a 38.3 Mb \textit{de novo} balanced inversion inv(9)(q22.32q34.2). One of the breakpoints was located in intron 2 of the \textit{PTCH1} gene. As a result, this inversion is predicted to result in either a truncated protein or mRNA subject to nonsense-mediated decay.

This finding was confirmed by karyotyping, FISH analysis of metaphase chromosomes, and Sanger sequencing. Notably, the initial WGS also identified a maternally-inherited VUS in the \textit{CHD7} gene (c.8077G>C; p.Gly2693Arg), which is associated with \textit{CHARGE} syndrome. The latter finding is likely to account for bilateral hypoplasia of the semicircular canals (not a known feature of GS), and possibly also for Kallman syndrome (reported in several cases of \textit{CHD7}-associated conditions).

MH either has a blended phenotype due to the combined effect of \textit{de novo} \textit{PTCH1} disruption and an inherited \textit{CHD7} variant or she represents an expanded phenotype of Gorlin syndrome. Pursuing and completing the diagnostic evaluation helped a) discover an unusual molecular etiology for GS; b) show the value of using novel tools to identify structural variants with WGS; c) provide more precise clinical management and counseling to the proband and her family.

This was part UDN (U01HG007672).

**Neonatology general IV**

**Concurrent session**

**10:15 AM**

**Friday, January 21, 2022**

**#330** MATERNAL MARIJUANA USE DURING PREGNANCY AND BREASTFEEDING: ASSESSING IN-HOSPITAL OUTCOMES OF EARLY PRETERM INFANTS

1MA Chang, 2S Curtis, 3NL Davis, 1University of Southern California Keck School of Medicine, Los Angeles, CA; 2University of Maryland School of Medicine, Baltimore, MD

10.1136/jim-2022-WRMC.326

**Purpose of Study**

Legalization and decriminalization of marijuana has been associated with increased use in women of child-bearing age. Since Tetrahydrocannabinol (THC) has been shown to pass into breastmilk, concern exists for effects of exposure prenatally and postnatally via mother’s milk (MM). Our neonatal intensive care unit (NICU) allows MM provision for infants regardless of THC status, though many will restrict MM use given concerns about effects on the neonate. Our objective was to evaluate differences in outcomes between neonates who did vs. did not have exposure to THC prenatally and postnatally via MM.

**Methods**

We performed a retrospective medical record review of early preterm (<34 weeks gestational age, GA) neonates born 9/2014–12/2020 and admitted to our Level IV NICU. Our institution conducts universal urine toxicology and substance use screens on admitted mothers. We compared maternal and neonatal characteristics in four cohorts: 1) THC positive moms providing MM, 2) THC positive moms who provided no MM, 3) THC negative moms who provided MM, 4) THC negative moms who provided no MM. We performed multivariable analyses to assess the effect of THC and MM status on significant outcomes.

**Summary of Results**

We identified 763 early preterm neonates, of whom 59% (n=448) were non-Hispanic Black and 31% (n=233) were non-Hispanic White. Seventeen percent (n=130) of mothers tested positive for THC. THC positive mothers were more likely to be late to prenatal care (PNC) while those who did not provide MM were more likely to receive no PNC (p<0.0001). There were no significant differences between the 4 cohorts in regards to growth restriction, oxygen requirement at 36 weeks, necrotizing enterocolitis, and ventilator requirement. However, we did find increased incidence of intraventricular hemorrhage (IVH) in infants who were THC exposed (p=0.0018), though the largest incidence overall and of grade III/IV IVH was in THC exposed infants without MM. We therefore performed logistic regression for predictors of IVH in early preterms and found that when controlling for variables such as birth weight, neither THC status (p=0.18) nor MM status (p=0.79) significantly predicted IVH in early preterms. The only significant predictors were antenatal steroids, delivery mode, and birth GA.

**Conclusions**

Providing MM from THC positive women to early preterm neonates is controversial as longer term effects of this exposure are unknown. In our study, we found no evidence that providing MM from THC positive mothers was detrimental to the health of this early premature population through hospital discharge. A better understanding of longer term perinatal outcomes associated with THC exposure would inform appropriate interventions to improve clinical outcomes and safely encourage MM provision for early preterm infants.

**#331** LATE PRETERM ANTENATAL STEROID USE AND INFANT OUTCOMES IN A SINGLE CENTER

MD Wahl*, C McEvoy, M Go. Oregon Health and Science University School of Medicine, Portland, OR

10.1136/jim-2022-WRMC.327

**Purpose of Study**

Antenatal steroids (AS) have been shown to lower respiratory complications in late preterm infants (LPI) and is now part of the management of pregnancies at risk for delivery between 34 0/7- 36 6/7 weeks gestational age (GA). We aim to determine our center’s adherence to the published Antenatal Betamethasone for Women at Risk for Late Preterm Delivery (ALPS) trial (PMID26842679) and assess neonatal outcomes related to its use.

**Methods**

We conducted a retrospective chart review of singleton mother-LPI dyads who delivered in our center from January 2019 to December 2020. We excluded those with major congenital anomalies. Data on AS use, maternal characteristics and neonatal outcomes were recorded and analyzed.

**Summary of Results**

There were 175 mother-infant dyads included, 92 (53%) received AS and 83 (47%) did not receive AS (No AS) prior to delivery. Maternal characteristics between the two groups such as age, race, ethnicity, delivery mode, smoking status, gestational hypertension and preeclampsia were similar. The incidence of preterm premature rupture of membranes was 66% in the AS group compared to 52% in the No AS group (p=0.02). Diabetes was present in 20% of
the women in the AS group, 11% of which had pregestational diabetes (an exclusion criterion in the ALPS trial). We found 90% adherence to the ALPS protocol within our center. The AS group had a mean GA of 35.5 weeks at delivery and an average time from admission to delivery of 58.2 hours, compared to the No AS group (36.1 weeks, p<0.05 and 19.96 hours, p=0.01, respectively). Neonatal outcomes such as birth weight, need for respiratory support, hypoglycemia, NICU admission and hospital length of stay, were similar between the two groups when controlled for GA.

Conclusions We continue to see some variation in the administration of AS in the late preterm gestation. In our study, AS administration did not improve neonatal outcomes such as the need for respiratory support, NICU admission, and overall hospital length of stay, although we are underpowered for these clinical results. Our study reflects that there remains variation in the approach to women with diabetes delivering during the late preterm period. We recommend that more studies are needed to evaluate the impact of AS use in women with diabetes, as well as overall long-term impact of AS on late preterm infants.
Abstract #333 Table 2

<table>
<thead>
<tr>
<th>Demographics and Hemoglobin levels at Birth Between Groups</th>
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<tbody>
<tr>
<td>Birth weight (g)*</td>
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<tr>
<td>--------------------------------------------------------</td>
</tr>
<tr>
<td>Gestational age (wks)*</td>
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<tr>
<td>Cesarean section, n (%)</td>
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<tr>
<td>Hispanic Race n (%)</td>
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<tr>
<td>Maternal Pre-eclampsia, n (%)</td>
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<tr>
<td>Hemoglobin at 24 hours of life (g/dL)*</td>
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</table>

*Median (25th percentile, 75th percentile)

Methods Used Retrospective study of term infants born at LAC+USC Medical Center and admitted to the neonatal intensive care unit between 2016–2019. Demographics, clinical data, duration of cord clamping and neurodevelopmental outcomes were collected from electronic medical records and paper records. Infants with ABO incompatibility were excluded. Ages and stages questionnaire was used to assess communication, gross and fine motor, problem solving and social skills at 9–12 months and 18–24 months of age. Data was analyzed with Wilcoxon Rank Sum and Chi Square tests.

Summary of Results There were 174 infants included in the study, however only 62/174 (36%) and 28/174 (16%) were assessed for neurodevelopmental outcomes at 9–12 months and 18–24 months of age respectively. Demographics were not different between the two groups except for Hispanic race. Hemoglobin level was significantly higher at birth in infants with DCC. (Table 1). Median neurodevelopmental scores in any domains of ASQ at 9–12 and 18–24 months (table 2) and median hemoglobin levels at 9–12 months of age were not significantly different between ICC and DCC [12.45 (11.7, 12.8) vs. 12.2 (11.6, 13.1) p=0.84]. No infant was treated with iron.

Conclusions Our preliminary data showed that delayed cord clamping in term infants resulted in a higher hemoglobin at birth. However, at 9–12 months of age, there was no difference in the hemoglobin levels between groups. Neurodevelopmental scores at 9–12 months and at 18–24 months were similar. No infant was diagnosed to have iron deficiency anemia. These results need to be verified in a larger sample size.

Nephrology and hypertension

Concurrent session
10:15 AM

Friday, January 21, 2022

Abstract #334 IMPACT OF MATERNAL CHRONIC ILLNESS AND PLACENTAL ABNORMALITIES ON NEONATAL OUTCOMES IN VERY LOW BIRTH WEIGHT INFANTS

K Ramm*, A Hisey, NS Nanduri, Y Shao, C Marquez, L Barton, R Ramanathan, M Birinwale. LAC+USC Medical Center, Los Angeles, CA

10.1136/jim-2022-WRMC.330

Purpose of Study Placental abnormalities and maternal chronic illnesses are often associated with adverse neonatal outcomes including small for gestational age (SGA) and stillbirth; however, there is limited information regarding its impact on morbidities in very low birth infants. This study aims to identify the impact of maternal factors, specifically chronic illness and placental abnormalities, on short term neonatal outcomes in VLBW infants.

Methods Used The data on VLBW infants from LAC+USC medical center between the years of 2009 to 2021 was retrospectively collected from electronic medical records after receiving IRB approval. Maternal factors included placental abnormalities, such as placenta previa, abruption, and placenta accreta and maternal chronic illnesses, such as sickle cell disease, hypertension, diabetes mellitus, hepatitis, cardiomyopathy, asthma, lupus erythematosus, obesity, and cancer. Neonatal outcomes associated with required resuscitation and ventilation were tested for significance using SPSS software version 28.

Summary of Results Of 383 VLBW infants, 48% were born to mothers with a chronic illness and 6.6% of these infants were born to mothers with placental abnormalities. Infants born to mothers with chronic illness were likely to be delivered by c section (88% vs 73% p<0.01) and were found to be SGA (34% vs 23% p=0.09). Only significant neonatal morbidity associated with the VLBW infants born to mothers with chronic illness was patent ductus arteriosus (PDA) (75% vs 65% p=0.02). Maternal illness did not increase the risk for any other NICU morbidities. Infants delivered due to placental factors were more likely to need resuscitation at the time of delivery (67% vs 36% p=0.002), invasive ventilation including high frequency ventilation and surfactant therapy. There was no statistically significant impact on long-term morbidities including bronchopulmonary dysplasia or abnormal MRI related to maternal illness or placental abnormality.

Conclusions Placental abnormalities were shown to significantly affect a VLBW infant’s need for resuscitation and ventilation after birth; however, maternal chronic illnesses cause growth restriction in VLBW infants and predisposes them to have PDA. These conditions do not seem to affect long term neonatal outcomes.

Abstract #335 ANALYSIS OF HUMORAL AND CELLULAR IMMUNE RESPONSES TO SARS-COV-2 VACCINATION (BNT162B2) IN IMMUNOCOMPROMISED RENAL ALLOGRAFT RECIPIENTS

SC Jordan*, B Shin, T Gadde, A Vo, N Ammerman, R Zhang. Cedars-Sinai Medical Center, Los Angeles, CA and Cedars-Sinai Medical Center, Los Angeles, CA

10.1136/jim-2022-WRMC.331

Purpose of Study The SARS-CoV-2 pandemic has resulted in more than 4 million deaths worldwide. Treatments have had limited efficacy, thus the last best hope is development of
effective vaccines. Currently available vaccines, including Pfizer (BNT162b2), demonstrated >90% efficacy against ancestral SARS-CoV-2 infection. However, this did not include an assessment of efficacy in immunocompromised individuals and was dependent on Spike RBD-IgG detection only. Recent data also demonstrate that immunocompromised transplant recipients have increased mortality after SARS-CoV-2 infection which makes efforts for the development of effective vaccine strategies more prescient.

Methods Used
Here, we report on an assessment of Spike-specific IgG and CD4+/CD8+ T-cell responses to BNT162b2 in immunocompromised kidney transplant patients compared to normal controls. We also analyzed the impact of immunosuppressive regimens on immune responses to BNT162b2 assessing patients on Tacrolimus + mycophenolate + prednisone (Tac+) v. patients maintained on belatacept + mycophenolate + prednisone (Bela+). Sixty-one kidney transplant recipients >1M post 2nd dose of the BNT162b2 mRNA vaccine had determinations of Spike-Receptor Binding Domain (RBD)-specific IgG levels and analysis of Spike-specific CD4+/CD8+ T-cell immune responses. Responses were compared to 41 age-matched healthy individuals (non-immunocompromised) controls. Fresh whole blood was collected in sodium heparinized tubes for T-cell stimulation assay. Plasma obtained was stored at -80°C for SARS-CoV-2 Spike RBD IgG analysis.

Summary of Results
Spike-RBD IgG responses were detected in 88% of normal individuals, but only 21% of Tx patients responded at 2M post-vaccination. No healthy controls showed CD4+ T-cell reactivity to Spike proteins prior to vaccination. However, there was a significant response at 1-month post-vaccination. CD4+ T-cell responses post 2nd dose of BNT162b2 were reduced in Tx patients (93% (38 of 41) positive in healthy controls vs 31% (16 of 52) n Tx recipients, p=0.0001). After 2–3 months post 2nd BNT162b2, ~62% of transplant recipients showed positive CD4+ T cell responses. CD8+ Spike-specific T-cells were detected in 56% healthy controls and 37% Tx recipients 1 month post-2nd-dose of the BNT162b2 vaccination (P=NS). No differences were seen by immunosuppression type (Bela+ v. Tac +).

Conclusions
In summary, we found a profound deficiency in SARS-CoV-2 Spike-RBD IgG responses after BNT162b2 vaccination (21% Tx v. 93% Normals) This is consistent with other reports. However, SARS-CoV-2 Spike-specific CD4+/CD8+ T-cell responses were seen in 47% of Tx patients. Analysis of T-cell responses to SARS-CoV-2 vaccination augments our ability to more properly define the scope and durability of immunity as T-cell immunity likely provides protection from a devastating disease.

#336
LEFT VENTRICULAR HYPERTROPHY, ACTIVATION OF MAMMALIAN TARGET OF RAPAMYCIN (mTOR) AND SUPPRESSED AUTOPHAGY IN THE HEART BEFORE HYPERTENSION IN MICE WITH POLYCYSTIC KIDNEY DISEASE (PKD)
OA Otto*, D Poddar, D Atwood, C Edelstein. University of Colorado Health, Aurora, CO

10.1136/jim-2022-WRMC.332

Purpose of Study
To determine cardiac structure and function, blood pressure (BP) and mTOR signaling in the heart in Pkd1RC/RC (RC) mice.

Methods
Used RC mice represent a hypomorphic Pkd1 gene knock in matching a human disease variant, Pkd1 p. R3277C. 120 day old RC mice were studied that have a kidney cyst density (% of kidney ± SEM) of 13.8 ± 2.8. Serial blood pressure measurements were taken in conscious animals using a noninvasive tail-cuff system. mTOR and autophagy proteins were determined in the heart by immunoblot analysis.

Summary of Results
See table 1. Heart weight corrected for body weight (HW/BW) was increased in RC mice. There was cardiac hypertrophy in RC mice as evidenced by increase in both intra-ventricular septum (IVS) and left ventricular wall (LVW) thickness. On echocardiogram there was increased LV mass, decreased ratio of peak velocity of early to late filling of mitral inflow (E/A ratio) indicating grade 1 diastolic dysfunction and decreased LV diastolic volume in RC mice. Blood pressure was not increased in RC mice. Increased mTOR signaling resulting in suppressed autophagy are processes that are known to influence cardiac hypertrophy Polycystin (PC-1), the gene product of the Pkd1 gene, is known to modulate the mTOR pathway. We hypothesized that PKD mice that also have the gene defect in the heart should have upregulation of mTORC1/2 in the heart. Thus, as it was unlikely that increased BP was contributing to the cardiac phenotype, we next looked at mTOR signaling and autophagy. In 70 day old mice, a time point before cardiac hypertrophy, there was an increase in mTORC1 (pAktThr308) and mTORC2 (pAktSer473) in the heart in RC mice. In addition, at 70 days old, there was suppressed autophagic flux in the heart in RC mice as evidenced by decreased LC3-II (a marker of autophagosomes) and no increase in LC3-II with bafilomycin.

Abstract #336 Table 1

<table>
<thead>
<tr>
<th></th>
<th>WT</th>
<th>RC</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>HW/BW</td>
<td>0.47</td>
<td>0.53</td>
<td>≤0.001</td>
</tr>
<tr>
<td>IVS (μm)</td>
<td>1045</td>
<td>1286</td>
<td>≤0.05</td>
</tr>
<tr>
<td>LVW (μm)</td>
<td>956</td>
<td>1218</td>
<td>≤0.01</td>
</tr>
<tr>
<td>LV mass</td>
<td>(mg/g)</td>
<td>1.2</td>
<td>1.5</td>
</tr>
<tr>
<td>E/A ratio</td>
<td>1.5</td>
<td>1.2</td>
<td>≤0.05</td>
</tr>
<tr>
<td>LV Diastol Vol (μL)</td>
<td>71</td>
<td>57</td>
<td>≤0.05</td>
</tr>
<tr>
<td>Systolic/Diastolic/Mean BP (mm)</td>
<td>116/88</td>
<td>109/84</td>
<td>NS</td>
</tr>
<tr>
<td>Hg</td>
<td>97</td>
<td>92</td>
<td></td>
</tr>
</tbody>
</table>

NS=not significant

Conclusions
There is increased heart weight, cardiac hypertrophy; increased LV mass, grade 1 diastolic dysfunction, activation of mTORC1/2 and suppressed autophagy in the heart in RC mice before the onset of hypertension

In conclusion, factors other than hypertension like increased mTORC1/2 signaling or suppressed autophagy may play a role in causing cardiac hypertrophy in PKD mice.

#337
ANTINEUTROPHIL CYTOLYSIC AUTOANTIBODY NEGATIVE PAUCI-IMMUNE GLOMERULONEPHRITIS IN A YOUNG FEMALE WITH SYSTEMIC LUPUS
S Garcia*, R Janival, N Sekhon, P Pierre, M Sukkar, J Bhandhol, S Mishra, S Eppanapally. Kern Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.333

Purpose of Study
To determine cardiac structure and function, blood pressure (BP) and mTOR signaling in the heart in Pkd1RC/RC (RC) mice.

Methods
Used RC mice represent a hypomorphic Pkd1 gene knock in matching a human disease variant, Pkd1 p. R3277C. 120 day old RC mice were studied that have a kidney cyst density (% of kidney ± SEM) of 13.8 ± 2.8. Serial blood pressure measurements were taken in conscious animals using a noninvasive tail-cuff system. mTOR and autophagy proteins were determined in the heart by immunoblot analysis.

Summary of Results
See table 1. Heart weight corrected for body weight (HW/BW) was increased in RC mice. There was cardiac hypertrophy in RC mice as evidenced by increase in both intra-ventricular septum (IVS) and left ventricular wall (LVW) thickness. On echocardiogram there was increased LV mass, decreased ratio of peak velocity of early to late filling of mitral inflow (E/A ratio) indicating grade 1 diastolic dysfunction and decreased LV diastolic volume in RC mice. Blood pressure was not increased in RC mice. Increased mTOR signaling resulting in suppressed autophagy are processes that are known to influence cardiac hypertrophy Polycystin (PC-1), the gene product of the Pkd1 gene, is known to modulate the mTOR pathway. We hypothesized that PKD mice that also have the gene defect in the heart should have upregulation of mTORC1/2 in the heart. Thus, as it was unlikely that increased BP was contributing to the cardiac phenotype, we next looked at mTOR signaling and autophagy. In 70 day old mice, a time point before cardiac hypertrophy, there was an increase in mTORC1 (pAktThr308) and mTORC2 (pAktSer473) in the heart in RC mice. In addition, at 70 days old, there was suppressed autophagic flux in the heart in RC mice as evidenced by decreased LC3-II (a marker of autophagosomes) and no increase in LC3-II with bafilomycin.
Purpose of Study Lupus associated glomerulonephritis (GN) typically presents as deposits of preformed immunoglobulins and complement within the glomeruli of the kidney. While, crescentic pauci-immune glomerulonephritis (CrGN) is commonly an anti-neutrophil cytoplasmic autoantibodies (ANCA) associated vasculitis causing rapidly progressive and necrotizing CrGN in the absence of immune complexes. We present the case of a systemic lupus erythematosus (SLE) patient with proteinuria and ANCA negative serology which was found to have pauci-immune GN on renal biopsy.

Methods Used A 23 year old woman with Systemic Lupus Erythematosus presented with generalized body aches, fatigue, and shortness of breath for 3 days. Physical examination revealed tachycardia, tachypnea, and bilateral costovertebral angle tenderness. Further testing revealed elevated Partial Thromboplastin Time and D-dimer. Labs also revealed acute kidney injury with nephrotic range proteinuria. Urine microscopy revealed granular casts of acute tubular necrosis and dysmorphic red blood cell cast leading to a possible diagnosis of proliferative glomerulonephritis or membranous nephropathy. A renal biopsy was performed revealing active and chronic pauci-immune crescentic glomerulonephritis characterized by 20% acute, 5% subacute fibrocellular and 20% remote fibrous crescents. There was 33% global glomerulosclerosis and mild-to-moderate tubulointerstitial scarring.

Summary of Results Pauci immune is a form of GN lacking immune reaction products except for minimal accumulation of fibrin and presents as a renal limited vasculitis. Distinguished by absence of circulating basement membrane antibodies and negative finding on renal immunofluorescence. The likelihood of a patient to have SLE in conjunction with ANCA associated pauci-immune GN is 2%.

Conclusions ANCA negative pauci-immune CrGN is an uncommon presentation of kidney damage in the setting of SLE. Despite source, progression of disease ends with end stage renal failure in ANCA positive/ANCA negative CrGN and LN. We believe ANCA negative CrGN warrants further investigation as it may be considered a separate entity from ANCA positive CrGN, but with no change in current management aside from extrarenal systemic considerations for ANCA positive pauci-immune GN.

Abstract #337 Figure 1 Glomerulus with cellular crescents (acute glomerular crescent)

#338 EATING DISORDERS, LAXATIVE ABUSE, URINARY TRACT INFECTION, AND NEPHROLITHIASIS: A CASE REPORT AND REVIEW OF LITERATURE

H Luo*, DM Vigil. University of New Mexico Health Sciences Center, Albuquerque, NM

10.1136/jim-2022-WRMC.334

Case Report Eating disorders are characterized by a persistent disturbance of eating that impairs health or psychosocial functioning. Renal stones are listed among the complications of eating disorders; however, very few cases have been reported.

We present a case of emphysematous pyelonephritis and nephrolithiasis associated with bulimia and laxative abuse. 49-year-old woman with a history of bulimia, laxative abuse, urinary tract infections, nephrolithiasis, post-traumatic stress disorder was referred to nephrology for her chronic kidney disease and nephrolithiasis. Patient was diagnosed bulimia and laxative abuse when she was 15 years old, which continued into adulthood. At age 37, she was diagnosed with left kidney hydronephrosis requiring a left ureteral stent. At age 46, she developed a 12mm left kidney stone complicated by emphysematous pyelonephritis, requiring an emergent ureteral stent. In our clinic the following labs and imaging were obtained: Creatinine 1.31 mg/dl (eGFR 44 ml/min), Potassium 2.9 mEq/l, HCO3 25 mEq/l, Calcium 9.2 mg/dl, Po4 4.3 mg/dl, Mg 2.1 mg/dl. Urinalysis had RBCs, WBCs, and LCE. Urine culture grew Escherichia coli. CT scan showed an atrophic left kidney and a 13mm staghorn calculus. Urology performed left ureteroscopy, laser lithotripsy and ureteral stent exchange. Stone analysis notable for mixed ammonium hydrogen urate/ carbonate apatite. Repeat CT scan demonstrated 2 small left non-obstructing lower pole nephroliths. 24-hour urorisk panel showed hypocitraturia and was started on potassium citrate for stone prevention. She was referred to an eating disorder specialist and a dietitian. Her clinical course continues to wax and wane with potassium ranging 2.4 to 4.2 mEq/l, and HCO3 18 to 33 mEq/l.

Eating disorders and laxative abuse often lead to extracellular fluid volume depletion, acid base disorders, hypokalemia, and hypocitraturia, all of which predispose the formation of kidney stones. We present this complex case of nephrolithiasis and pyelonephritis associated with an eating disorder and laxative abuse, attempting to explain the potential mechanisms of formation of staghorn ammonium renal stone involving hypocitraturia and frequent urinary tract infections.

Abstract #339 ARE RENAL SPARING PROTOCOLS SAFE AND EFFECTIVE IN LIEU OF RANDOMIZED TRIALS IN HEART TRANSPLANT?

1J Hu*, 2T Singer-Englar, 3N Patel, 4S Kim, 5M Hamilton, 1J Kobashigawa. 1Western University of Health Sciences College of Osteopathic Medicine of the Pacific-Northwest, Lebanon, OR; 2Cedars-Sinai Smidt Heart Institute, Los Angeles, CA

10.1136/jim-2022-WRMC.335

Purpose of Study The calcineurin inhibitors (CNIs), including tacrolimus and cyclosporine, have revolutionized outcomes of heart transplant (HTx) patients but unfortunately have many side effects, which include nephrotoxicity, hypertension, and malignancy. It has been recently demonstrated through two large, randomized trials, the SCHEDULE Trial and the
Abstracts

MANDELA Trial, that weaning of CNIs has resulted in an increase of rejection episodes. Most recently, many HTx centers have been using a modified CNI wean protocol in order to prevent further nephrotoxicity and to avoid kidney dialysis. It appears that an earlier (<2 years) renal sparing protocol (RSP) may have more problems compared to later initiation of the protocol (2–5 years after transplant). Therefore, we reviewed our experience in CNI weaning.

Methods Used Between 2010 and 2020, we assessed 34 HTx patients who underwent our RSP beginning after 6 months post-HTx and divided them into those initiated <2 year and between 2–5 years after heart transplant. Our protocol for CNI weaning is to decrease the CNI by one-half for 2 weeks, then by an additional one-half for 2 weeks, and then stop. Meanwhile, patients are also started on a proliferation signal inhibitor (PSI), mostly sirolimus, beginning at 1 mg/day and the dose is increased accordingly to maintain a PSI level between 4 and 8 ng/mL. Endpoints included subsequent freedom from any treated rejection (ATR, including acute cellular rejection [ACR], antibody-mediated rejection [AMR], and biopsy-negative rejection [BNR]), survival, and freedom from cardiac dysfunction (hemodynamic compromise rejection [HCR], defined as left ventricular ejection fraction ≤40%) in the 1 year following RSP initiation. As CNI wean has also been demonstrated to have beneficial aspects, we also assessed the need for antihypertensive medications, diabetes in terms of hemoglobin A1c, and glomerular filtration rate (GFR).

Summary of Results 89% of patients were successfully weaned off CNI at <2 years and 94% at 2–5 years after transplant. However, RSP was not sustained in 50% of RSP <2 year patients and in 31% of 2–5 year patients. The reasons included rejection, medication intolerance, and death. GFR for both groups was markedly improved at 1-year post RSP initiation. There was no difference between groups in anti-hypertension medications or hemoglobin A1c levels.

Conclusions CNI weaning at a large, single center appears to be safe and efficacious for select patients but sustaining RSP continues to be a challenge. Further development of other RSPs is warranted.

Abstract #330

TRIPLE WHAMMY CAUSING NEPHRONOPHTHISIS AND ATYPICAL OPHTHALMOLOGIC & NEUROLOGIC MANIFESTATIONS

J Froud*, A Egense, SP Shankar. UC Davis Children’s Hospital, Sacramento, CA

10.1136/jim-2022-WRMC.336

Case Report We report on an adolescent male with end-stage renal disease, septo-optic dysplasia (SOD), degenerative high myopia, and developmental delays (DD) with 2q13 homoyz-
gous deletion and 17q12 duplication. Nephronophthisis (NPH) is a genetically heterogeneous cystic renal disorder progressing to end-stage renal disease. Mutations in NPHP1 at 2q13 cause Joubert syndrome (JS) and NPH. JS is characterized by abnormal brain MRI (molar tooth sign), hypotonia, DD, and other multi-systemic features in <30% of cases. The 17q12 micro-
duplication syndrome is characterized by variable features including intellectual disability, hypotonia, seizures, autism, microcephaly, structural brain anomalies (focal cortical dysplasia, periventricular leukomalacia), and eye anomalies such as strabismus, amblyopia, cataract, coloboma, and microphthalmia.

Case Presentation An 11-year-old boy was referred to genetics for molecularly confirmed NPH. He also had DD, learning disabilities, congenital nyctagmus, SOD, amblyopia, degenera-
tive high myopia, and hypothyroidism. A brain MRI in infancy showed heterotopia, lissencephaly, and hypoplastic septum pellucidum. Family history was non-contributory. He was diagnosed with stage 3 chronic kidney disease at age 11 when he presented with polydipsia. Renal biopsy confirmed NPH. A nephronophthisis genetic testing panel revealed homozygous NPHP1 deletion. Given his clinical features were unusual for NPHP1 disorder alone, a chromosomal microarray (CMA) was performed. This identified a homozygous 110 kb deletion at 2q13 involving the NPHP1, MALL, and SMIM37 genes and an additional pathogenic 1.45 Mb duplication at 17q12. Parental genetic testing was declined.

Discussion We describe the first reported case, to the best of our knowledge, of a patient with a dual diagnosis of NPH due to homozygous NPHP1 deletion and the 17q12 microduplication syndrome with unique ocular and neurologic findings. Although JS was considered as a potential diagnosis, SOD, degenerative high myopia, and absence of typical findings such as molar tooth sign made JS unlikely, prompting additional genetic evaluation. Homozygous deletions in NPHP1 account for approximately 21% of all NPH cases and are not typically associated with brain and eye involvement. The 17q12 duplication syndrome is the likely cause of his DD and learning disabilities. However, his specific MRI and eye findings have not been reported in 17q12 duplication. Neither NPHP1 deletion nor 17q12 duplication are known to be associated with SOD. This individual is an example of multiple genetic variations leading to a unique and complex phenotype. Future investigations and similar cases are needed to better understand the genotype/phenotype correlations and interactions.
EMERGING ANTICOAGULANTS FOR THE THROMBOPROPHYLAXIS OF ATRIAL FIBRILLATION IN PATIENTS WITH CHRONIC KIDNEY DISEASE

1L Berger*, 1B Bartnik-Olsen, 1S Ashwal. 1Western University of Health Sciences, Pomona, CA; 2Albert Einstein College of Medicine and Montefiore Medical Center, Bronx, NY

10.1136/jim-2022-WRMC.337

Purpose of Study The mainstay treatment for atrial fibrillation (Afib) thromboprophylaxis is anticoagulants. However, their indication in patients with concomitant end stage renal disease (ESRD) remains complex due to increased risks of bleeding and drug-drug interactions. While the American heart association (AHA) guidelines only recommends warfarin in this context, the European Heart Rhythm Association’s (EHRA) guideline holds no recommendations currently but promotes an personalized approach factoring in patient preferences and clinical situations. This review aims to bridge this contrast and limitation by investigating novel therapeutic options that may allow for safe anticoagulation in patients with Afib and ESRD.

Methods Used We performed a comprehensive literature search for novel anticoagulants studied in the context of Afib and CKD. The databases Pubmed, ScienceDirect and Googler scholars were utilized with articles limited to a publication date of 2010 onward. Further exclusion criteria included all drugs before 2018 that are FDA approved for use in ESRD and already reviewed in the current US and European national guidelines.

Summary of Results The four anticoagulants we found matching our inclusion criteria includes Betrixaban, Fondaparinux, Tecarfarin and Ichorcumab. Tecarfarin has most of the benefits associated with warfarin but with less drug-drug interaction and a decrease kidney dependent metabolism. Fondaparinux and betrixaban have both demonstrated advantages in the context of lower dosage control and a lack of need for laboratory monitoring. Studies have also demonstrated a reduction in duration of hospital stay with Fondaparinux. Ichorcumab has also demonstrated the potential as a safer alternative for those at higher risks of bleeding.

Conclusions Our review suggests there are multiple novel anticoagulants that may diversify the current EHRA and AHA recommendations for patients with concomitant Afib and ESRD. Although these therapies have demonstrated non-inferiority to warfarin, higher level trials are needed to further establish their therapeutic index and efficacy in the context of Afib and ESRD.

Neuroscience II
Concurrent session
10:15 AM
Friday, January 21, 2022

WHITE MATTER METABOLITE RATIOS PREDICT COGNITIVE OUTCOME IN PEDIATRIC TRAUMATIC BRAIN INJURY

1L Berger*, 1B Bartnik-Olsen, 1S Ashwal. 1Loma Linda University School of Medicine, Loma Linda, CA; 2Loma Linda University Medical Center, Loma Linda, CA

10.1136/jim-2022-WRMC.338

Purpose of Study The rapid acceleration-deceleration motion in traumatic brain injury (TBI) generates widespread mechanical shearing forces resulting in diffuse axonal injury and downstream metabolic consequences. This mechanism predisposes regions with high axon density (i.e. white matter (WM) to greater burden of injury. Metabolite changes are well described in adult TBI, however less is known about the longitudinal consequences in the pediatric population. The purpose of this study was to evaluate the prognostic ability of global WM and GM metabolite ratios following pediatric TBI and their relationship to 12 month neuropsychological assessments of IQ, attention, and memory.

Methods Used 3D proton magnetic resonance spectroscopic imaging (MRSI) in pediatric patients with complicated mild (cMild), moderate, and severe TBI was acquired acutely (6–18 days) and 12-months post injury and compared to age-matched normal developing adolescents. A global linear regression model, co-registering MRSI metabolite maps with 3D high resolution magnetic resonance images was used to identify longitudinal white and gray matter metabolism ratio changes.

Summary of Results Acutely, GM NAA/Cr, WM NAA/Cr and WM NAA/Cho ratios were significantly lower in all TBI groups, compared to controls (ANCOVA, p ≤ 0.02; figure A, B). GM NAA/Cho was reduced only in the severe TBI group (p ≤ 0.001). At 12 months, all metabolite ratios normalized to control levels in each of the TBI groups (figure C, D). Acute GM and WM NAA ratios were strongly correlated to 12-month assessments of IQ, attention, and memory (Pearson correlation with FDR, q < 0.001).

Conclusions These findings suggest that whole brain GM and WM metabolite ratios reflect longitudinal changes in neuronal metabolism following TBI, which can be used to predict neuropsychological outcome in pediatric patients.
Abstracts

#343 METABOLIC-DRIVEN ANALYTICS OF PYRUVATE TREATMENT AFTER TBI DETECT MAJOR BENEFITS IN ANTI-OXIDANT AND ANTI-INFLAMMATORY PATHWAYS

N Golovachev*, 1L Siebold, 1J Tan, 2N Harris, 1B Bartnik-Olsen. 1Loma Linda University School of Medicine, Loma Linda, CA; 2University of California Los Angeles David Geffen School of Medicine, Los Angeles, CA

Purpose of Study The purpose of this study was to elucidate the metabolic pathways affected by ethyl pyruvate (EP) given 24 hours after experimental traumatic brain injury (TBI).

Methods Used 18 male rats were randomized into vehicle-treated controlled cortical impact (CCI; N=9) or ethyl pyruvate-treated CCI (N=9) groups. Moderate to severe CCI injury was induced in the left parietal cortex following a 2.0 mm cortical deformation via pneumatic piston. The vehicle group (CCI Veh) was injected with 0.1M phosphate-buffered saline (PBS) and the EP treatment group (CCI EP) was injected with EP (40mg/kg, i.p.) at 0, 1, 3 and 6 hours post-injury. At 24 hours post-injury, rats were euthanized by decapitation under deep isoflurane anesthesia and tissue samples were collected from the left cortex and frozen powdered in liquid nitrogen. Samples were shipped to Metabolon, Inc., (Durham, NC) and prepared for analyses of global metabolic profiles using Ultrahigh Performance Liquid Chromatography-Tandem Mass Spectrometry (UPLC-MS/MS) or Gas Chromatography-Mass Spectrometry (GC-MS). MetaboAnalyst 5.0, GraphPad Prism (Version 9.2), and Sketch (Version 75) were used for statistical analyses and figure composition.

Summary of Results Sample analysis produced 380 detected metabolites. Cystine, ascorbate (vitamin C), nicotinamide ribonucleotide (NMN), adenosine 5‘-monophosphate (AMP), and adenosine from the ipsilateral cortex were all significantly altered following EP treatment 24 hours post-injury (greater than 2-fold change compared to vehicle-treated animals). The top 15 metabolites with the greatest importance from random forest analysis, along with a heatmap, are depicted in figure 1. The top 15 altered pathways associated with EP treatment are depicted in table 1, with pyruvate metabolism and TCA cycle metabolism demonstrating the lowest Holm corrected p-value. Other significant metabolic pathways include purine, cysteine and methionine, and glutathione metabolism.

Conclusions This dosing regime of EP alters metabolic pathways associated with anti-oxidation, anti-inflammation, and metabolism that may contribute to its neuroprotective effects.

Abstract #343 Table 1 Significantly altered pathways after EP treatment 24 hours after TBI. Pathways with >1 metabolite hit and >5 impact score. Table sorted by Holm p value. A metabolite hit is defined as a significantly altered metabolite in the pathway of interest. The pathway impact is calculated as the sum of the importance measures of the matched metabolites normalized by the sum of the importance measures of all metabolites. Acetyl-CoA is seen as a key metabolite present in multiple pathways.

<table>
<thead>
<tr>
<th>Enriched Pathways</th>
<th>Holm p</th>
<th>Impact</th>
<th>Metabolite Hits</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pyruvate metabolism</td>
<td>0.0056112</td>
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<td>Acetyl-CoA; Phosphoenolpyruvate</td>
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<tr>
<td>Citrate cycle (TCA cycle)</td>
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<td>0.03668</td>
<td>Acetyl-CoA; Phosphoenolpyruvate</td>
</tr>
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<td>Fatty acid elongation</td>
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<td>0.03668</td>
<td>Acetyl-CoA</td>
</tr>
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<td>Glyoxylate and</td>
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#344 METHYLprednisolone INDUCES INCREASED NEURONAL TISSUE SPARING AFTER SPINAL CORD INJURY IN A RODENT MODEL

M Solorio*, T Prasse, J Bergquist, ZZ Kaing, CP Hofstetter. University of Washington, Seattle, WA

Purpose of Study Currently, there is no universally accepted neuroprotective drug for traumatic spinal cord injuries (tSCI) in the clinic. Our objective is to identify a treatment to use during the acute phase aimed at limiting neuronal tissue loss after SCI. Previous studies have shown that...
methylprednisolone (MPSS) has neuroprotective effects, possibly through its anti-inflammatory properties. Here we analyze the effects of acute MPSS administration on tissue sparing after a cervical SCI in a rodent model.

Methods Used We used 16 adult female rats randomized into three different groups: laminectomy only (n=3), SCI only (n=8), SCI+MPSS treatment (n=5). They underwent a longitudinal incision over C3-C7, then a laminectomy to expose the spinal cord at the C5 level. The rats that were in the SCI only group and the SCI+MPSS group, underwent a midline incision using an Infinite Horizon impactor at 200 kdyn. Rats that were in SCI+MPSS treatment group received one immediate post injury (60 mg/kg body weight) dose and three additional intravenous boluses (30 mg/kg body weight) of MPSS every two hours after injury. After the SCI, all animals had behavioral testing using functional locomotor (FL) scoring done 7, 10 and 14 (days post injury) dpi. Tissue sparing analysis was performed by measuring the lesion area of cross-sectional slices after immunohistochemistry (IHC) staining, which visualized anti-GFAP and anti-CD68.

Summary of Results All the injured animals showed forelimb motor function improvement over time after injury. However, there was no statistically significant difference in between any of the injured groups. Tissue sparing analysis showed that animals in the SCI+MPSS group had a significantly higher mean tissue sparing percentage of 86% [± 14.2]. Animals in the SCI only group had a mean tissue sparing percentage of 76% [± 18.5] at the lesion area (p < 0.0001). Lesion area analysis also showed a significantly larger area of damage detected in SCI only animals with a mean difference of [0.82 mm²] compared to animals in SCI+MPSS group (p < 0.0001).

Conclusions These results show promising neuroprotective properties of MPSS becoming effective by promoting acute tissue sparing when given immediately after SCI, which implies increased neuronal function. Although there was no statistically significant difference in behavioral analysis between treatment groups, this is most likely due to the comparably short survival time and a similar extent of injury in all groups. More research is needed to identify new strategies to minimize the systemic side effects of MPSS and optimize the local effective dose.

#345 ATYPICAL PARSONAGE-TURNER SYNDROME INVOLVING CRANIAL NERVES 8, 9, 10, 11 AND THE PHRENIC NERVE

1C Munhall*, 2K Helland. 1The University of Arizona College of Medicine Phoenix, Phoenix, AZ; 2HonorHealth, Scottsdale, AZ

10.1136/jim-2022-WRMC.345

Case Report Parsonage-Turner syndrome, also commonly referred to as Neuralgic Amyotrophy (NA), is a condition leading to symptoms centered around the brachial plexus. It often arises as abrupt unilateral shoulder pain, followed by paresthesias and progressive weakness. Despite unclear etiology, NA has been associated with postinfectious, postsurgical, post-vaccination, and posttraumatic states. Patients are usually treated with pain management and physical therapy and usually exhibit good long-term recovery.

Although classically involving the brachial plexus, cases have been documented involving a variety of other nerves. A 2016 review in the journal of Neurology and Neurophysiology found only 22 cases with cranial nerve involvement reported in the English literature to that date. CN 10 was most commonly involved and presented most often as a laryngeal nerve palsy. The authors noted phrenic nerve involvement in 6 out of 22 cases, three of which were asymptomatic and found via chest x-ray. This case report highlights a unique presentation of NA which exhibited involvement of cranial nerves (CN) 8, 9, 10, and 11 as well as phrenic nerve involvement.

Case A 70 year old man with a history of moderate bilateral sensorineural hearing loss (SNHL) vaccination. Flexible laryngoscopic exam demonstrated a sluggish true left vocal fold with impaired adduction. Imaging of the chest and neck showed no lesion impinging on CN 10, but demonstrated diaphragmatic paralysis confirmed with fluoroscopy.

The patient followed up several months later reporting new dysphagia and microaspiration. The patient was noted to have velopharyngeal insufficiency with reflux of liquids into the nasopharynx, left soft palate paresis, and right uvular deviation suggesting impairment of left CN 9 in addition to CN 10. Audologic evaluation revealed worsening of left SNHL from moderate to severe to now profound, representing involvement of CN 8. A neurologist additionally noted weakness of the left trapezius muscle consistent with a CN 11 deficit.

After 2 years of ongoing dysphonia and microaspiration despite speech therapy, the patient received a left true vocal fold injection which improved his dysphonia and aspiration risk. The patient noted ongoing dysphagia, and subsequent transesophageal manometry demonstrated absent esophageal peristalsis consistent with left CN 10 dysfunction. The involvement of left CN 8, 9, 10, 11, and phrenic nerve in addition to brachial plexopathy was a uniquely atypical presentation of NA. The atypical involvement of multiple cranial nerves in a variety of neurologic syndromes, including NA, Bell’s palsy, Guillain-Barré, and many others should be a clinical consideration for physicians when evaluating patients with multiple cranial nerve palsies.

#346 SCREEN TIME AND CHANGES TO OCULO-PHYSIOLOGICAL MEASUREMENTS

K Inouye*, P Davey. Western University of Health Sciences, Pomona, CA

10.1136/jim-2022-WRMC.342

Purpose of Study Due to the COVID-19 pandemic, the average amount of time spent in front of a screen has spiked considerably. Prior studies suggested that reading standardized text/font from an iPad causes measurable physiological and visual changes compared to paper. This is particularly evident if digital devices used at nighttime are leading to sleep disturbances. What is not known is if the use of digital devices during the day leads to measurable physiological changes compared to reading print media.

Methods Used We recruited 30 volunteers to evaluate whether these changes could be observed in 30 minutes using a cognitively demanding reading task. These volunteers were asked to do a total of 3 readings: iPad in well-lit room, paper in well-lit room, and iPad in dark room. Physiological and visual measurements were taken before and after readings, and patients were asked to fill out a dry eye questionnaire before and after each visit.
Summary of Results Our results showed a significant increase in dry eye symptoms across all readings (p<0.05), although there was no significant difference between symptoms for iPad and paper when reading in a well-lit room (p=0.89). We also noted decreased blood pressure across all three readings, increased contrast sensitivity with paper and decreased contrast sensitivity with iPad in a well-lit room, and decreased blue cone sensitivity with iPad in a dark room.

Conclusions Based on the parameters measured, we can surmise that 30 minutes of reading a collectively demanding reading task is not enough time to evoke the previously observed physiological responses to screens, although the visual changes were consistent. The results of the dry eye symptom questionnaire suggest that for shorter periods, ambient light might alleviate ocular fatigue that was noted when reading from the iPad in a dark room.

Purpose of Study Chronic cough (CC), defined as cough lasting at least eight weeks, is a complaint of up to 46% of patients referred to specialist clinics. Previously, studies have reported a relationship between CC and voice disorders. However, studies have not reported on the specific diagnosis of the voice disorder or on the time relationship between the patients’ first voice complaint and the voice diagnosis made by an otolaryngologist. The data in this study examine complaints of voice changes (hoarseness) and the length of time before a diagnosis of the voice disorder.

Methods Used The first complaint of hoarseness or voice changes in 105 patients with a diagnosis of CC was identified from medical records between January 2015 and February 2020. The diagnoses of the voice disorders were determined following referral to an otolaryngologist. Inferential statistics were used to determine differences in gender, age, types of voice disorders, and the time between voice complaint and the voice diagnosis. Common comorbidities were also documented.

Summary of Results Of the 105 patients, 18 males and 87 females, data were available for 88 patients to determine that the average time between complaint of hoarseness and diagnosis of a specific voice disorder was 32.3 months (Range: 1–180 months). There was no difference in age between males and females (p=0.05). For the group, the most common diagnoses were neurogenic voice disorders, namely, vocal fold paresis (n=47) and vocal fold atrophy (n=28). Common comorbidities included allergies, gastroesophageal reflux, and airway disorders.

Conclusions This study adds to the complexity of treating chronic cough. The results of the present study suggest that neurogenic voice disorders are the most common voice diagnoses in patients with CC. Other treatments that did not resolve the CC were explored prior to referral to an otolaryngology clinic. In reviewing the medical records, we found that patients were often referred to other specialties for treatment of gastroesophageal reflux, allergies, and various airway disorders despite their initial voice complaints. This study points out the need for team management, including an otolaryngologist, in patients experiencing hoarseness, dysphonia, or other common voice complaints along with the primary complaint of cough. The range of voice disorders and comorbidities in the cohort also demonstrates the complexity of diagnoses of chronic cough.

Purpose of Study Pain related to knee osteoarthritis (OA) is a common and growing presentation to medical facilities. Intra-articular hyaluronic acid (HA) injections along with intraarticular corticosteroid injections represent minimally invasive and cost-effective treatment options. In rare cases, the use of hyaluronic acid has resulted in pseudo sepsis. Our study evaluated the efficacy of combined corticosteroid and HA injection in treatment of pain associated with knee OA. In addition, we hypothesize that corticosteroid preceding HA injections can reduce the incidence of pseudo sepsis.

Methods Used A retrospective chart review was conducted to identify patients with pain related to knee OA treated with corticosteroid and HA (177 patients, average age 75 years, 80 females and 97 males) as well as corticosteroid alone (296 patients, average age 65 years, 216 females and 80 males). Under sonographic guidance, a corticosteroid (Group #1 - 40 mg Depo-Medrol, 25 patients, 40 mg Kenalog, 152 patients; Group #2 - 40 mg Depo-Medrol, 51 patients, 40 mg Kenalog, 241 patients, 4 mg Dexamethasone, 1 patient) was injected into the knee joint. In Group #1, under sonographic guidance, HA (Synvisc-One® 48mg) was injected 10 ± 8 days following the corticosteroid. A 0–10 patient self-reported pain scale was used as the primary outcome. For statistical analysis a repeated measures mixed effects model was used to fit the percent pain reduction values. Secondary outcomes included adverse events (pseudo sepsis) and additional treatments. Pain scores were collected prior to treatment and at 1 month, 3 months, 6 months, and 12 months post-treatment.

Summary of Results Reported pain level was reduced at all time points in both groups compared to pre-treatment levels. The overall percent reduction in pain was greater in Group 1 compared to Group 2 at all recorded time points. At one year post treatment, there was an average of 45% pain reduction of Group 1 compared to 34% in Group 2. No adverse events were reported.

Conclusions Combined intraarticular corticosteroid and HA injections demonstrated improvement in pain reduction without increasing the incidence of pseudo sepsis.
compared to corticosteroid alone up to one year post treatment. In addition, the combined therapy resulted in no reported cases of pseudo sepsis. Based on our study, combined corticosteroid and HA injection represents a safe and effective treatment option in patients suffering from knee OA pain.

**Abstracts**

**#349 RURAL GENERAL SURGERY CHALLENGES AND RESOURCE LIMITATIONS IN WASHINGTON STATE**

KC Larson*, V Lobova, S Lewis, AS Hopp. Washington State University Elson S Floyd College of Medicine, Spokane, WA

Purpose of Study Rural general surgeons operate in an environment that differs significantly from colleagues practicing in urban or academic settings. Challenges experienced by this surgical subgroup will be unique and may not be elucidated unless intentionally identified. Understanding these challenges provides insight into their research priorities and how we can best support this distinct group of surgeons going forward. As an initial starting point, we seek to form connections with and among these surgeons through personal interviews and site visits. Our end goal is creating a consortium of rural general surgeons for Washington state.

Methods Used We conducted a pilot mixed-methods study to explore the challenges and research priorities of rural general surgeons in Washington State. Critical access hospitals (CAH) were identified by the Washington State Department of Health having met the criteria outlined by federal designation under the Rural Hospital Flexibility Program. Surgeons practicing at CAHs were identified using online facility directories and through connections via faculty and CEOs/CMOs. Communication included email, phone calls, and mailed letters. Once identified, part 1 consisted of virtual, semi-structured interviews and surveys. Part 2 included comprehensive in-person interviews and site visits to the facility. Part 3 will include creating a consortium of rural general surgeons and further analysis of the qualitative data.

Summary of Results We contacted 79 rural general surgeons in Washington resulting in 25 interviews, 17 completed surveys, and 3 in-person site visits. From the interviews, Washington rural general surgeons identified the following areas of concern: blood bank and operating room supply limitations, adequacy of resident training for solo rural practice, limited staff and nurse training, minimal peer support, restricted medical subspecialty consultation, and high surgeon attrition due to high turnover of both surgeons and staff.

Conclusions We established a myriad of systematic healthcare concerns regarding providing quality rural general surgery care in the state of Washington. We hope to utilize this research to raise awareness of the current challenges and investigate solutions to mitigate the challenges in rural general surgery.

**#350 COMPARISON OF MULTIPORT VERSUS SINGLE PORT VIDEO ASSISTED THORACOSCOPIC SURGERY FOR PEDIATRIC PNEUMOTHORAX**

MA Sacks*, L Goodman, YS Mendez, FA Khan, A Radulescu. Loma Linda University Medical Center, Loma Linda, CA

Purpose of Study The purpose of this study was to compare the operative outcomes of single port and multiport video assisted thoracoscopic surgery (VATS) for primary spontaneous pneumothorax (PSP).

Methods Used After Institutional Board Review (IRB) approval, a retrospective study of a single surgeon’s experienced using the thoracoscopic technique during a two-year period reviewed thoracoscopic cases for PSP in children (<18 years). Seven cases classified into three traditional multiport VATS (MP-VATS) and four single port VATS (SP-VATS). Times were compared between initial chest tube placement, surgery, chest tube removal, discharge, and follow up. Morphine Milligram Equivalents (MME) were calculated for opioids using the opioid conversion guide by the Center for Disease Control. Summary of Results Seven patients were included in the study. The median age was 14.88 years [range 12–17 years], but similar between the groups. Both groups had predominant male presence 67% MP and 75% SP. The average time from chest tube placement to surgery was 3.6 days (MP 4.32 vs SP 3.06, p=0.21). Operating time averaged 1.2 hours and was similar between the groups: MP 1.04 vs SP 1.33 hours (p=0.09). The estimated blood loss was higher for MP 5.33ml vs SP 2.25ml (p=0.11). Intraoperative pain medications were delivered weight-based including intravenous acetaminophen and intradermal 0.25% bupivacaine hydrochloride and epinephrine. MME were similar intraoperative and postoperative until chest tube removal. The chest tube removal occurred at 5.54 vs 3.59 days MP vs SP respectively (p<0.05). Length of stay was 10.46 vs 8.33 days for MP and SP (p=0.30). In an average of 10 months follow up, one SP patient had a recurrent pneumothorax after chest tube removal, requiring replacement of chest tube, and one MP patient required an additional surgery for contralateral PSP.

Conclusions In this small case series, outcomes were not significantly different for single port when compared with traditional multiport VATS for PSP. This provides a convincing basis to expand the usage for this technique and analyze with a larger group.

**#351 FAMILY PRESENCE IN THE OPERATING ROOM**

1SK Kamer*, 2Z Quinnones, 3K Pyke-Grimm, 3–5 D Char. 1Promona College, Claremont, CA; 2Stanford Medicine, Stanford, CA; 3Lucile Salter Packard Children’s Hospital at Stanford, Palo Alto, CA

Purpose of Study Care locations are becoming more fluid, with patients often moving between emergency room (ER), imaging, operating room (OR), and intensive care unit (ICU) for

## Abstract #350 Table 1

<table>
<thead>
<tr>
<th>Timing</th>
<th>Procedure</th>
<th>All Cases</th>
<th>OR Time (hours)</th>
<th>CT removal (days)</th>
<th>Overall LOS (days)</th>
<th>24 hours Postoperative removal</th>
<th>Postoperative Surgery and postoperative course</th>
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<tbody>
<tr>
<td>Anaesthesia in terms of Morphine Milligram Equivalent (mg/kg)</td>
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<tr>
<td>MP-VATS</td>
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<td>0.30</td>
<td>0.17</td>
<td>0.40</td>
<td>0.41</td>
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</table>

*p CT is chest tube (thoracostomy), *p LOs is length of stay, **significant (p<0.05)
resuscitations and diagnostic procedures which will inform acute decisions. Locations like the ICU and ER often have spaces that can be rapidly converted from bedspace into operating rooms. With family presence (FP) already allowed in some of these locations, should (and so in what contexts) families be allowed to be present in the OR similar to how they are being allowed in these other care locations?

Congenital Heart Disease (CHD) is the most common type of birth defect leading to critical illness and death in the US. Given the complexity of disease, these children repeatedly undergo invasive diagnostic and therapeutic procedures. Families often want to be present to support their children through these procedures, and want as much information as they can gather to inform difficult clinical care choices. We assessed the perceptions of clinicians caring for children with CHD about FP in the OR.

Methods Used We conducted semi-structured interviews of 7 clinicians (anesthesiologists, nurses, and surgeons) at a high-volume pediatric cardiac surgical center. We discussed previous experience with and understanding of FP, perceptions toward FP in the OR, and support needed for implementing FP to the OR. Responses were analyzed using grounded theory and a codebook was developed.

Summary of Results Several significant themes emerged including: the role of family member emotions on FP success; potential benefit of FP on reducing patient anxiety and fear, with concomitant reduction in need for sedation medications; whether FP is needed in contexts when the patient is unconscious; and, if FP might distract clinicians. Clinicians also expressed the need for structural changes to support FP particularly cultivating necessary staff who could explain what is going on to the families and policy for who is responsible for the family if they are present during procedures.

Conclusions This pilot study suggest potential benefits of FP in OR for children with CHD: 1) enhancing family understanding of clinical care to inform decision-making; 2) reducing patient anxiety and need for anesthetics or restraints; and, 3) end-of-life or serious situations where family perceptions of grief may be positively impacted by FP. However, these benefits are mitigated by possible harms: 1) medical-legal concerns will need to be addressed so clinicians feel comfortable being observed; and, 2) distractions to the care team by FP including self-consciousness, lack of training managing families’ emotions, and the possibility that certain families’ emotional states may worsen patient and team distress with consequent impact on clinicians and critical clinical workflows.

Abstract #352 Figure 1

CASE REPORT: ANESTHESIA MUMPS FOLLOWING OPERATIVE REPAIR OF A TRAUMATIC ORTHOPEDIC INJURY

1H Hajeh*, 2J Miller, 1M Gill, 1J Bhandohal. 1Kem Medical Center, Bakersfield, CA; 2Ross University School of Medicine, Miramar, FL

Case Report A 25-year-old male presented to the emergency department after sustaining a gunshot wound to the left elbow. On presentation, the patient endorsed elbow pain and physical exam revealed swelling and bleeding from the site of injury. X-ray imaging showed a comminuted, intra-articular fracture of the distal humerus with fragments of shrapnel. Patient was sent to the operating room for open reduction and internal fixation, ulnar nerve neuroplasty and fragmented bullet extraction. The surgery lasted 2 hours with no operative complications. After 8 hours postoperatively, the patient complained of swelling of the right cheek. He denied any pain or loss of sensation other than mild ‘soreness and tension’. Physical examination showed a right parotid swelling with 9.3 cm cranioaudal and 6.2 cm ventrodorsal dimensions.

Facial nerve assessment was normal with no restriction in the temporomandibular joint movement. The enlargement was firm, mobile, with no warmth nor erythema. No tenderness was elicited on palpation. Immunization records weren’t available and the patient was unable to recall his vaccination history including Mumps, Measles or Rubella. Mumps antibodies in serum test showed Mumps IgG antibodies concentration of 27.70 AU/mL and IgM antibodies titre of <1:20 denoting immunity and no acute infection. Of note, the patient denied any post-surgical complications including similar parotid enlargements. A clinical diagnosis of ‘anesthesia mumps’ was made and the patient was managed with supportive care. On postoperative day two, the parotid gland remained swollen, however markedly decreased in size from the day prior. The patient remained otherwise asymptomatic, denying pain or discomfort. With this significant clinical improvement, the patient was discharged before complete resolution of the swelling due to the benign nature of the condition. He was scheduled for a follow up in the clinic, however the patient cancelled the appointment. A phone call appointment was arranged and the patient endorsed reduction in the size of the swelling and complete resolution 2 days after discharge.

Conclusion Although cases of anesthesia mumps are usually benign with no treatment required, they remain stressful for the patient and the physician alike if the benign nature of the condition wasn’t recognized. This warrants spreading awareness among physicians on this case and assuring the patients that such conditions are benign and complete resolution is expected.

UNUSUAL CASE OF PRIMARY SMALL BOWEL BEZOAR CAUSING INTUSSUSCEPTION

1MM Won*, 2MA Sacks, 1RM Leigh, 2L Goodman, 2E Tagge, 2A Radulescu. 1Loma Linda University School of Medicine, Loma Linda, CA; 2Loma Linda University Children’s Hospital, Loma Linda, CA

Case Report Intussusception, the prolapse of one section of intestine into another, is a common cause of small bowel obstruction in pediatric patients. Bezoars are concretions of ingested foreign material. Trichobezoars, bezoars made of hair, more commonly occur in female pediatric patients. If gastric
trichobezoars grow to sufficient size, ‘Rapunzel syndrome’ may occur as the bezoar extends into the intestines, creating lead points for intussusception to occur. Rarely, hair passes completely through the stomach and forms a trichobezoar within the small bowel. This obstruction can also create lead points and cause intussusception. This is the third recorded case of intussusception due to a primary intestinal bezoar.

Case Presentation We present the case of an 8-year-old male with a preliminary diagnosis of appendicitis. Upon further imaging, bowel obstruction related to a small bowel to small bowel intussusception was discovered to be the probable cause. Diagnostic laparoscopy revealed an ileo-ileal intussusception caused by an ileal bezoar, with subsequent evacuation of the causative bezoar.

Conclusion This is the second reported case of an isolated ileal trichobezoar causing intussusception. While intussusception secondary to a trichobezoar most commonly occurs due to Rapunzel syndrome, it is still possible for an intestinal trichobezoar to be present without an associated gastric trichobezoar.

Case Report Limb length discrepancies (LLD) pose a challenge to the orthopedic community with their multivariate etiologies; ranging from congenital malformation to acquired conditions that include bone loss from open fractures, nonunion, osteomyelitis, tumor, malunion, bone debridement due to infection, or comminution. Limb lengthening as a corrective measure occurs via distraction osteogenesis, a process entailing cutting and slowly separating the bone via a device called a distractor, allowing for the bone healing process to fill the gap over time. While historical Ilizarov principles of lengthening at a rate of 1 mm per day continue to be viable, the complications and inconveniences associated with Ilizarov method external fixation devices drove surgeons to develop alternative corrective interventions. To address this issue, Paley et al developed the PRECICE (Nuvasive) system, a magnetic intramedullary (MI) lengthening nail and the corresponding technique that would allow for internally based limb lengthening. We report one patient with history of complex left lower limb discrepancy with evidence of a combination of post-traumatic and congenital etiology presenting with genu valgum. With these indications, the patient underwent a corrective osteotomy of the left distal femur with retrograde placement of a PRECICE MI lengthening nail system. The patient had successful LLD correction with robust healing 9 months follow-up.

A MASSIVE PHYLLODES TUMOR OF THE BREAST

Case Report Phyllodes tumors are fibroepithelial neoplasms that make up less than 1% of breast tumors with an average size of 6 cm. They usually arise in the third and fourth decade of life in women and the mainstay of treatment is surgical excision. Here, we present a rare case of a highly vascularized benign phyllodes tumor that weighted 7.98 kg.

Methods Used Retrospective case report

Summary of the results A 55-year-old female without family history of breast or ovarian cancer presented for evaluation of a non-tender growing mass in her right breast. She initially noted the mass after a mechanical injury to her right breast two years prior to presentation. On physical examination, a firm non-mobile mass was palpated. There were not any skin changes or nipple discharge.

Ultrasound of the right breast revealed a large heterogeneous lobulated mass measuring 27.5 x 21 cm, replacing a great portion of her right breast. CT chest with contrast demonstrated a large soft tissue density mass measuring 25 cm. Core needle biopsy was completed, and showed highly cellular spindle cell proliferation composed of fusiform cells with hyperchromatic nuclei and increased mitotic figures.

The Immunohistochemistry panel revealed that the lesional cells were positive with antibodies directed against Vimentin+, SMA f+, and GATA3 f+. Patient underwent a right breast mastectomy. Tumor was found to be massive with weight of 7.98 Kg and highly vascularized with venous hypertension. This led to a high amount of blood loss for which the patient received 2 units of packed red blood cells.

Abstracts

#354 COMPLEX LOWER LEFT LEG DEFORMITY CORRECTION WITH PRECICE SYSTEM: A CASE REPORT AND REVIEW OF THE LITERATURE

1M Porter*, 1AE Tellis, 1MG Johnston, 1Washington State University Elon S Floyd College of Medicine, Spokane, WA; 2Providence Orthopedics and Sports Medicine, Spokane, WA

10.1136/jim-2022-WRMC.350

Abstract #354 Figure 1 A: Lower Extremity X-Ray standing scanogram on blocks, 2.5/8-inch block under left foot date 11/19/2019; B: Lower Extremity X-Ray 2 months status post-surgery 03/16/2020.

#355 A MASSIVE PHYLLODES TUMOR OF THE BREAST

1F Moshaghi Shari*, 1J Davis, 1Western University of Health Sciences, Pomona, CA; 2Arrowhead Regional Medical Center, Colton, CA

10.1136/jim-2022-WRMC.351

Abstract #355 A MASSIVE PHYLLODES TUMOR OF THE BREAST
Abstracts

Gross examination of the mastectomy specimen showed multiple focal areas of ulceration, necrosis and cystic areas occasionally filled with hemorrhagic gelatinous material.

Morphological evaluation revealed spindle tumor cells arranged in long sweeping fascicles with herringbone pattern and storiform areas consistent with a benign Phyllodes tumor.

Discussion Phyllodes tumors are usually painless tumors with an average size of 6 cm. However, as seen in our patient, these tumors can rarely be larger than 20 cm. Etiology of these tumors remains unknown. Trauma has been previously hypothesized to be involved in development of phyllodes breast tumors through possible fibroblast growth stimulators such as endothelin-1. Our patient with a phyllodes tumor reported a breast injury preceding her diagnosis of breast cancer. Most reports of phyllodes tumors of the breast endorse surgical resection of the neoplasm as a primary treatment without focusing on intraoperative complications. During our surgery, we encountered a highly vascular tumor that led to the need for blood transfusion after implementing clamping and electrocautery technique.

CASE REPORT

Human papillomavirus (HPV) is a sexually transmitted infection (STI) acquired through oral, skin-to-skin, or genital contact. The CDC estimates 79 million Americans are infected with HPV, making it the most prevalent STI in the country. This virus persists in 10% of people, in the epithelial linings of the oropharynx and anogenital tract. Although HPV may persist asymptomatically, it has the potential for oncogenesis.

Increasing HPV prevalence has led to a rise in related cancers. In 2018, head and neck squamous cell cancer (HNSCC) was the sixth most common cancer worldwide, with nearly 900,000 new cases and approximately 450,000 related deaths. The incidence of HNSCC is expected to increase by 30% by 2030. While tobacco and alcohol are important risk factors for HNSCC, increasing rates of HNSCC in the U.S. are primarily due to oropharyngeal HPV infection. The diagnosis, management, and surveillance of HPV-driven HNSCC and related cancers is thus an increasingly relevant clinical consideration. In this case, we present a patient who developed a concomitant HPV-driven anal cancer, atypical after HNSCC, detected and treated early through serial PET-CT surveillance.

Case A 70-year-old female with a history of cervical dysplasia status post hysterectomy presented to an otolaryngologist with globus sensation and oropharyngeal swelling. Exam demonstrated a left tonsillar mass and left neck lymphadenopathy. The tonsillar mass was biopsied and found to be p16-positive squamous cell carcinoma, indicating HPV involvement. Computed tomography (CT) of the neck demonstrated two enlarged lymph nodes. She was treated with surgical resection of the mass and ipsilateral neck dissection. Serial whole-body positron emission tomography (PET-CT) scans showed no evidence of recurrence or metastases, but scans just more than 2 years after treatment revealed a second primary cancer (SPC) of the anus. This anal cancer also demonstrated p16 positivity.

The patient was treated with chemoradiation therapy with a complete response.

Discussion The rising prevalence of HPV infection raises the likelihood of SPVs in patients with a history of HPV-driven cancers. While approximately 90% of local recurrences occur within two years of initial treatment, this does not account for SPC development. Patients with primary HNSCC have a high risk of developing SPVs of the head and neck and rarely metastases. SPVs of the anogenital tract following HNSCC are less common, but people with any HPV-driven primary exhibit 7-fold increased risk for a SPC development. This paper hopes to highlight early detection and treatment of a SPC of the anus through whole-body PET-CT surveillance. The role of such scans in evaluating for local recurrence, metastases, and SPVs is an important question in managing patients with HPV-driven HNSCC.

Joint plenary session

WAFMR, WAP, WSCI, AND WSPR

Friday, January 21, 2022
1:30 PM – 4:15 PM

CASE REPORT

A MASSIVE PHYLLODES TUMOR OF THE BREAST

1K. Murhall, 2K. Heiland, 1J. Newell. 1The University of Arizona College of Medicine Phoenix, Phoenix, AZ; 2HonorHealth, Scottsdale, AZ

10.1136/jim-2022-WRMC.352

Purpose of Study Due to human papillomavirus (HPV) associated oropharyngeal squamous cell carcinoma (OPSCC) having significantly improved prognosis, the American Joint Committee on Cancer (AJCC) introduced a distinct staging system for p16+ OPSCC in its 8th edition. The clinical nodal staging system removed multiple pathologic factors present in the 7th edition, including nodal quantity, and extracapsular extension (ECE). This study aimed to characterize whether the simplification of the staging system resulted in the loss of prognostic value by using the National Cancer Database (NCDB).

Methods Used The NCDB was queried for patients diagnosed with p16+ OPSCC. Patients with no staging information, metastatic disease, who did not receive definitive surgery or radiation, or had unknown follow-up were excluded. The prognostic impact of nodal size, nodal quantity, nodal laterality, and ECE on overall survival (OS) were assessed using survival analysis with the Kaplan Meier method, univariable, and multivariable Cox proportional hazards regression.

Summary of Results A total of 21,868 patients met the inclusion criteria. On Kaplan Meier analysis, patients with more than one positive lymph node had significantly inferior OS (p<0.001; 5-yr OS: 82% vs 86%). Patients with ECE also had inferior outcomes (p<0.001; 5-yr OS: 82% vs 75%). A largest nodal size of >6 cm was also associated with inferior OS (p<0.001); 5 yr OS: 66% vs 82%). Lastly, patients with contralateral or bilateral nodal involvement also had inferior OS (p<0.001; 5 yr OS: 71% vs 84%). On multivariable Cox regression, having more than one positive node (p<0.001; HR [95%CI]: 1.17 [1.07–1.28]), ECE (p<0.001; HR [95%...
TELEMEDICINE INCREASES ACCESS TO PEDIATRIC TREATMENT MANAGEMENT AMID THE COVID-19 PANDEMIC

From January to March 2020, the COVID-19 pandemic emerged in the US and NM forcing healthcare practices to reimagine patient care. NM shelter-in-place orders started in March. Initially following these orders, telehealth visits were the only option for healthcare services. The Healthy and Fit Children’s Clinic (HFCC) at UNM leveraged telemedicine for medical management of their pediatric patients with overweight and obesity through partnership with established telemedicine clinics, transitioning within one month of the pandemic’s onset to a 100% virtual healthcare model. This study assessed this transition in relation to four objectives: patient volume, billing/reimbursement, no-show rates, and patient satisfaction.

Methods Used Improvement science methodology guided the project throughout. The clinic partnered with the HRSA-funded telemedicine project (#H2ARH3037) to implement telemedicine in the care and prevention of childhood obesity. Telemedicine clinic encounters were billed through UNM hospital. Multiple iterative convergent PDSA cycles informed step by step telemedicine implementation workflow. Information was collected from scheduled clinic visits and now shows/cancellations, billing and reimbursement, and a six-question patient satisfaction survey about their clinic visits.

Summary of Results Following telemedicine implementation (April-August 2020), the median number of patients seen per week at HFCC increased by 120% and no-shows declined overall by 20%, which indicated a sustained positive shift above the established pandemic baseline (Jan-March 2020). Telemedicine clinic visits were billed at $122 to $177, an increase from $24 to $35 for telephone visits, leading to a 74% increase in total monthly median billing from April 2020 which indicates a sustained positive shift above the established baseline, surpassing even pre-pandemic billing (April 2019-March 2020.) Over the study period, satisfaction surveys were completed by 26 patients ages 7-15 years old, eight of whom were monolingual Spanish speakers. These patients rated satisfaction with clinic visits ≥90% across all six questions compared to 83% previous global provider rating for in-person visits.

Conclusions With rising rates of childhood obesity exacerbated by substantial pandemic weight gain among youth greater than pre-pandemic rates (Woolford et al, 2021), finding an effective model for consistent feasible healthcare is crucial for accessible quality care continuity particularly for the more vulnerable underserved populations seen in NM. This study demonstrated that telemedicine has potential to improve patient outcomes, improve patient satisfaction, and receive reimbursement for services. Future steps should assess a hybrid clinic model to maximize benefits of face-to-face visits with convenience and reliability of telemedicine.
importance of screening and treating neuropsychiatric symptoms in this disorder. Future directions clinically include cohort expansion and deep phenotyping of neuropsychiatric symptoms. We are also developing 3D neural culture models of CACNA1C variants to understand the role of calcium flux in neuronal migration and cortical development.

**Abstracts**

#360 SYSTOLIC DYSFUNCTION AT THE TIME OF DIAGNOSIS CORRELATES WITH AMYLOID BURDEN AND PREDICTS MORTALITY IN TRANSTHYRETIN CARDIAC AMYLOIDOISIS

1S Bukhari*, 1V Malayala, 2Z Shahid. 1Temple University, Philadelphia, PA; 2Jeanes Hospital, Philadelphia, PA

10.1136/jim-2022-WRMC.356

**Purpose of Study** Transthyretin cardiac amyloidosis (ATTR-CA) is generally considered to be associated with diastolic heart failure; however, systolic heart failure (HFrEF) can also be seen, particularly in patients with advanced disease. The mechanisms and prognostic implications of systolic dysfunction in patients with ATTR-CA have not been systematically studied, which we tried to explore in our study.

**Methods Used** In our prospective registry, the diagnosis of ATTR-CA was based on positive PYP scintigraphy and negative serum studies for AL amyloidosis. Patients were classified as ATTR-rEF (ejection fraction, EF<40%) or ATTR-pEF (EF>40%) based on cardiac magnetic resonance imaging (CMR). CMR was used to measure global extracellular volume (ECV) for quantification of amyloid burden. Kaplan-Meier survival analysis and adjusted cox proportional hazard analysis were performed.

**Summary of Results** Of the 124 ATTR-CA patients (mean age 79.9 ± 7.4, 87% men, 90% Caucasian), 51 (41%) were ATTR-rEF. Compared to ATTR-pEF, ATTR-rEF patients were more symptomatic ( NYHA-FC ≥ 3, 61% vs 26%, p<0.001), had lower prevalence of obstructive coronary artery disease (CAD) (37% vs 55%, p=0.05), worse mean diastolic dysfunction (3 vs 2.15, p<0.01), lower tricuspid annular plane systolic excursion (TAPSE <1.7, 59% vs 25%, p<0.001) and worse renal function ( creatinine, 1.63 ± 0.85 vs 1.27 ± 0.55 mg/dl, p<0.01). On CMR, ATTR-rEF group had higher ECV compared to ATTR-pEF (mean ECV, 62% vs 51%, p<0.01). Over a mean follow up period of 1.5 years, 27 (22%) patients died. ATTR-rEF was associated with higher mortality compared to ATTR-pEF (35% vs 12%, p=0.002; HR 3.7, 95%CI 1.62–8.63, p<0.01). On Cox proportional hazard model adjusting for TAPSE and creatinine, reduced EF was an independent predictor of mortality (HR 3.02, 95% CI 1.30–7.10, p=0.01). When divided into EF ≥ 50%, EF 41–49% and EF ≤ 40%, there was stepwise increase in risk of mortality (p<0.01).

**Conclusions** HFrEF is present in more than one-third of patients with ATTR-CA at the time of diagnosis, and is an independent predictor of mortality in ATTR-CA. ATTR-rEF patients also have higher ECVs, indicative of higher amyloid burden and potentially more advanced disease.

**Behavior and development II**

**Concurrent session**

**8:00 AM**

**Saturday, January 22, 2022**

#361 EFFECTS OF EXERCISE ON REALIGNING A DISRUPTED MOLECULAR CLOCK

1ET Scholten*, 2E McCoy, 2EE Schmitt. 1University of Washington School of Medicine, Seattle, WA; 2University of Wyoming, Laramie, WY

10.1136/jim-2022-WRMC.357

**Purpose of Study** Approximately 18% of the US population work irregular shifts. This can lead to a circadian misalignment of our body’s internal molecular clock. Our clock is critical for biological processes such as hormone secretion, body temperature regulation, and sleep/wakefulness cycles. Shift work is a constant in our society so it is valuable to study mechanisms that could mitigate its negative effects. Previous research has shown that voluntary exercise has promising effects on regulatory gene frequency in peripheral tissues in mice. Therefore, the purpose of this study was to determine the effect of forced treadmill exercise on circadian rhythm realignment.

**Methods Used** 24 (n=12 males, n=12 females)C67/B1 four-month-old mice were used in this experiment. The 4 experimental conditions (n=3) included: 1) disrupted + exercise (DE), 2) disrupted + sedentary (DS), 3) non-disrupted + exercise (NDE), and 4) non-disrupted + sedentary (NDS). The disrupted groups were circadian disrupted in a separate vivarium by changing their 12 hr wake and sleep cycle by one hour every day. Pre and post dual-energy X-ray absorptiometry (DEXA) measurements were also recorded on all mice to determine changes in body composition of the experimental groups. The gastrocnemius and soleus skeletal muscles (SM) were extracted, RNA was isolated, then qRT-PCR was performed on key circadian clock genes (Bmal, Per2, Clock) to...
Summary of Results

Analysis of body composition showed that the males in the DE, DS, NDE, and NDS all lost percent body fat of 14.4%, 9.7%, 10.35%, and 11.3%, respectively. Analysis of the female mice displayed a decrease in the DE and DS groups of 3.3% and 11.5%, respectively, while NDE and NDS groups of female mice both increased in percent body fat of 19.75% and 10.65%, respectively. Analysis of key circadian clock genes in the SM of male mice showed no significant differences in Per2 or Bmal expression between groups, yet there was a trend toward significance (p=0.09) in Clock gene expression in DE mice. In the SM of female mice, there was no change in Per2 expression, but there was a trend towards significance (p=0.07) in Bmal expression in the DS group, and significance (p=0.01) in Clock expression in the DE group.

Conclusions

Our results demonstrate that exercise does play a role in the re-entrainment of a misaligned molecular clock when examining circadian genes. The results also reveal sex-differences in gene expression, which is an important factor when considering an exercise intervention protocol for individuals that work the night shift or have a disrupted clock. We will continue to analyze gene expression in other key peripheral tissues (heart, lung, kidney) to continue to identify positive responses to an exercise protocol to re-align a disrupted circadian rhythm.

**Abstract #362**

**Development and Pilot Study of a Booklet to Support Conversations of Quality of Life and Goals of Care Among Parents Expecting a Baby with a Fetal Anomaly**

**M Coquillette, 2H Nsier*, 2AG Dempsey, 1Children’s Hospital Colorado, Aurora, CO; 2University of Colorado Denver School of Medicine, Aurora, CO**

**Purpose of Study** To develop a booklet supporting conversations of quality of life and goals of care with parents expecting a baby with a fetal anomaly. Aims: 1) assess acceptability, feasibility, demand, implementation, and practicality; 2) evaluate and refine the booklet; and 3) evaluate participants’ qualitative feedback regarding preliminary satisfaction with the booklet.

**Methods** Used A mixed-methods study involving N=20 expectant mothers presenting in a fetal care center at a children’s hospital, approached at 30–36 weeks gestation. At enrollment, participants completed a demographic and background questionnaire and were given the booklet. The booklet was adapted for fetal care from an existing aid used by surrogate decision-makers, with input from multidisciplinary healthcare staff at the study site. Participants were sent surveys 2-weeks after enrollment and 4-weeks after delivery. Outcome measures included characteristics of recruited and enrolled parents, refusal rate for participation, completion rate of

<table>
<thead>
<tr>
<th>Study Aim</th>
<th>Question/Prompt (Question Type)</th>
<th>Participant Quotes (2-Week Survey)</th>
<th>Participant Quotes (4-Week Survey)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aim 1: Assess feasibility, acceptability, demand, implementation, and practicality</td>
<td>Who did you talk to about the questions? (Multiple choice)</td>
<td>'My partner' (50%), 'Alone' (38%), 'My family' (13%)</td>
<td>'Alone' (100%)</td>
</tr>
<tr>
<td></td>
<td>When is the best time to give the guide to parents? (Multiple choice)</td>
<td>'Soon after fetal diagnosis' (33%), 'At a follow-up appointment' (33%), 'At the delivery planning meeting' (33%)</td>
<td>This question was not asked on the 4-week survey.</td>
</tr>
<tr>
<td>Aim 2: Evaluate and refine the booklet and its processes</td>
<td>The guide helped me to… (Multiple choice)</td>
<td>'Identify family strengths' (75%), 'Understand by baby’s condition' (25%), 'Facilitates conversations with health care' (25%), 'Talk to friends' (29%), 'Talk to the medical team' (29%), 'Helps with communication' (50%)</td>
<td>'Too long' (25%)</td>
</tr>
<tr>
<td></td>
<td>The problems of the guide are… (Multiple choice)</td>
<td>'Topics are too stressful' (60), 'Too long' (20%)</td>
<td>None reported</td>
</tr>
<tr>
<td></td>
<td>What topics are missing? (Free response)</td>
<td>None reported</td>
<td>'None, each person is looking for different things'</td>
</tr>
<tr>
<td>Aim 3: Evaluate preliminary responses of participants to the booklet</td>
<td>Please give any comments about problems of the guide or how we can improve it (Free response)</td>
<td>'I wish it talked more about what to expect during the birth and time in the CCU' 'Topics or the way they are worded could be triggering to certain people'</td>
<td>None reported</td>
</tr>
<tr>
<td></td>
<td>Please give any other comments about the guide (Free response)</td>
<td>'Helped to ask more informed questions to our care team' 'There was good information to think about and address in the book' 'Even though it makes her [the mother] nervous, it helps prepare her for the future' 'The guide helped me come up with a plan and helped me understand the importance of being aware of any change' 'Good for those that like to journal and write down their feelings' 'I didn’t find any problems in the guide; it just really helps you think deeply'</td>
<td>'I got this really close to delivery, so I had already thought all this through. Would have been great to get this when we got the diagnosis' 'We were pretty prepared, though I can see it may be helpful for families who are not' 'It helped her [the mother] talk to her family and identify strengths and weaknesses'.</td>
</tr>
</tbody>
</table>
Abstracts

surveys, and qualitative feedback about parent experience using the study booklet and parent perception of booklet utility.

Summary of Results Expectant parents expressed willingness and interest in these topics and using this booklet, with 20/21 recruited parents enrolling in the study. Feedback survey response rates were low; 50% of mothers returned at least one feedback survey. Responding participants reported multiple benefits and few concerns with the booklet. At 2-weeks, 100% of respondents indicated they had looked at the booklet and 75% of respondents at 4-weeks indicated they looked at the booklet after the baby’s birth. At 2-weeks, participants indicated moderate satisfaction (78/100 on a visual analog scale) and moderate helpfulness (63/100 on a visual analog scale). At both 2 and 4-weeks, participants indicated they would recommend this booklet to other parents in the fetal care center. Participants’ qualitative and quantitative responses about the booklet are listed in Table 1.

Conclusions The results of this mixed-methods study suggest this parenting population is interested in engaging in conversations about quality of life and goals of care pre-delivery, even with low-resource tools. Attention was high, likely due to the stress of parenting a medically complex child. Yet, a majority indicated it helped prepare them for their infant’s hospital stay and identify family strengths and care goals. Most indicated they would recommend the booklet to other parents of medically complex children. Future study is needed to assess this intervention style’s impact on parent mental health, parenting experience, and ability to engage in medical decision-making for their child. Future studies should investigate the efficacy of ongoing interventions given with clinician facilitation/guidance rather than a one-time intervention.

Impact of Harm Reduction Treatment With or Without Pharmacotherapy on Concurrent Substance Use Among People Experiencing Homelessness and Alcohol Use Disorder

N Mos confronted, 1,2 SE Collins. 1University of Washington School of Medicine, Seattle, WA; 2Washington State University, Pullman, WA

Purpose of Study A prior randomized controlled trial showed that behavioral harm reduction treatment for alcohol use disorder (HaRT-A) – with or without pharmacotherapy (i.e., extended-release naltrexone) – was effective in improving alcohol outcomes and quality of life for people experiencing homelessness and alcohol use disorder. Because nearly 80% of the sample also reported polysubstance use at baseline, this secondary study tested whether HaRT-A also positively impacted other substance use outcomes.

Methods Used In the parent study, 308 adults with current alcohol dependence and homelessness were randomized to receive either HaRT-A plus intramuscular injections of 380 mg extended-release naltrexone (XR-NTX; HaRT-A + XR-NTX), HaRT-A plus placebo (HaRT-A + placebo), HaRT-A alone, or community-based supportive services as usual (control). In this secondary study, we used multilevel mixed-effects models to detect changes in substance use before and after exposure to HaRT-A. For less prevalent behaviors, outcomes included any past-month use (cocaine, methamphetamine, opioids), and for more prevalent outcomes (i.e., polysubstance use and cannabis use), outcomes were past-month frequency (i.e., number of days of use).

Summary of Results Compared to control participants, HaRT-A (with or without XR-NTX) participants showed significantly reduced 30-day frequency of cannabis use (IRR=0.59, 95% CI=0.40–0.86, p = .006) and polysubstance use (IRR=0.64, 95% CI=0.43–0.98). No other significant changes in prevalence or frequency of use were detected.

Conclusions Compared to services as usual, receipt of HaRT-A is associated with reduced cannabis and polysubstance use frequency. These findings suggest that the benefits of HaRT-A may extend beyond its impact on alcohol and quality of life outcomes to positively reshape individuals’ overall substance use patterns. Future studies are needed to further investigate the efficacy of harm reduction treatment for polysubstance use and to develop guidelines for polysubstance use treatment.

Expanding Behavioral Phenotypes in Sex Chromosome Trisomy with Parent Reported Character Strengths

TG Thompson, 1 J Rubin*, 1,2 N Tartajiga. 1Children’s Hospital Colorado, Aurora, CO; 2University of Colorado Denver School of Medicine, Aurora, CO; 3University of North Carolina at Chapel Hill College of Arts and Sciences, Chapel Hill, NC

Purpose of Study Children with sex chromosome trisomy (SCT; XXY/Klinefelter syndrome, XYY, XXX) are known to have increased risks for behavioral health problems including deficits in social cognition, inattention, and increased rates of anxiety and depression. However, a more balanced description of the phenotype that includes strengths is lacking. Limited data on the strengths of school-aged children with SCT has documented profiles of extraordinary kindness, creativity, and perseverance. To date, there has been no holistic description of behaviors, including the assets, of young children with SCT. This study aimed to describe parent reported behavior and strengths in 18–36 month old children with SCT.

Methods Used Data were collected for the eXtreamiday Baby longitudinal natural history study. Parents rated behaviors of their children with prenatally identified SCT (n=140) on the Child Behavior Checklist (CBCL) at ages 18m(n=67), 24m(n=103), and 36m(n=54). Descriptive statistics portrayed Internalizing (IB), Externalizing (EB), and Total Problem (TPB) behaviors at each time point. One-sample T-tests were used to compare the sample to the norming population (M=50, SD=10); one-way ANOVA compared groups (XXY=n=97, XYY=n=13, XXX=n=30). Parents provided free-text responses to the CBCL question: ‘Please describe the best things about your child’. Positive psychology content analysis was used to code qualitative responses for character strengths.

Summary of Results Results for the pooled sample revealed relatively low rates of IB(18m=39.59; 24m=41.34; 36m=44.70), EB(18m=41.34; 24m=43.00; 36m=44.96), and TPB(18m=40.53; 24m=42.43; 36m=45.72), all significantly lower than the norming population at all time points (p<.05). At 36m, boys with XYY showed significantly higher EB than boys with XXY(59.00 v 43.16; p=.012); small sample size in XYY limits interpretation. There were no other group differences. Parents described their children’s strengths effusively; the most frequent codes were loving, happy, sweet, humorous, curious.
Conclusions This study expands the SCT phenotype by providing a more balanced description of behavior. Results indicate that toddlers with SCT show fewer problem behaviors than the general population and parents describe many strengths of character in their young children, including loving, happy, and sweet demeanors. Future analyses will examine how behavior and strengths profiles in SCT develop over the lifespan and how they relate to a variety of psychosocial and developmental outcomes. A more thorough understanding of the phenotype, including a robust understanding of assets, can improve clinical practice and inform development of interventions.

Purpose of Study Adverse childhood experiences (ACEs) are associated with increased risk of diabetes in adulthood. However, the impact of ACEs on clinical outcomes among adults with diabetes is not well understood. We aim to elicit a better understanding of which diabetes co-morbidities have been linked to ACEs from the current literature.

Methods Used In this systematic review, we assessed the literature on relationships between ACE and diabetes clinical outcomes. We scanned references of retrieved articles, and two authors independently evaluated article eligibility, extracted data, and assessed quality. We used meta-analytic methods for outcomes that were included in at least four studies to estimated pooled odds ratios (ORs) using random effects models and the inverse variance method for pooling.

Summary of Results Our search located 458 articles, of which 13 studies were eligible for inclusion, comprising a total of 16,118 individuals. Most studies used clinical samples (n/N=8/11) of African American or white patients (n=5, n=3) with type II diabetes (Type I= 18.2%; pooled= 27.3%). Depression was the most studied co-morbidity (n=6), followed by cardiovascular disease, obesity, HbA1c, and post-traumatic stress disorder (each were assessed twice). The pooled OR for depression among diabetic patients with moderate/high (≥2) relative to low (0–1) ACE exposure, was 2.67 (95%CI: 1.16–6.12; p=0.0537, k=5) with high heterogeneity (I²=0.95).

Conclusions We found a strong association between ACEs and depression among diabetics: additional research is needed before pooled estimates on other clinical outcomes can be calculated. Potential benefits of ACE-informed psychotherapeutic interventions for diabetic patients with high ACEs merits investigation.

#365 ADVERSE CHILDHOOD EXPERIENCES (ACE) AND CO-MORBIDITIES AMONG ADULTS WITH DIABETES MELLITUS: META-ANALYSIS OF THE ASSOCIATION BETWEEN ACE AND DIABETIC DEPRESSION
M Malwane*, J Abad, M Riddle, M Thompson. University of Nevada Reno, Reno, NV
10.1136/jim-2022-WRMC.361

Purpose of Study Adverse childhood experiences (ACEs) are associated with increased risk of diabetes in adulthood. However, the impact of ACEs on clinical outcomes among adults with diabetes is not well understood. We aim to elicit a better understanding of which diabetes co-morbidities have been linked to ACEs from the current literature.

Methods Used In this systematic review, we assessed the literature on relationships between ACE and diabetes clinical outcomes/co-morbidities. We searched Embase, PubMed, and ProQuest through August 12th, 2020 using keywords related to ACEs and autoimmune disease or diabetes. We scanned references of retrieved articles, and two authors independently

#366 ADVERSE CHILDHOOD EXPERIENCES AND DIABETES IN ADULTHOOD: AN UPDATED SYSTEMATIC REVIEW AND META-ANALYSIS
J Abad*, M Malwane, M Thompson, M Riddle. University of Nevada Reno, Reno, NV
10.1136/jim-2022-WRMC.362

Purpose of Study There is growing evidence that adverse childhood experiences (ACEs) increase the risk of diabetes in adulthood through inflammatory and behavioral pathways: previous meta-analyses require updating and exploration of sub-groups to explain heterogeneity of effect in the observed literature.

Methods Used We searched PubMed, Embase, and ProQuest through August 12, 2020, using PRISMA guidelines and key words related to ACEs, autoimmune disease, and diabetes. We included English language peer-reviewed articles and graduate dissertations reporting associations between ACE and diabetes on subjects ≥18 years old. Two authors independently screened studies for inclusion, extracted data, and assessed quality. For studies that estimated the association between ≥4 ACEs and diabetes, we used random effects meta-analysis to calculate the pooled odds ratio (OR), and meta-regression methods to test sources of heterogeneity among pooled effect estimates.
Abstracts

#365 ADVERSE CHILDHOOD EXPERIENCES (ACE) AND CO-MORBIDITIES AMONG ADULTS WITH DIABETES: META-ANALYSIS OF THE ASSOCIATION BETWEEN ACE AND DIABETIC DEPRESSION

10.1136/jim-2022-WRMC.361

Purpose of Study

10.1136/jim-2022-WRMC.362

Conclusions

10.1136/jim-2022-WRMC.363

Purpose of Study

10.1136/jim-2022-WRMC.364

Conclusions

EATING TO HONOR YOUR BODY AND LIFE: RECIPES AND INFORMATION FOR THE BLACKFEET COMMUNITY IN MONTANA

K Jones*. University of Washington School of Medicine, Seattle, WA

Purpose of Study

On the Blackfeet Reservation, food insecurity and chronic disease are prevalent and reduce residents’ life expectancies. When American Indians were forcibly relocated, their communities began to rely on processed foods provided by the government, and over more than a century, cooking with processed foods rather than traditional foods was adopted. This project’s goal was to offer culturally-responsive nutrition education and recipes.

Methods Used

An asset-based, community engagement approach was utilized during the development of this project, and involved consulting Southern Piegan Health Center, the Blackfeet Food Access and Sustainability Team (FAST), the Food Delivery Program on Indian Reservations (FDPIR), and staff at Blackfeet Community Hospital (BCH). These organizations do valuable work on nutrition education and improving the food environment on the reservation, and the goal was to supplement their efforts. The project was influenced by conversations with patients about nutrition as well as by exposure to the local grocery stores and food environment. It was also shaped by literature on nutrition interventions on reservations.

Summary of Results

Average FDPIR food boxes meet a relatively strong Healthy Eating Index value, indicating that healthy choices are available to community members who use FDPIR. Participants may need help choosing these options or tips for utilizing those foods. Additionally, research on another reservation found a need for skill-based nutrition education that addresses time and monetary constraints. So, a nutrition guide highlighting healthy tips was made, and 4 recipes were compiled - 2 simple ways to enliven canned vegetables, and 2 Indigikitchen recipes using traditional Blackfeet foods. Nutrition guides were placed in exam rooms at BCH, where patients often sit waiting for providers. Guides and recipes were provided to partners at FAST Blackfeet, the Blackfeet FDPIR, and BCH.

Conclusions

Using an asset-based approach enhanced this project’s outcome. The materials were shared with organizations that work in the community and can display and distribute them at their food pantries. The recipes can be made available in proximity to the ingredients they require. Feedback from community partners, such as the type of seasonings preferred by community members, was utilized. Examining how the resources are received by the community is the next step.

Diversity, equity and inclusion III

Concurrent session

8:00 AM

Saturday, January 22, 2022

THE PRIORITY OF GENDER & RACIAL DISPARITY IN NEUROLOGY JOURNALS: A BIBLIOMETRIC ANALYSIS, 2016–2021

Authors: T Layne*, M Shaheen, K Schrode.

Institutions: Charles Drew University of Medicine and Science, Los Angeles, CA

Purpose of Study Medicine and the field of neurology are not immune to gender and racial disparities present more broadly in society. Even within academia there remains a persistent lack of representation of women and racial minorities, but with the recent spotlight on discrimination it is more necessary than ever that prevailing disparities are acknowledged by medical journals and reflected in their publication priorities. This bibliometric study assesses whether social justice trends have influenced the number of articles published that discuss gender and racial disparity in the past 5 years within the top three leading neurological journals.

Methods Used The leading journals in neurology were chosen based on the SCImago Journal Rank Indicator, which measures the impact or influence of a journal, and total citations in the past 5 years. The three highest neurology journals based on those two parameters were The Lancet Neurology, Nature Reviews Neurology, and Alzheimer’s and Dementia. Online archives of each of these journals were used and a search was conducted using the key words ‘gender’, ‘race’, and ‘disparity’, to find articles related to such topics. Data was collected from September 2016 to August 2021 by counting the number of articles discussing gender or racial disparity for each month. The data was then processed into charts to provide a visual representation of how important the journal deems topics related to gender and racial inequality.

Summary of Results The results indicated that although there was a slight overall increase in coverage of the aforementioned topics, it was negligible compared to the broad impact of current social justice movements. Between the three leading journals, Nature Reviews Neurology had the fewest publications on both gender and racial disparity. Our data suggests an inadequate publication priority for scholarly work on gender and racial disparity in the chosen journals, over the last five years reviewed.

Conclusions Our assumption is that any trend towards more publications related to gender and racial disparities are reflective of the publication priorities of, and thus the perceived relevance to, the editorial boards of the chosen journals. As physicians whose job is to advance and better healthcare for all our patients, it is necessary that the journals we subscribe and read in the field also reflect the topics that are pertinent today. There appears to be a disconnect between current events and leading journals in neurology. We believe that this bibliometric study, albeit limited by a relatively small sample size, reflects the larger lack of diversity and coverage on socially relevant topics in leading neurology journals. This study can be replicated to other journals and other fields as well to establish a broader view of changes or the lack thereof in medicine.

RACIAL/ETHNIC DIFFERENCES IN ACCESS TO FORMAL HOME CARE AND ITS RELATION TO COGNITION AND ACTIVITY OF DAILY LIVING FUNCTIONS

Authors: T Layne*, M Shaheen, K Schrode.

Institutions: Charles Drew University of Medicine and Science, Los Angeles, CA

Purpose of Study Elderly individuals prefer to age in their home. Formal Home Care Services (FHCS) helps to support an individual’s ability to age at home. Studies have illustrated that FHCS are associated with improved Activity of Daily Living (ADL) functioning and cognition, however, there is a disparity in accessing FHCS among lower socioeconomic groups. Little research exists in the U.S. that endeavor to understand the factors associated with the lack of access to FHCS and its impacts amongst the disadvantaged groups. We aim to examine access to FHCS among African American (AA) and Latino American (LA) geriatric populations and its effect on ADLs, IADLs and cognition.

Methods Used We analyzed data related to receiving FHCS for 1,552 subjects ≥ 65 years old from the National Health and Aging Trends Study (NHATS). The outcomes included cognition, depression, injuries, mobility, and ADL function. Data was analyzed using chi square test and multiple logistic regression.

Summary of Results 25% of study subjects received FHCS [Hispanics (37%) and African Americans (36%) compared to Whites (24%), (p<0.05)]. There was no significant difference between FHCS recipients and facility care recipients in ADLs,
IADLs, cognition and depression (p>0.05). FHCS recipients were more likely to have higher injuries/hospital stays since baseline as well as an increased need for ADL assistance compared to facility care recipients (p<0.05).

Conclusions Although most Medicare beneficiaries who received home care were minorities, further analysis and investigation is needed to understand the factors associated with the poorer care outcomes observed in those who received home care.

#371 INTERVIEWING SPANISH SPEAKING PARENTS ABOUT COMMUNICATION IN THE NICU

E Batton*, L Catalan, M Freeman, C Ramos, S Hurst, K Marc-Aurele. University of California San Diego, La Jolla, CA

10.1136/jim-2022-WRMC.367

Purpose of Study To explore from whom and in what manner parents with Limited English Proficiency (LEP) receive information about their baby and to assess parent satisfaction with communication in the NICU.

Methods Used We conducted in-person interviews of Spanish speaking NICU parents using a semi-structured interview format. Parents of newborns admitted to the NICU for at least one week who identified Spanish as their preferred language were eligible. Interviews were performed by one of two bilingual providers. All interviews were audio-taped and transcribed. Transcripts were reviewed for errors by certified bilingual Spanish speakers and then translated to English. A directed content comparison approach was taken to identify contextualized segments from each interview that correspond to targeted questions. We collected newborn demographic data, length of stay, and SNAPPE II (Score for Neonatal Acute Physiology-Perinatal Extension II), a predictor of neonatal mortality.

Summary of Results Fourteen interviews were conducted. Average gestational age at birth was 33 weeks, 4 days [25.5 – 41.2] and 43% were female. Eighty-eight percent of SNAPPE-II scores at 12 hours of life were between 0 to 20 (low). The average length of stay was 45 days [9–120 days]. At the time of the interview, the infants were 32 days old on average [3–118 days]. The average interview duration was 8 minutes (5–22 minutes). Parents primarily received information from nurses and reported rarely speaking with doctors. They were more likely to receive information when they were present in the NICU and when they asked. Parents feared that interpreters were inaccurate, but no parent reported ever refusing an interpreter. They reported difficulty expressing doubts to the medical team. Parents often experienced a delay in communication, which was perceived to be as a result of not speaking English. Parents expressed variable satisfaction with the frequency of information received but almost all reported being extensively involved in their child’s care. Parents requested more bilingual providers be available in addition to electronic interpreters.

Conclusions Parents with LEP face unique challenges communicating in the NICU. Our findings can inform neonatal quality initiatives to decrease delays and burdens for families with LEP to receive timely and good communication. Future studies are needed to determine the best strategies to facilitate communication with Spanish speaking families.

#372 HOW IMPORTANT IS IT TO BE IN A COMMUNITY THAT REFLECTS YOUR RACIAL IDENTITY? EXPLORING THE EXPERIENCES OF BLACK, INDIGENOUS AND PEOPLE OF COLOR (BIPOC) STUDENTS IN RURAL & UNDERSERVED CLINICAL SETTINGS

B Cedeno Betancourt Sil*, T Keys. University of Washington School of Medicine, Seattle, WA

10.1136/jim-2022-WRMC.368

Purpose of Study Medical educators have recently begun to research clinical learning environments as experienced by BIPOC Medical students. The limited results demonstrate that BIPOC students regularly experience mistreatment and discrimination, often as microaggressions. This qualitative study aims to explore how the type of clinical settings, particularly rural and urban underserved locations impact BIPOC medical student’s experiences.

Methods Used Current self-identified BIPOC University of Washington medical students who participated and completed the Rural/Urban Opportunities Program within the last three years were invited to participate in a 90-minute focus group discussion via Zoom. The sessions were facilitated by a current medical student who utilized guided questions to explore the experiences of BIPOC students in urban/rural clinical learning environments. The guided questions were developed by the research team, and utilized in all five focus groups. From 8/4 – 9/1, 2021, five focus groups were conducted with two to four participants per focus groups and one individual interview. There was a total of 12 participants. All focus groups were transcribed and are being reviewed for thematic content. Preliminary themes were identified by the research team.

Summary of Results BIPOC students expressed general apprehensions of being placed in a rural clinical setting, more so than students who were placed in an urban underserved clinical setting, with BIPOC preceptors that serve BIPOC patients. Students placed with either a BIPOC preceptor, or within a BIPOC community tended to have a more positive clinical experience. However, being placed in an urban underserved setting is not necessarily a protector against microaggressions in the clinical setting. Students placed in a rural community with non-BIPOC preceptors who acknowledged possible microaggressions had a positive clinical experience. In general, BIPOC medical students preferred urban underserved clinical experiences over rural clinical experiences due to the belief that a rural clinical setting would result in a greater number of microaggressions.

Conclusions We found that microaggressions can take place in both urban and rural underserved clinical settings. More research needs to be done to further understand the experiences of BIPOC students to improve diversity in medical education. Immediate systemic changes include acknowledgments of the experiences BIPOC students could encounter in Rural/Underserved clinical settings, and an avenue for safely bringing the experiences up to preceptors and faculty.
#373 THE IMPACTS OF RACE ON PREGNANCY OUTCOME IN WOMEN OF COLOR: OSTEOPATHIC MEDICAL STUDENTS’ PERSPECTIVE

I Adesina*, V Liederbauer, B Novak, AL Nelson, E Guenther. Western University of Health Sciences, Pomona, CA

10.1136/jim-2022-WRMC.369

Purpose of Study The main goal of this study is to evaluate the perceptions and attitudes of medical students about the impact race has on access to quality of care as well as pregnancy outcomes in non-Hispanic Black women compared to other races.

Methods Used A voluntary, anonymous 17-question survey was emailed to students at Western University of Health Sciences in Pomona, CA. Responses to most questions were rated on a 5-point Likert scale from disagree strongly to strongly agree. There was also an open-ended question for participants to share their thoughts about these issues. IBM SPSS Statistics 27 was used to run descriptive statistics and conduct chi-square analysis (p value <0.05 was considered statistically significant). Responses from Caucasian students were compared to those who declared other races. This abstract reports the results for four of the main questions relating to these outcomes.

Summary of Results The response rate was 21% (250) from the 1200 invited students. Racially, 48.0% were white, and 67% were female. First-year students made up 39.6% of the population with subsequent years in declining proportions: 23.7%, 20.8%, and 15.7% respectively. Over 85% of respondents believed that race is a strong predictor of pregnancy outcome; 66% disagreed with the statement that health problems of women of color are primarily due to personal choices they make. Over 87% of respondent agreed the color of a person’s skin can influence the quality of medical care that a person receives.

Conclusions Even though studies show that Black women have a higher maternal mortality risk than white women, 13% of respondents did not recognize the impact that race had on maternal outcomes. One-third of students did not disagree with the statement that Black women are primarily responsible for their own poor health. Importantly, one in seven students was not aware of the increased mistrust women of color have of the medical system. The findings demonstrate remaining education gaps that despite limited response rate, may serve as a baseline to measure the impact of more extensive curriculum changes to teach medical students more about equality and diversity in the future.

#374 VARYING RATES OF DISABILITY EXPOSURE REPORTED ON CORE ROTATIONS WHILE OSTEOPATHIC MEDICAL STUDENTS REPORT SUPPORT FOR FORMAL DISABILITY TRAINING

1KM Lucara*, 1S Henderson, 1M Blair, 2AL Nelson, 1E Guenther. Western University of Health Sciences College of Osteopathic Medicine of the Pacific-Northwest, Lebanon, OR; 1Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA

10.1136/jim-2022-WRMC.370

Purpose of Study In the 2014 US Census, 17.6% of adults were severely disabled. People living with a disability have self-reported higher rates of dissatisfaction regarding unmet needs in their healthcare experience and there are no standards in the Commission on Osteopathic College Accreditation curricula for treating those with disabilities. Without the accreditation standards, we sought to quantify the exposure to caring for patients with disabilities that 3rd and 4th year osteopathic medical students (OMS) get while on clinical rotations, to investigate how prepared students felt they were to care for this population. We also studied the attitudes of OMS towards formal disability training.

Methods Used An anonymous, beta-tested, IRB-approved survey and clarifying disability definitions were distributed electronically to all 3rd and 4th year OMS on clinical rotations in the spring. Descriptive statistics were calculated for each question based on the denominator of individuals who responded to the specific question.

Summary of Results 600 surveys were distributed, and 59 responded with 40 completed. 51% of students reported exposure to caring for people with disabilities prior to medical school. 30% reported being exposed to greater than 20 patients with disabilities on rotations. Reported exposure rates to caring for patients with disabilities differed by core rotation: internal medicine (86%), psychiatry (84%), family medicine (79%), pediatrics (77%), osteopathic manipulative medicine (76%), surgery (53%), and obstetrics and gynecology (36%). Students reported the most comfort with patients with physical/ambulatory disabilities (85%) and the least comfort with sensory difficulties (50%). Using the Likert-scale, 68% reported feeling probably (4/5) or definitely (5/5) prepared to care for patients with disabilities. However, 68% also felt the current curriculum probably or definitely did not fully prepare them and thought formal disability training should be added to pre-clinical curriculum.

Conclusions In this pilot study, experience with the care of individuals with disabilities was observed on all core medical rotations, but some offered minimal exposure. Overall, students were in strong support of implementing formal training in caring for patients with disabilities and the majority felt the current curriculum did not prepare them to care for these patients.

#375 AN ASSESSMENT OF ACCESS TO TOPICAL CALCINEURIN INHIBITORS FOR PEDIATRIC VITILIGO

1H Munzing*, 1,3H Brandling-Bennett. 1University of Washington School of Medicine, Seattle, WA; 2Seattle Children’s Hospital, Seattle, WA; 3UW Medicine, Seattle, WA

10.1136/jim-2022-WRMC.371

Purpose of Study Vitiligo is an autoimmune disease that presents with depigmented patches of skin and hair. Untreated childhood vitiligo can have lasting psychosocial effects on patients and their caregivers. Many insurance companies, Medicaid in particular, consider vitiligo to be a ‘cosmetic disease’ without considering the substantial psychosocial health implications. Absence of an FDA-approved therapy for repigmentation of vitiligo offers further challenge, though off-label therapy with topical calcineurin inhibitors (TCIs) is a first-line therapy. This study examined access to TCIs among patients with Medicaid insurance compared with private insurance.

Methods Used A retrospective cohort study was conducted at Seattle Children’s Hospital’s (SCH) which examined the medical records of patients (180) that were <18-years-old, seen between January 2010 and December 2020, had at least two

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medical visits with a diagnosis of vitiligo, and were prescribed treatment with tacrolimus or pimecrolimus (TCI). Their records were reviewed for prescribed treatment plan, ability to access the TCI medications, and insurance type. Associations of Medicaid vs. private insurance and quality of TCI access were studied. Chi-squared statistical testing was used to determine significant differences between treatment access in Medicaid versus private-insurance patients.

**Summary of Results** Patients with Medicaid represented 66.9% of the study cohort while private insurance accounted for 33.1%. Of the Medicaid patients, 50.4% experienced ‘good access’ (no insurance issues reported) to the prescribed TCI compared to 89.5% of private-insurance patients. ‘Challenging access’ (received medication but experienced: denials initially or during treatment, limited access to more potent TCI’s, or necessity to pay out-of-pocket) was reported for 41.7% of Medicaid patients and 7.0% of private-insurance patients. ‘No access’ was experienced by 7.8% and 3.5% of patients with Medicaid and private insurance, respectively. There is a statistically significant difference in the ability to appropriately access the prescribed treatment for vitiligo for private insurance versus Medicaid patients (p ≤ 0.001). Of the 29 documented prior authorization denials, 18 referenced ‘cosmetic’ conditions for the coverage denial.

**Conclusions** Insurance carriers frequently consider vitiligo to be a cosmetic condition, limiting access to prescribed TCI medications for pediatric patients. Inequity in access to prescribed treatment exists for patients with Medicaid versus private insurance coverage. Medicaid patients are much more likely to experience insurance denial, lapses in treatment coverage, out-of-pocket cost, and limited coverage of stronger TCI formulations.

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**Abstracts**

**#376** TWENTY YEARS ON—HAS PATIENT-CENTERED CARE BEEN EQUALLY WELL INTEGRATED AMONG MEDICAL SPECIALTIES?

S Lim*, A Khormali, RJ Wassersug. The University of British Columbia Faculty of Medicine, Vancouver, BC, Canada

10.1136/jim-2022-WRMC.372

**Purpose of Study** The concept of ‘patient-centered care’ (PCC) was endorsed by a US Institute of Medicine 2001 publication, now cited >5000 times. PCC emphasizes autonomy of patients, prioritizing their subjective assessment of healthcare needs. In this study, we assessed how 6 medical specialties—pediatrics, OB-GYN, orthopedics, radiology, dermatology, and neurosurgery—have engaged with PCC plus two derivative concepts: ‘person-centered care’ (PeCC) and ‘family-centered care’ (FCC) over the past 20 years. We hypothesize that large differences in use of these terms reflect gender differences among physicians within the disciplines. Specifically, female dominated fields will have greater engagement with PCC, PeCC and FCC.

**Methods Used** Reference to PCC, PeCC and FCC in the medical literature for the 6 specialties was used as a proxy for clinical engagement of the concepts in those disciplines. We gathered our sample by cross referencing MeSH terms of PCC, PeCC & FCC and the medical subspecialty in PubMed, CINAHL, and PsycInfo. In each specialty, a Pearson correlation test elucidated the change in number of publications of each concept versus percent of female physicians.

**Summary of Results** There is significant correlation (all p < 0.0001) between reference to PCC in the literature and the number of women in each field except neurosurgery (p > 0.5). Pediatrics showed the most extensive reference to PCC followed by OB-GYN, with a significant difference between all disciplines (p < 0.001). When correlating for the number of papers published in the different fields, reference to PCC is 18X more common in pediatrics than in neurosurgery. PeCC is used in pediatrics and OB-GYN far more than any other specialty. PCC use grew exponentially each year until 2016, when the trend started to flatten/decline. Conversely, PeCC references have only climbed. If this continues, within 3 years, PeCC will overtake PCC as the dominant care centricity term in academic literature. FCC is exclusively used by pediatrics.

**Conclusions** Differences in engagement with PCC, PeCC, and FCC appear to reflect true differences in face-to-face interactions within the disciplines, with high uptake in person-oriented specialties (pediatrics & OB/GYN) and low uptake in technique-oriented specialties (surgery). Our data show these differences correlate with the percentage of females in each specialty. Over the last 20 years, order of PCC uptake in the fields has not changed; the female bias has persisted. Our results do not endorse nor reject PCC/PeCC/FCC ideology. Rather, they recognize stereotypical cognitive differences between men and women. Particularly, Baron-Cohen et al. (2013) have shown women are more attentive to patients’ facial expressions and underlying emotional needs. Thus, as gender proportions continue to shift, we anticipate changes to medical culture. Ideal care will embrace both stereotypical male & female modes of medical practice.

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**Endocrinology and metabolism III**

**Concurrent session**

**8:00 AM**

**Saturday, January 22, 2022**

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**#377** SENSITIVITY AND SPECIFICITY OF THE MONTREAL COGNITIVE ASSESSMENT FOR DETECTING CLINICALLY SIGNIFICANT COGNITIVE IMPAIRMENT IN OLDER ADULTS WITH TYPE 1 DIABETES

R. Kudrna*, J. Choi, L. Forseca, N. Chaytor. Washington State University Elson S Floyd College of Medicine, Spokane, WA

10.1136/jim-2022-WRMC.373

**Purpose of Study** The Montreal Cognitive Assessment (MoCA) is a screening test used to detect cognitive impairment. Data for using MoCA in older adults with type 1 diabetes (T1D) is limited despite increased risk for cognitive impairment in this population.

**Methods Used** Older adults with T1D were administered a battery of neuropsychological measures and the MoCA. Clinically significant cognitive impairment was defined as 2 or more test performances ≥1.5 SD below demographically corrected normative data. Receiver operating curve (ROC) analysis was performed and the Youden index was used to select an optimal MoCA cutoff score.
Summary of Results

201 older adults with T1D (mean age = 68.3, 47% female, 57% with bachelor’s degree or higher, 92% non-Hispanic white) completed both the neuropsychological test battery (47.8% impaired) and MoCA (mean = 25.55, SD = 3.1). ROC area under the curve was 0.745 [95% CI 0.678 - 0.812, p < 0.05]. The standard cutoff score of <26 resulted in sensitivity of 0.604 and specificity of 0.714, while a cutoff score of <27 resulted in a sensitivity of 0.750 and specificity of 0.610. The Youden indices for these cutoff values are 0.318 and 0.360 respectively. Minimally acceptable sensitivity (i.e., >0.80) was obtained when using a cutoff score of <28, while >0.80 specificity was obtained with a cutoff score of <25.

Conclusions
The MoCA appears to have moderate overall utility as a cognitive screening instrument in older adults with T1D. Use of the standard cutoff score of <26/30 may not adequately detect those requiring further evaluation (60% sensitivity). The optimal MoCA cutoff score (based on Youden’s Index) for detecting cognitive impairment defined by neuropsychological testing was <27/30. A score of <28 resulted in acceptable sensitivity, but was accompanied by low specificity (42%), necessitating further cognitive evaluation.

Abstract #377 Table 1 Summary of results

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<th>Specificity</th>
<th>Youden index</th>
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#378
SIDE EFFECT SYNERGISM BETWEEN METFORMIN AND GLP-1 RECEPTOR AGONISTS – AND A SOLUTION

JM Miles*. University of Kansas Medical Center, Kansas City, KS

10.1136/jim-2022-WRMC.374

Case Report
Metformin (MET) is the most widely prescribed medication for type 2 diabetes (T2DM), but it has unfortunate gastrointestinal (GI) side effects (chiefly diarrhea, in 30% or more of patients) that limit its use in some individuals. MET raises circulating concentrations of glucagon-like peptide-1 (GLP-1), increases satiety, and in some studies delays gastric emptying. Roughly half of an oral dose of MET is not absorbed; and is thought to be responsible for bile acid malabsorption. GLP-1 receptor agonists (GLP-1RAs) are often used in combination with MET and have their own GI side effects; as many as 10–20% of subjects receiving GLP-1RAs experience nausea and/or diarrhea. We report 6 patients with type 2 diabetes who developed GI side effects when taking MET and a GLP-1RA in combination. Included were 4 males and 2 females, 70 ± 2 years old, with BMI 34 ± 3 kg/m2 and hemoglobin A1c 7.5 ± 0.8%. All were taking MET 2000–2500 mg/day in two divided doses before meals or otherwise on an empty stomach, with no GI side effects. When exenatide ER 2 mg weekly was added to MET, a 79 y.o. woman developed severe diarrhea, nausea and vomiting. These symptoms resolved completely when she started taking both doses of MET immediately after eating (PC). Initiation of semaglutide 0.25 mg weekly resulted in intolerable nausea in a 68 y.o. woman; her symptoms disappeared with PC administration of MET, and she subsequently had no side effects on the 0.5 mg dose. A 68 y.o. man was given dulaglutide 0.75 mg/week and tolerated it without difficulty, but when the dose was increased to 1.5 mg/week he developed severe diarrhea and nausea that resolved completely when he took a 48 h holiday from MET; symptoms did not recur when he resumed MET immediately PC. A 64 y.o. man tolerated dulaglutide 0.75 mg/week but had abdominal cramping and diarrhea with the 1.5 mg dose. There was complete remission of symptoms with PC administration of MET, and he eventually tolerated 3.0 mg/week with no GI side effects. When a 70 y.o. man increased the dose of dulaglutide from 1.5 mg/week to 3.0 mg/week he began having near-daily diarrhea. He started taking MET on a full stomach and eventually was able to take 4.5 mg dulaglutide with no adverse GI effects. A 69 y.o. man began having diarrhea and abdominal discomfort when the dose of dulaglutide was increased from 0.75 mg/week to 1.5 mg/week. With PC administration of MET, he eventually was able to take 4.5 mg/week with no GI side effects. Conclusions: These 6 cases demonstrate side effect synergism between GLP-1RAs and MET that resolved when MET was taken PC. When diarrhea occurs with initiation of a GLP-1RA, it may thus be an indirect effect mediated via MET, since GLP-1RAs do not cause bile acid malabsorption. After-meal administration of MET allows some patients to tolerate a full dose of both medications, with potential greater benefit in the treatment of T2DM.

#379
METFORMIN INDUCED LACTIC ACIDOSIS: A CASE REPORT

1P Chan*, 2E Deemer, 3L. Mosavi, 2A Heidari. 1Ross University School of Medicine, Miramar, FL; 2Kern Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.375

Purpose of Study
To present a rare case of metformin induced lactic acidosis (MALA)

Methods Used
Retrospective study

Summary of Results
Metformin is the most commonly prescribed and preferred initial drug therapy for type 2 diabetes. Metformin is the only FDA-approved biguanide, due to its lower risk for lactic acidosis. Although rare, metformin-associated lactic acidosis (MALA) has a mortality rate of 31%, which underscores the importance of early diagnosis and treatment.

We report on a 75-year-old female with diabetes mellitus type 2 with HbA1c 13.6% and hypertension presented to the emergency room complaining of 2 episodes of clear, watery diarrhea, nausea, and 2 episodes of nonbloody nonbilious emesis for 2 days. Labs were significant for acute kidney injury (AKI) with creatinine of 5.65 mg/dL, anion gap metabolic acidosis (Na of 129 mmol/L, CI of 96 mmol/L, HCO3 of 20 mmol/L, and corrected anion gap of 19 mmol/L), and severe lactic acidosis of 8.4 mmol/L. Over 3 days, the lactic acid levels were labile at 8–9 mmol/L, despite aggressive fluid resuscitation. Computerized tomography (CT) abdomen/pelvis, retroperitoneal ultrasound, comprehensive stool panel, blood cultures, and wound cultures were negative for infection, while urinalysis and culture were positive for yeast. Patient’s kidney function progressively worsened with oliguria, requiring

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hemodialysis (HD). After 2 HD sessions, the patient’s lactic acidosis and anion gap metabolic acidosis resolved.

Of note, the patient was admitted and treated 1 week prior for intractable nausea/vomiting, acute kidney injury, and urinary tract infection. At discharge, the patient’s labs showed blood urea nitrogen of 18 mg/dL and creatinine of 0.86 mg/dL. She was discharged with ciprofloxacin 500 mg twice daily, Metformin 500 mg twice daily, and lisinopril 2.5 mg daily.

Conclusions Metformin is a first-line diabetes medication with a well-known, rare, side effect of lactic acidosis. As a diagnosis of exclusion with a high mortality risk, it is imperative physicians can quickly identify and treat. The treatment for MALA is HD or continuous veno-venous hemofiltration (CVVH) and should be initiated urgently to prevent further morbidity or mortality.

Ultimately, this patient’s rapid decline following metformin initiation illustrate the importance of early recognition and treatment of MALA.

### Abstracts

**VITAMIN D STATUS AND CYSTIC FIBROSIS RELATED DIABETES: A RETROSPECTIVE CHART REVIEW**

G. Gunawardana*, Y. Peng, M. Wu, J. Alvarez, Y. Tangpricha, Emory University, Atlanta, GA; Emory University School of Medicine, Atlanta, GA

Purpose of Study Cystic fibrosis, caused by a mutation of the cystic fibrosis transmembrane conductance regulator (CFTR) gene, often results in chronic infection and retention of mucus in the lungs. This drastically decreases the functionality of epithelial cells and has been found to be comorbid with organ dysfunctions such as the pancreas, liver, and even vas deferens. Pancreatic disorder in patients with CF has been associated with decreased absorption of fat-soluble vitamins, including A, D, E, and K, but its relationship with vitamin D hasn’t been reported [1]

Methods Used In this retrospective chart review of a longitudinal cohort study of patients identified from the Emory Clinic Data Warehouse, it was attempted to determine the relationship between vitamin D levels and the onset of CFRD. Vitamin D levels were measured via serum 25-hydroxyvitamin D (25(OH)D), with measurements taken between January 1st, 2008, and December 31st, 2012. Patients included as part of the study were CF patients in the Emory Clinic and Hospital from 2002–2012, and were stratified based on decreased vitamin D levels. Log-rank (Mantel-Cox) Tests compared the relative risk of time to CFRD onset by vitamin D status, while Chi-square tests assessed the association between the development of CFRD and vitamin D status.

Summary of Results The chi-square tests concluded that 25(OH)D < 20ng/mL and CFRD development are not independent events (P=0.03*), and Log-rank (Mantel-Cox) Test showed a significant hazard ratio between time to CFRD onset and vitamin D status stratified by deficiency at 25(OH)D < 20ng/mL (95%CI: 1.2, 2.7, P < 0.05**).

Conclusions Adults with CF and 25(OH)D levels below 20ng/mL were at an increased risk of developing CFRD overall, while those below 30 ng/mL didn’t show similar correlations. As such, concentrations of 25(OH)D above 20 ng/mL may decrease risk of progression to CFRD, in those with CF.

**THE DEVELOPMENT OF FOURNIER’S GANGRENE IN A PATIENT WITH A HISTORY OF SCROTAL ABSCESS AND UNDERLYING TYPE 2 DIABETES TREATED WITH A SODIUM-GLUCOSE COTRANSPORTER-2 INHIBITOR**

S Sevilla*, J McBale, A Alkasir, D Curie, University of Nevada Reno, Reno, NV

Purpose of Study Diabetic kidney disease (DKD) as a cause of end-stage kidney disease is increasing over time. DKD is projected to increase by 54% in the US. Glomerular filtration rate (GFR) and Creatine are known biomarkers to detect DKD. This study aims to examine vitamin D and C-reactive protein (CRP) and Triglyceride-Glucose-index (T-G index) as biomarkers for the detection of DKD. Find potential biomarkers to aid in the accuracy of the diagnosis of diabetic kidney disease

Methods Used We analyzed data from The National Health and Nutrition Examination Survey 2001–2010. We classified DKD by GFR or albumin-creatinine ratio. Biomarkers tested were vitamin D, CRP, and T-G index. We analyzed the data using multiple logistic regression and produced a receiver operating curve (ROC) to determine the accuracy of the predictions adjusting for the confounding variables. Data were analyzed using SAS9.4 accounting for the design and sample weight.

Summary of Results Of the 1,691 subjects, 54% had DKD by GFR, 6% had DKD by albumin-creatinine ratio, 52% had high-risk CRP levels, 74% had high-risk T-G index, and 2% had low vitamin D.

Subjects with high-risk vitamin D, high-risk CRP, and the combined high-risk CRP and T-G index were 3, 2, 2.6, 7 times more likely to have DKD (by albumin-creatinine ratio) respectively relative to the other groups (p<0.05). These biomarkers can predict subjects with DKD with 67% to 68% accuracy.

Conclusions CRP, T-G index, and Vitamin D biomarkers can be used to detect patients with DKD. Longitudinal prospective large sample studies would further the study for possible use in the early detection of DKD.
empagliflozin after an A1c of 10.6. Three years later, he was admitted for polymicrobial perineal and scrotal necrotizing fasciitis including part of the bulbar urethra. The extent of his infection required serial debridements and split thickness skin grafts to salvage exposed areas of the left testicle and penile shaft. The veteran was under inpatient care for a total of 45 days during which time empagliflozin was stopped. Subsequent outpatient follow up revealed progressive wound healing and stabilization of his A1c.

Discussion FG is a rapidly progressive disease with a reported mortality rate of 20% to 40%. SLGT2 inhibitors are a relatively new class of blood sugar lowering medications that work by decreasing glucose reabsorption in the proximal tubule. The glycosuria that is facilitated by this mechanism increases the risk of genitourinary infections. In 2018, the FDA sent out a warning regarding rare occurrences of FG with the use of SLGT2 inhibitors. Although a following meta-analysis of randomized controlled trials showed no difference in risk of FG between SGLT2 inhibitors and controls, the number of events was small and led to a wide confidence interval. Thus, an increase in FG risk cannot be ruled out. Further data is needed to support the development of refined guidelines for prescribing SLGT2 inhibitors. A comprehensive history and physical, including past genital infections requiring surgical intervention, may better risk stratify patients for the development of FG prior to initiation of empagliflozin. Alternative treatment options outside of class, such as GLP-1 agonists, may be favorable in the setting of a history of such infections.

Conclusion There is a substantial number of reported cases involving FG and the use of SLGT2 inhibitors in diabetic patients. We intend to add to the pool of data and increase prescriber awareness in order to decrease the incidence and preserve the quality of life of these patients. The implementation of refined drug use criteria for SLGT2 inhibitors may be beneficial.

Genetics II

Concurrent session

8:00 AM

Saturday, January 22, 2022

#383 CREATING A ZEBRAFISH MODEL USING CRISPR TO MODEL PHENOTYPIC CHARACTERISTICS OF CNKSR2-RELATED DISEASE

1SL Cole*, 2S Nishizaki, 1L Aki Higa, 3M Dennis, 1JJ Shen. 1UC Davis, Sacramento, CA; 2University of California Davis, Davis, CA

10.1136/jim-2022-WRMC.379

Purpose of Study The field of genetics is rapidly evolving and more patients are receiving genetic diagnoses for their conditions. Greater understanding of the underlying pathophysiology of genetic diseases is important for improving clinical management and investigating treatment options for affected individuals. Pathogenic variants in Connector Enhancer of Kinase Suppressor of RAS2 (CNKSR2) cause an X-linked condition characterized by epilepsy, sleep disturbances, and developmental delay particularly affecting speech. Even though this is a very rare condition with only ~40 patients identified worldwide, there is more information known clinically compared to our knowledge about the pathogenic mechanism of disease. We aim to use a zebrafish model to provide further insights.

Methods Used We are establishing a zebrafish model to evaluate CNKSR2-related disease. Due to a genome duplication event there are two paralogues, Cnksr2a and Cnksr2b, that are being disrupted individually and in combination, to determine the extent to which the human disease can be recapitulated. Embryos are injected with knockout DNA designed using CRISPR technology. Once hatched, several tests are performed using FishInspector to evaluate for dysmorphology, and Zebrawax to assess for seizures. FishInspector software is used to measure characteristics of wildtype zebrafish against the CNKSR2 disease model. Features evaluated include head size, eye size and spacing. For seizure evaluation, zebrafish are evaluated first in a normal state and then taken through procedures to reduce the seizure threshold. The movement and activities during each cycle are compared between wild type and the disease model.

Summary of Results At a few days of life the zebrafish were evaluated for phenotypic differences in dysmorphology and movement. Preliminary findings indicate that loss of one of the paralogues results in a statistically significant larger head size (macrocephaly) and that the CNKSR2 disease model zebrafish appear to exhibit more activity consistent with seizures compared to wild type.

Conclusions Studying human disease through zebrafish allows for comprehensive phenotyping of rare genetic conditions, and this model represents a valuable translational medicine tool in investigating future treatments such as gene therapies.
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in the mother. This X-inactivation testing revealed a ratio of 100:0, consistent with a highly skewed pattern with the maternal X chromosome active while the paternal X chromosome is inactive.

Conclusions Variation in PTCHD1 has previously been reported in association with a broad phenotype of autism and developmental delay/intellectual disability (DD/ID), however this X-linked disease is rarely reported in females. Here we report a case in which a truncating variant in PTCHD1 was inherited from an unaffected mother to an 8-year-old female proband with DD/ID, and for which X-inactivation studies demonstrated significant skewing towards the truncated allele. This case highlights the utility of genome sequencing and X-inactivation studies to diagnose challenging cases of X-linked disease in females that may otherwise be overlooked.

#385 CALCULATION OF BRAIN ORGANOIDS GLUCOSE CONSUMPTION. A CRITICAL FIRST STEP IN UNDERSTANDING ORGANOID METABOLISM

L Forero,* AE Lee, LM Bird, C Sneathlage, A Muotri. University of California San Diego, La Jolla, CA

10.1136/jim-2022-WRMC.381

Purpose of Study Cortical organoids are in-vitro models of the human cerebral cortex. They are derived from human subjects, cultured as pluripotent stem cells, and then differentiated into self-organizing neuronal tissue. They offer a unique way to explore neuroscience in an accessible and flexible environment free of the constraints of in-vivo testing. However, their fidelity as a model for the human brain is still not entirely clear. With the goal of future use in translational research, further study is needed to compare the metabolism and physiology of organoids to functional human brain tissue. The goal of this study was to find the average glucose consumption of an in-vitro organoid and compare this with the glucose consumption of an in-vivo human brain.

Methods Used We will be using 232 established mature brain organoids derived from a healthy subject divided into six wells with 3mL each of standard neurobasal media. We will estimate weight per organoid based on previously established average values. We will measure at Time A the glucose within three 1mL samples of the organoid media using a YSI analyzer. Approximately 48 hours later, at Time B, we will take another three samples from the other three wells and again assess the glucose content via YSI analyzer. We will then analyze the glucose consumed by calculating glucose per 1 gram of tissue per day. Finally, we will compare this to known values of human cortical glucose consumption.

Summary of Results The average glucose organoid consumption per organoid was 0.035 mg of glucose in 24 hours. On average, an adult human brain consumes 120 grams of glucose in 24 hours. Estimating an average human brain weight of 1300 grams, we estimated that an adult human brain consumes 90 mg of glucose per gram of brain tissue per day. Per our findings, organoids consumed on average 26 mg of glucose per gram of brain organoid tissue per day.

Conclusions We have performed an experiment revealing the average glucose consumption of a mature cortical organoid, something that has never before been published in the medical literature. We have discovered that organoids consume approximately 1/3 of the glucose that the average human brain consumes. This information will help establish further translational potential as well as in any future experiments with organoids involving metabolism.

#386 NEONATAL LETHALITY AND GENETIC MODIFIERS IN A NEW MOUSE MODEL OF NIEMANN-PICK DISEASE, TYPE C

1,2,3JL Rodriguez-Gil*, 1OE Watkins-Chow, 3FM Platt, 1WJ Pavan. 1NHGRI, Bethesda, MD; 2Stanford University School of Medicine, Stanford, CA; 3University of Oxford Medical Sciences Division, Oxford, UK

10.1136/jim-2022-WRMC.382

Purpose of Study Niemann-Pick Disease, Type C (NPC) is a fatal neurodegenerative disorder that exhibits intracellular accumulation of unesterified cholesterol in late endosomes/lysosomes and marked accumulation of glycosphingolipids in neuronal tissue. NPC disease is extremely heterogeneous in the timing of clinical presentation (prenatal to adulthood) and severity. Currently there are no FDA-approved therapies that effectively increase lifespan or slow disease progression. For rare disorders such as NPC where sample size is limited, model organisms have played an essential role in the identification of genetic modifiers. We hypothesize that strain-specific variants between inbred mice will have an impact on disease severity, enabling us to identify potential genetic modifiers of NPC disease.

Methods Used We generated a new mouse model for NPC1 harboring a novel allele (Npc1T1084M) using CRISPR/Cas9-mediated gene targeting.

Summary of Results Neonatal lethality: During characterization of this mouse model, a significant reduction of viable homozygotes was observed (10.5% vs. 25%, p<0.0001). Interestingly, the expected genotype frequency was observed at E19.5 (25%) suggesting Npc1 mutants exhibit a previously uncharacterized lethal phenotype postnatally. We confirmed that the majority of Npc1 mutants died shortly after birth and showed signs of respiratory insufficiency. Subsequent electron microscopy analysis of lung tissues showed atypical surfactant morphology.

Genetic modifiers: We also identified strain-specific QTL affecting lifespan. Interestingly, Npc1 mutants on a B6J genetic background had a more severe visceral pathology of foam cell accumulation than Npc1 mutants on a BALB/cJ background. These underlying pathological changes translated into B6J mutants having a significantly shorter lifespan (70 days) than mutants on a BALB/cJ background (84 days; p<0.0001), suggesting strain-specific modifiers contribute to disease severity. QTL analysis of backcross N2 mutants detected significant linkage to markers on chromosomes 1 (LOD=5.57) and 7 (LOD=8.91).

Conclusions The generation of our new NPC1 mouse model facilitated the first analysis of neonatal lethality in an NPC1 model organism. This was also associated with abnormal lung surfactant pathology. A similar phenomenon has been reported in NPC patients with pre/perinatal onset. This model may serve as the first animal model for neonatal-onset NPC1. Furthermore, our study is also the first to identify genomic regions in Npc1 mutant mice containing potential modifier variants associated with changes in lifespan. Identification of modifiers will contribute to our understanding of the highly variable phenotype observed in NPC patients and advance our efforts to improve patient therapy.
Purpose of Study Hyperammonemia is a documented phenomenon in patients with fibrolamellar hepatocarcinoma (FH). There are several proposed mechanisms for hyperammonemia in these patients, including intrahepatic shunting, increased cell breakdown from chemotherapy initiation, decreased expression of the OTC gene in tumor cells undergoing treatment with chemotherapy, or a paraneoplastic process disrupting the urea cycle. We hypothesize that the etiology of hyperammonemia in patients with FH is due to acquired ornithine transcarbamylase (OTC) deficiency.

Methods Used Chart review, biochemical laboratory analysis, and literature review.

Summary of Results Here we describe a 9-year-old girl with newly diagnosed fibrolamellar hepatocarcinoma (FH) with secondary liver failure, hyperammonemia, renal insufficiency with acute kidney injury, gastrointestinal (GI) bleed, and large portal vein thrombosis. Physical exam demonstrated evidence of portal congestion, including dilated veins on the abdomen as well as an enlarged liver. Ammonia level peaked at 370 umol/L prior to initiation of continuous renal replacement therapy (CRRT). Biochemical laboratory analysis while on CRRT included plasma amino acids with normal glutamine, normal acylcarnitine profile, and significantly elevated orotic acid in the urine. She was experiencing persistent hyperammonemia ranging from 200–250 umol/L despite CRRT, so she was treated with intravenous arginine and Ammonul (sodium phenylacetate and sodium benzoate). These medications are used in the treatment of primary OTC deficiency, and they produced a decrease in ammonia to normal levels in our patient. Other medication options, including lactulose, rifaximin, and other oral medications were not able to be given due to her ongoing GI bleed. Repeat plasma amino acids were sent once CRRT was discontinued and showed elevated glutamine.

Conclusions Our patient demonstrated biochemical findings (hyperammonemia, elevated glutamine, significantly elevated urine orotic acid) and response to treatment that were suggestive of acquired OTC deficiency. Acquired OTC deficiency is defined by decreased OTC activity that is independent of an OTC disease-causing variant. This evidence supports the hypothesis that acquired OTC deficiency is the etiology of hyperammonemia in FH. Surjan et al. hypothesized that the molecular mechanism is related to increased expression of a fusion protein that activates expression of Aurora kinase A. Aurora kinase A then upregulates c-Myc expression, an oncogene that targets ODC. ODC in turn catalyzes the decarboxylation of ornithine in polyamine synthesis, shunting ornithine away from the urea cycle and inhibiting OTC activity. In order to investigate this proposed mechanism, we intend to conduct further studies to measure polyamine synthesis, a downstream effect of the aforementioned mechanism.

Purpose of Study Analyzing the transcriptome of the maternal-fetal interface can provide insight into placental development and pregnancy complications such as spontaneous preterm birth. Placental and decidual gene expression is cell-type specific, so a molecular dissection of cell and tissue types may improve our understanding of the maternal-fetal interface and reveal differences related to disease. We compiled cell and tissue type specific genes from single cell RNA sequencing (scRNA) studies and investigated relative gene expression in bulk RNA sequencing data of the decidua and placenta.

Methods Used We performed a literature search of placenta publicly available scRNA studies published after 2015. We compiled the results of these studies into a specific gene list which described genes uniquely expressed in placental and decidual cells. We tested the accuracy of this gene list by using it to differentiate between paired chorion villous trophoblast (CVT) and decidua basalis (DB) samples from publicly available bulk RNA dataset GSE73714. We then evaluated the ratios of decidual/placental gene expression in other publicly available bulk RNA datasets from the placenta and decidua using t-tests.

Summary of Results We compiled 2,158 genes from 5 scRNA studies, with 1,103 genes from 9 cell types reported as placenta specific, and 1,208 genes from 9 cell types reported as decidua specific. Many genes reported as placenta-specific in scRNA sequencing datasets were highly expressed in both CVT and DB samples from GSE73714. We evaluated the ratios of decidual-placental gene expression in 5 bulk RNA datasets, containing 17 decidua-related and 29 placenta-related samples total. In a dataset containing matched DB and CVT samples, DB samples had a higher decidua-specific gene expression than samples taken from CVT (t-test, p < 0.05).

Conclusions We curated a gene list of placental and decidual specific genes using scRNA studies which differentiated between paired DB and CVT samples. Our findings can be used to study cell composition changes in pregnancy complications. For example, placenta samples from spontaneous preterm births often have upregulated immune signaling genes, and our results could help characterize immune cells associated with those genes. Our study also highlights research gaps including a lack of specificity in scRNA datasets and limited decidual scRNA data. More research on decidual tissue is needed to better understand the maternal side of the maternal-fetal interface.
Purpose of Study University of Nevada, Reno School of Medicine’s (UNR Med) Medical First-Year Intensive Transition (MedFIT) is a two-week orientation program that introduces matriculating students to the academic and extracurricular rigor of medical school. The established program includes a curriculum of lectures intended to introduce students to a normal week of medical school culminating in a low-stakes, professor-written lecture exam and anatomy practical. In addition, students are placed into groups and introduced to their Pack Mentor, a 2nd year medical student that serves as a longitudinal peer mentor. Previous research on UNR Med’s MedFIT has evaluated the curriculum structure through qualitative feedback with student perceptions. However, there is a lack of quantitative data analyzing the real efficacy of each session through objective assessment.

Methods Used A survey was used to evaluate the subjective efficacy of the MedFIT program as well as the objective assessment of their gained knowledge from the program, both pre- and post-MedFIT in July 2021. The survey contained a mix of Likert-style and free response questions for the perceived efficacy and receptivity to various sessions. Additionally, multiple choice questions were included to assess true efficacy through student’s retention of the information. The data collected will be evaluated using a paired t-test to compare pre- and post-MedFIT survey scores of each question. Longitudinal retention of this material and relevance to performance in medical school will be evaluated with a post Block 1 and Block 2 survey in early October and December 2021, respectively.

Summary of Results From preliminary analysis of our results, students viewed the lectures, mock lecture exam, and anatomy practical as the most helpful in preparing them for medical school. The nutrition lecture and student panels were deemed least helpful. We saw large improvements between our pre- and post-medFIT survey scores in students’ confidence to find reputable medical sources, understand the Medical Student Performance Evaluation, and which extracurriculars are available to get involved in. Scores in comfortability regarding the Pack Mentor mentorship program or how to manage stress in medical school lacked significant change. Our largest improvements in actual efficacy include information on student body leadership positions as well as supplemental training sessions with a 51% and 49% increase in scores, respectively.

Conclusions Preliminary results indicate that MedFIT is effective at presenting salient administrative information and policies while acclimating students to medical school but that the Pack Mentor program could be an area for improvement in the future. Potential solutions might be adding in more structured interactions to allow for cohesiveness across all groups and more time for the incoming students to meet their classmates through social wellness activities.
THE QUALITY OF YOUTUBE VIDEOS ON RADIOTHERAPY AND PROSTATECTOMY FOR PROSTATE CANCER

1N Wong*, 2P Ingledew, 1St George’s University School of Medicine, St. George, Grenada; 3British Columbia Cancer Agency, Vancouver, BC, Canada

Purpose of Study Prostate cancer is the 3rd leading cause of death among Canadian men with radiotherapy and prostatectomy as options for each disease stage. Studies show that information about treatments and management of side effects are the most needed information for cancer patients. With ~22 billion people visiting YouTube and many cancer patients using it as a source for cancer related information, it is prudent to understand the quality of information available. This study aims to describe the quality of YouTube videos with specific focus on radiotherapy and prostatectomy and the management of treatment side effects.

Methods Used Videos were selected by searching phrases such as ‘Prostate Cancer Radiotherapy’ or ‘Prostatectomy side effects and/or management’. The first 50 videos were recorded. A rating tool derived from similar tools for the evaluation of websites was adapted to analyse the videos for: currency, attribution, content, coverage and accuracy. DISCERN was used to score attribution and content. Materials from NCCN, UpToDate and cancer.ca were used to develop a consensus document and to evaluate accuracy and coverage of the information presented. Two raters were involved in the review to ensure consistency.

Summary of Results Of the videos analyzed, video length ranged from one minute to one hour long and the dates of creation from 2012 to 2021. Videos were led by physicians (74%), patients (16%) or other health professionals (8%). Of the presenters, physician Video Popularity Index = 23.5 while patient presented videos = 61.4. 57% Radiotherapy videos described the procedure or the risks/benefits of the procedure while 33% provided a description of both the procedure and described risks/benefits. 53% Prostatectomy videos described the procedure or the risks/benefits of the surgery while 33% provided both a procedural description and risk/benefits. 83% Radiotherapy videos covered side effects, 60% covered quality of life, and 70% covered special considerations. In comparison, 89% prostatectomy videos covered side effects, 75% covered quality of life, and 78% covered special considerations. No misinformation was found in radiotherapy videos, however 13% had missing or ill-elaborated information for side effects. Similarly, no misinformation was found in prostatectomy videos but 19% did not discuss the procedure in depth.

Conclusions This study provides a description of online resources available to prostate cancer patients. While most of the information is accurate, not all videos cover quality of life, or could be improved to better explain technical aspects of treatment & management of side effects. This information can be helpful for physicians and patients to navigate educational needs, improve patient understanding and increase patient independence in effective coping and management of side effects.

WHY ARE SOME PARENTS AGAINST OR UNSURE ABOUT VACCINATING THEIR CHILDREN AGAINST COVID-19?

1,2M Bera*, 1,2M Kung, 2K He, 1,2B Afghani. 1University of California Irvine, Irvine, CA; 2Children’s Hospital of Orange County, Orange, CA; 3Children’s Hospital of Los Angeles, Los Angeles, CA

Purpose of Study The COVID-19 pandemic continues to evolve. Increasing community vaccination rates will be an important factor in curbing the spread of the virus. As children return to schools and other activities, they represent a large group of potential vectors. It is important to understand what concerns parents possess regarding vaccinating their children so that we may help address these hesitations.

Methods Used An anonymous electronic survey was distributed via REDCap to parents of children who were seen in the outpatient or inpatient setting in an academic center between September 2020 and September 2021. Parents were asked to qualify their willingness to vaccinate their children against COVID-19 by choosing ‘yes’, ‘no’, or ‘not sure’. They then had the opportunity to share an open-ended response explaining their answer. These responses were then analyzed for common themes among those who responded ‘no’ or ‘not sure’.

Summary of Results Of 189 parents who completed surveys, 15% responded ‘no’ and 30% responded ‘not sure’ when asked if they would vaccinate their children against COVID-19. When asked an open-ended question about their hesitancy, 42 wrote the reason for their concern. The figure 1 below shares the common themes expressed in these responses. Some responses included multiple themes and thus were attributed more than once in the final totals. The most common themes among those who were unsure was the fear of side effects (56%), and the most common them among those not wanting the vaccine was the need for more information and research about the vaccine (47%).

Conclusions It is crucial to be aware of the parents’ concerns regarding the COVID-19 vaccine. Our study suggests that the most common reasons for hesitancy are fear regarding the potential side effects and the lack of adequate research and data. Knowledge about the determinants of hesitancy provides insight as to the most important topics to focus when educating families who struggle to make a decision. As the COVID-19 pandemic continues to evolve, larger studies are warranted to identify reasons for vaccine hesitancy in different populations.
### Abstracts

#### #393 STRATEGIES TO IMPROVE ALCOHOL USE DISORDER POST-DISCHARGE TREATMENT ENGAGEMENT IN MCCALL, IDAHO

N Massey*. University of Washington School of Medicine, Seattle, WA

**Purpose of Study** Discussions with local healthcare workers from McCall, a small resort town of 3,000 nestled in the mountains of central Idaho, regarding public health issues highlighted concerning high rates of alcohol use within the community. These observations were supported by the St. Luke’s 2019 Community Health Needs Report that reported 2% of the population engaged in binge drinking in 2016 compared to the national average of 16%. Further investigation highlighted the ways a culture of substance abuse, rising costs of living, and a significant shortage of treatment options contribute to high rates of alcohol use within McCall.

**Methods Used** An asset-based approach was selected to investigate possible interventions to address alcohol use disorder (AUD). Input from staff at St. Luke’s McCall pointed to Recovery Oriented Community (ROC), the only local resource helping individuals with AUD. ROC is led by a group of peer-recovery coaches (PRC), individuals with lived experience in recovery that are focused on providing recovery coaching and resource navigation. Initial discussions with ROC centered around existing ideas to begin introducing hospitalized patients with AUD to PRC. This concept aims to help patients connect with recovery resources after discharge. A literature review of available data was selected to begin investigating the viability and efficacy of such a system.

**Summary of Results** Research thus far has demonstrated the feasibility of the proposed system and shown encouraging results including improved post-discharge treatment engagement and lower rates of alcohol use following hospitalization. This information was delivered to ROC in an effort to bolster existing discussions with St. Luke’s to develop specific policies that incorporate PRC in hospital care for patients with AUD.

**Conclusions** The asset-based approach used for this project allowed opportunities to assist ROC through providing evidence and support for ongoing efforts to address AUD in McCall. Most meaningfully, this project was done in a practical and relatively expedient manner with little initial capital. Future work with ROC will focus on educating providers on the benefit of PRC, the development of a strong relationship between these providers and ROC, and ongoing research into the efficacy of this system.

#### #394 SUPPORT FOR PEOPLE WITH DEMENTIA AND THEIR CAREGIVERS IN RAVALLI COUNTY, MONTANA USING AN ASSET-BASED APPROACH

IM Starke*. University of Washington School of Medicine, Seattle, WA

**Purpose of Study** Ravalli County, Montana has an older population, with 26.5% being 65 or older compared to just 19.3% of the state of Montana. Hospice staff note that many people move to Ravalli County to retire for the rural lifestyle and outdoor recreation. In this study, an asset-based approach was used to identify a need in Ravalli County for more support for people with dementia and their caregivers. A review of literature found that Memory Cafes are a realistic intervention.

**Methods Used** The Ravalli County Health Assessment was reviewed to identify epidemiological factors unique to the county. Next, interviews with a primary caregiver and hospice staff were conducted to learn about specific needs. Additionally, an interview was conducted with the Ravalli County Council on Aging (RCCOA) to recognize existing assets in the community and where there is need for more growth. Finally, literature was reviewed to identify a realistic intervention.

**Summary of Results** A primary caregiver and hospice staff noted that there is a need for more support for caregivers of the elderly population. The director of RCCOA informed of a plethora of services they offer to the elderly and said there is a specific need for more support for people with dementia and their caregivers. They also noted that RCCOA has resources to expand services in the community. A qualitative study from London found that Memory Cafes, which are a weekly monthly gathering in the community for people with dementia and their caregivers, provides socialization, decreases isolation, normalizes living with dementia, and supports caregivers of people with dementia.

**Conclusions** People with dementia and their caregivers in Ravalli County, Montana could better be supported through the implementation of Memory Cafes. RCCOA has the resources and connections in the community to organize this service, making Memory Cafes a realistic intervention for Ravalli County. The asset-based approach was successful because it worked from the perspective of both community members and community organizations as well as the literature to identify a need and solution. Further research is needed to understand how to address barriers to implementing Memory Cafes, especially geographic barriers considering Ravalli County has a low population density.

#### #395 IMPROVING OUTREACH FOR UNHoused INDIVIDUALS EXPERIENCING OPIOID DEPENDENCE IN SHORELINE, WASHINGTON

AN Phan*. University of Washington, Seattle, WA

**Purpose of Study** Unhoused individuals are one of the communities most affected by substance use disorders, but few seek treatment due to stigma, negative attitudes about treatment, lack of motivation, or other challenges such as finances, treatment location, hours of operation of the clinics, etc. Shoreline, a city located in King County, Washington, has been identified by International Community Health Services (ICHS) as a community with a high large number of patients who suffer from opioid abuse. This report analyzes a plausible method for improving outreach for opioid abuse in unhoused individuals that can be implemented by community health partners in Shoreline.

**Methods Used** Therapeutic Health Services (THS), one of the community health partners addressing opioid abuse located in Shoreline, has made efforts in improving outreach by partnering with programs that house homeless individuals. Different outreach method studies were analyzed in this public health report and peer-recovery was chosen as the most potentially successful outreach method to implement into programs such as THS. One of the studies analyzed was an intervention
implemented as part of the Illinois Opioid State Targeted Response project. This study implemented those who were at least 1 year in stable recovery to contact populations that were considered ‘hidden’. The method used to determine efficacy in this study was determined by the percentage of those that showed up to a linkage meeting.

**Summary of Results** The results of the Illinois Opioid State Targeted Response project showed that of those who showed up, 92% agreed to have the linkage manager schedule them a treatment intake appointment and of those, 86% initiated treatment for an overall treatment initiation rate of 47%. This is significantly better than the overall treatment initiation rate of 28% found in prior SAMHSA-sponsored 12 site HIV risk reduction projects.

**Conclusions** Because of the work already underway to expand outreach, implementing peer recovery workers as part of outreach projects is something that can very realistically be done. Additionally, THS has already implemented peer-recovery albeit part of counseling rather than outreach, so this is a plausible method of expanding outreach. Next steps would include seeking out potential candidates and gathering resources needed for implementation of this program.

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**Purpose of Study** The COVID-19 pandemic affected healthcare communities from the institutional level to the personal level. Current research focuses on the experiences of physicians during the COVID-19 pandemic; however, few studies have investigated the professional and personal impact of COVID-19 on non-physician employees of healthcare centers. Additionally, many of the narratives surrounding the COVID-19 pandemic have been centered around large urban institutions; there has been little information discussing the impact of the pandemic on rural communities. The objective of this study is to gain insight into the perspectives of non-physician healthcare workers in a rural town and bring awareness to their experiences.

**Methods Used** A deductive qualitative study of semi-structured interviews was conducted with employees from a healthcare center in a rural town. Placement site was determined by University of Washington School of Medicine’s Office of Rural Programs. A total of five non-physician employees of Jefferson Healthcare Medical Center in Port Townsend, WA were interviewed. Interviews were conducted from June 2021 through August 2021. Interview questions were formed through multiple sessions with colleagues of the National Humanities Center and the University of Washington School of Medicine. As transcripts were reviewed, themes were identified and analyzed.

**Summary of Results** Analysis revealed the following themes: (1) impact of vaccination on personal and professional identities; (2) communal responsibility throughout the COVID-19 pandemic; and (3) consequences of institutionally implemented safety measures. Additional themes centered around the emotions related to COVID-19, change, and uncertainty.

**Conclusions** This study brought insight into the experiences of non-physician healthcare workers in a rural town during the COVID-19 pandemic. Utilizing open ended questions with focused listening can give a voice to those in the community. The themes identified demonstrate the importance of recognizing the diverse experiences of community members and the value of community in a rural environment. Lastly, there is much to be gained from using a narrative-based interview style within and outside research settings.

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**Neonatology general V**

**Concurrent session**

**8:00 AM**

**Saturday, January 22, 2022**

**#397 IMPROVING SCREENING, DIAGNOSIS, AND, MANAGEMENT OF TRANSIENT HYPOGLYCEMIA IN A LEVEL III NEONATAL INTENSIVE CARE UNIT**


**Purpose of Study** The physiologic transition from fetal to neonatal glucose homeostasis has a nadir by 2–4 hours of life before a self-recovery. There is emerging evidence that treatment of transient hypoglycemia with intravenous (IV) dextrose may negatively impact neurodevelopmental outcomes. We explored our neonatal intensive care unit (NICU) care practices and identified practice patterns that suggest an over-diagnosis and over-treatment of normal physiologic events. We aimed to modify practice patterns to optimize screening, diagnosis, and management of transient hypoglycemia.

**Methods Used** The quality improvement population was neonates born ≥35 weeks gestational age at the University of Utah NICU. Infants excluded are those with congenital anomalies, hypoxic-ischemic encephalopathy, mechanical ventilation, or vasopressor infusion. Interventions were obtaining the first blood glucose measurement at 2 hours of life, standardizing the blood glucose concentration for intervention to <45mg/dL, and providing the first enteral feed by 2 hours of life. Process measures were time to first blood glucose measurement, the number of infants receiving an IV dextrose infusion or bolus, time to first enteral feed, and total hypoglycemic events in the first 24 hours of life.

**Summary of Results** Pre-intervention n = 40 and post-intervention n = 153. Mean time from birth to first blood glucose measurement increased from 49.5 (SD ±17.8) to 122 minutes (SD ±37.1) (p < 0.001). IV dextrose infusion rate decreased from 97.5% of admissions to 41.2% (p < 0.01), with a decreased rate of boluses from 32.5% of patients to 5.8% (p < 0.001). Time to first enteral feed decreased significantly (per Xmr chart) from 15.8 hours of life to 3.2 hours of life (p < 0.001). Blood glucose measurements <45mg/dL in the first 24 hours of life had a non-statistically significant rise from 10.1% to 13.9%.

**Conclusions** Our practice pattern changes better identified hypoglycemia that persistent beyond the physiologic nadir

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Abstracts
while preventing treatment of normal physiologic events. This evidence shows that early enteral feeds over IV dextrose infusions can provide initial management of transient hypoglycemia without a significant rise in total hypoglycemic events within the first 24 hours of life.

**Abstracts**

INVESTIGATING RELATIONS BETWEEN THE NICU SPEECH ENVIRONMENT AND WEIGHT GAIN IN INFANTS BORN VERY PRETERM

M Chan Morales*, K Kumar, VA Marchman, M Scala, KE Travis. Stanford University, Stanford, CA

Purpose of Study Children born preterm, especially those born very preterm (<32 weeks gestational age, GA) are at risk for adverse neurodevelopmental outcomes, including language and related learning disorders. Adverse language outcomes in preterm children have been attributed, in part, to the minimal amounts speech exposure that neonates are exposed to while in the neonatal intensive care unit (NICU). Experimental studies that directly expose preterm infants to maternal speech sounds in the NICU find significant improvements in health factors relevant for neurodevelopment. However, few studies have examined whether natural variations in the speech environment of the NICU are related to short-term health outcomes in preterm infants. Such data are important for optimizing the speech environment of the NICU. The purpose of this study was to examine relations between the NICU speech environment and rate of weight gain during hospitalization, an important determinant of neurodevelopmental and physical health outcomes for preterm infants.

Methods Used Participants were infants born very preterm (<32 weeks GA in the NICU at Stanford’s Lucile Packard Children’s Hospital (n = 20). The auditory environment of each infant was assessed at 32–36-week post-menstrual age (PMA) using a speech-counting device known as a Starling, which continuously estimates the amount of speech spoken near to the infant per 5 minute interval. Speech rates were averaged for each infant over the entire 4-week period. Average rates of weight gain (Average grams/kilo/day) were ascertained via electronic medical chart review over the same period. Zero-order correlations and linear regressions controlling for the time infant spent in an incubator versus an open crib and starting weight at 32 weeks PMA were performed to examine associations between speech counts and average rates of weight gain.

Summary of Results Results indicated that infants who heard more speech, on average, gained weight significantly more quickly than those who heard less speech (r(20) = 0.64, p < .01). Importantly, speech rate accounted for more than 40% additional variance after covariates (r²-change = .43, F(1,16) = 13.6, p < .001). Analyses further exploring rates of visits and caloric intake as possible confounders did not reduce the size of the effects.

Conclusions The present findings suggest that enhancing the amount of speech exposure that preterm infants experience in the NICU may be beneficial for promoting physical growth, an important determinant of neurodevelopment. As such, NICU policies should be modified to increase speech exposure.

LIMITED ENGLISH PROFICIENCY AND HISPANIC VERY LOW BIRTHWEIGHT INFANT HEALTH OUTCOMES

1J Feister*, 2P Kan, 1H Lee. Lucille Salter Packard Children’s Hospital at Stanford, Palo Alto, CA; 2Stanford University, Stanford, CA

Purpose of Study Limited English proficiency (LEP) has been shown to impact patient care. There is a gap in knowledge on the impact of LEP on NICU patients. The primary aim of this study was a) characterize the prevalence of LEP in Hispanic families with VLBW infants and b) determine the association between LEP to various outcomes related to preterm birth.

Methods Used This was a cohort study comparing outcomes in patients with primarily Spanish-speaking families (a proxy for LEP) to patients with primarily English-speaking families. Data from the California Perinatal Quality Care Collaborative with linkage to Office of Statewide Health Planning and Development database were used. Inclusion criteria were VLBW infants (birthweight < 1500 g) of Hispanic ethnicity born at participant hospitals from 2009–2012 with documented primary language as either Spanish or English. Exclusion criteria were infants with major congenital anomalies or early mortality. Multivariable regression models accounting for infant factors (birthweight, gestational age) and maternal factors (age, parity, insurance status, and education) were created and odds ratios (OR) were estimated.

Summary of Results Of 7,020 infants who met inclusion criteria, 28% of families had LEP. Infants of families with LEP were more likely to receive human milk at discharge [crude OR 1.25 (95% CI 1.12–1.39), aOR 1.55 (1.36–1.77)] less likely to have necrotizing enterocolitis [crude OR 0.74 (0.58–0.94), aOR 0.62 (0.47–0.82)], and more likely to have a hospital readmission within the first year of life [crude OR 1.21 (1.09–1.34), aOR 1.00 (0.88–1.12)]. There were no significant differences in in-hospital mortality, need for oxygen at discharge, retinopathy of prematurity, prolonged length of stay, or high risk infant follow-up clinic referrals.

Conclusions LEP is common among Hispanic families in California with VLBW infants. Infants with LEP are more likely to receive breast milk at discharge and less likely to have NEC than those from families who identify English as their primary language. Families with LEP may be more likely to be recent immigrants, a group that is sometimes demonstrated to have better infant health outcomes than their native born counterparts. Infants of families with LEP were more likely to be readmitted to the hospital, suggesting that LEP may be a surrogate marker of re-admission risk, potentially highlighting communication barriers.

BONE QUALITY ASSESSMENT IN PRETERM INFANTS

A Tarell†, S Malone Jenkins, M Grinsoel, B Yoder. University of Utah Health Hospitals and Clinics, Salt Lake City, UT

Purpose of Study Diagnosis of metabolic bone disease (MBD) in preterm infants is challenging. The standard for diagnosis is dual energy x-ray absorptiometry (DXA) to measure bone mineral content (BMC), which has barriers limiting its use. Quantitative ultrasound (QUS) is an alternative mode of assessing bone status via measuring speed of sound (SOS).
SOS is determined by bone cortical thickness, density, microstructure, and elasticity. The purpose of this study is to determine the relationship between BMC, SOS, and growth parameters in preterm infants.

Methods Used In this observational prospective study, preterm infants born at ≤ 32 weeks gestational age (GA) or birth weight (BW) ≤ 1800g underwent both SOS and BMC measurements. BMC (Hologic) and SOS (Omnisense 8000P) were assessed ≤ 7 days of each other at either discharge or 40 weeks corrected GA (CGA). Absolute values were assessed for both tests as well as standardized z-scores for SOS. Pearson correlations ($R^2$) and linear regression were used to determine the relationship between demographics, growth parameters and the two modalities.

Summary of Results 41 preterm infants with a mean GA of 29.8 weeks (SD ± 2.2) and mean BW 1315 g (± 429) had bone quality assessment at a mean CGA 37.7 weeks (± 2.1). SOS was positively correlated with BW (R² 0.261, $p < 0.05$), BW (R² 0.090, $p = 0.057$), and birth head circumference (R² 0.101, $p < 0.05$). SOS was negatively correlated with CGA at the time of the test (R² 0.240, $p < 0.05$). BMC did not correlate with birth GA, BW, or head circumference. However, BMC positively correlated with CGA, weight, and length at the time of the test (R² 0.342, 0.794, 0.239 and $p < 0.05$ respectively). SOS and BMC were weakly inversely correlated (R² 0.163, $p < 0.05$).

Conclusions SOS and BMC poorly correlate to each other in preterm infants assessed close to term GA. SOS, but not BMC, correlated with birth GA and BW. SOS showed an inverse correlation with advancing postnatal age, with z-scores decreasing from birth to several weeks of age. BMC showed a positive correlation with advancing postnatal age. Given these findings, and the availability to perform repeated SOS at the bedside, we speculate that SOS may be a safer alternative and offer more information about bone health status in growing preterm infants compared to BMC. More research is needed to better understand the roles of SOS and BMC in the assessment of preterm infant bone health.

### #401 FACTORS ASSOCIATED WITH ABNORMAL BRAIN MAGNETIC RESONANCE IMAGING IN VERY LOW BIRTH WEIGHT INFANTS

NS Nanduri*, A Hisay, C Marquez, L Barton, M Binivale, R Ramanathan. Keck Hospital of USC, Los Angeles, CA 10.1136/jim-2022-WRMC.397

Purpose of Study Preterm very low birth weight infants (VLBW) are at greater risk of neurodevelopmental morbidity due to impaired brain development. Though it is clinically difficult to predict the outcome of these infants when discharged from the NICU, abnormal brain MRI findings may predict long-term consequences. Existing research shows very limited diagnostic testing done on these infants to assess neurodevelopment in the NICU. Term-equivalent MRI may indicate the need for further developmental assessments and close follow up. There is no consensus on how factors associated with various intensive care interventions link to abnormal MRI findings when these infants reach term gestation. This study aims to evaluate these factors associated with abnormal MRI findings in preterm VLBW infants.

### Abstract #401 Table 1 Factors associated with brain MRI abnormalities

<table>
<thead>
<tr>
<th>Brain MRI abnormal (%)</th>
<th>Brain MRI normal (%)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intubation in delivery room</td>
<td>40.7</td>
<td>59.3</td>
</tr>
<tr>
<td>Vaginal delivery</td>
<td>32.3</td>
<td>67.7</td>
</tr>
<tr>
<td>Highest pCO2 &gt;50 in first 24 hours of life</td>
<td>36.3</td>
<td>63.7</td>
</tr>
<tr>
<td>Invasive ventilation in first 24 hours</td>
<td>39</td>
<td>61</td>
</tr>
<tr>
<td>Invasive synchronized intermittent mandatory ventilation (SIMV)</td>
<td>42</td>
<td>57.9</td>
</tr>
<tr>
<td>High frequency oscillatory ventilation</td>
<td>30.9</td>
<td>69.1</td>
</tr>
<tr>
<td>High frequency jet ventilation</td>
<td>23.4</td>
<td>76.6</td>
</tr>
<tr>
<td>Patent ductus arteriosus</td>
<td>40.9</td>
<td>59.1</td>
</tr>
</tbody>
</table>

Methods Used Retrospective data of preterm VLBW infants (<1500 g) born between 2009 and 2021 at LAC + USC Medical Center was evaluated for MRI brain findings. Institutional IRB approval was obtained. An abnormal MRI was defined as abnormalities documented by pediatric neuroradiologists, including hemorrhage, periventricular leukomalacia, infarct or ischemia. Maternal factors, resuscitation at delivery, and early neonatal factors were analyzed for significant association.

Summary of Results Out of 356 infants matching inclusion criteria, 191 (52.3%) had abnormal brain MRIs. Lower gestational age ($p < 0.001$) correlated strongly with abnormal MRI. Infants delivered by spontaneous vaginal delivery as well as those intubated in the delivery room and with low 5 min Apgar scores were more likely to have abnormal MRI. VLBW infants with higher ventilation and oxygen requirements within the first 24 hours of life were at higher risk for developing abnormal MRI. Infants requiring invasive ventilation support beyond 24 hours of life were also more likely to have abnormal MRI, including those requiring high frequency oscillatory ventilation and jet ventilation. Incidence of abnormal MRI in infants with hemodynamically significant patent ductus arteriosus (PDA) was also significantly higher. These infants were also more likely to have initial low hematocrit ($p = 0.003$) and severe intraventricular hemorrhage on cranial ultrasound ($p = 0.016$). NICU duration of stay was longer for infants with abnormal MRI ($p < 0.001$). Over 50% of abnormal MRI findings were related to cerebellar abnormalities.

Conclusions VLBW infants with factors including need for intubation in the delivery room, higher ventilation and oxygen requirement in the first 24 hours of life, longer invasive ventilation, hemodynamically significant PDA, low hematocrit and abnormal cranial ultrasound predict abnormal brain MRI findings performed at term.

### #402 USE OF ELECTRONIC TEXT MESSAGING OF NRP PRINCIPLES TO PEDIATRIC TRAINEES & IMPACT ON RETENTION

1E Eskandar-Afsahi*, 1M Martinez Gomez, 1S Liu, 1K Tedesco, 2AM Yeh, 2R Ramanathan, 3M Binivale. 1LAC+USC Medical Center, LA, CA; 2USC Keck School of Medicine, LA, CA 10.1136/jim-2022-WRMC.398

Purpose of Study Pediatric trainees have decreased exposure to neonatal resuscitation. The purpose of this study was to...
provide distance learning via text messaging of NRP principles 8th edition and assessing retention and integration of NRP skills among pediatric residents.

**Methods Used** This is a prospective, randomized-controlled study including Pediatrics and Medicine-Pediatrics residents. An initial survey and quiz were completed to assess baseline knowledge and confidence in neonatal resuscitation. Twenty participants were included and randomized to control group, no text messages, and intervention group, receiving biweekly NRP content pearls (Image 1) for 6-weeks. Subsequently, there was a 6-week period of no contact. All participants completed a repeat quiz at 6-weeks and 12-weeks to assess their knowledge retention. Two high fidelity simulations were completed by all participants who were scored on skills, knowledge and teamwork using a validated tool.

**Summary of Results** Of the 20 study participants, 16 were Pediatric residents and 4 were Medicine-Pediatrics residents. Thirty-five percent were PGY-1, 40% were PGY-2 and 25% were PGY-3. Fifty percent had less than three resuscitations for term infants and 70% had less than three resuscitations for preterm infants in the last year. Majority agreed that they needed more knowledge and experience in neonatal resuscitation. Seventy percent stated that they not confident in leading a neonatal resuscitation. As for the initial quiz scores, Pediatric PGY-1, 2, and 3 scored 72.5%, 69.25%, and 69%, respectively. Medicine-Pediatrics PGY-1 and 2 scored 63.4% and 80%, respectively. Questions missed were about secondary apnea, pre-birth questionnaire, volume of intravenous/intravascular flush to be given and the chronologic sequence of transitional physiology after birth.

**Conclusions** Based on our initial results, it is evident that Pediatric trainees do not get enough experience, lack knowledge, and do not feel comfortable leading a neonatal resuscitation. With ongoing text messaging of NRP content pearls, we hope to show an increase in Pediatric trainees’ retention of knowledge and confidence level in caring for neonates requiring resuscitation at birth.

**Abstract #402**

**Figure 1** NRP Content pearl example

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**Abstract #403**

**Comparison of Procalcitonin and High Sensitivity C-Reactive Protein as Screening Tests in Neonates for Early Onset Sepsis and Late Onset Sepsis**

A. Vachhani*, A. Chambliss, M. Durand, R. Ramanathan, R. Cayabyab. LAC+USC Medical Center, Keck School of Medicine of USC, Los Angeles, CA

10.1136/jim-2022-WRMC.399

**Purpose of Study** Early detection of both early onset sepsis (EOS) and late onset sepsis (LOS) in newborns is difficult as the first signs of infection are often nonspecific. A laboratory test to rapidly identify these infants would add value in improving outcomes and limiting unnecessary antibiotic exposure. The purpose of this study was to compare the utility of procalcitonin (PCT) and high sensitivity CRP (hsCRP) as screening tests for EOS and LOS.

**Methods Used** Retrospective study of infants admitted to the neonatal intensive care unit from August 2018-February 2021. Infants with two paired PCT and hsCRP values obtained 12–24 hours apart and with blood, urine or cerebrospinal fluid culture results were included. EOS workup was performed at <=72 hours of life while LOS workup at >72 hours of life. Infants on antibiotics at the time of workup and those diagnosed with presumed infection, but negative cultures were excluded. Infants were classified as non-infected or infected (positive culture). Data were analyzed with Fisher-exact test or Chi Square and Wilcoxon rank sum where appropriate. Diagnostic value was determined by calculating sensitivity, specificity, positive predictive value (PPV) and negative predictive value (NPV) using cutoff of 10mg/L and 1.0ng/mL for hsCRP and PCT respectively.

**Summary of Results** A total of 359 infants received workups for EOS and 126 infants for LOS. There were 12 infants (3%) who had confirmed EOS while 21 (17%) with confirmed LOS. Demographics and outcomes of infants are shown in table 1. The sensitivity, specificity, PPV and NPV for serial measurements of PCT and hsCRP are shown in table 2. For EOS, subsequent measurements of both biomarkers led to...
an increase in sensitivity, but a decrease in PPV while in LOS sensitivity and PPV increased. The areas under the ROC curve were higher for both biomarkers for LOS in comparison to EOS (table 2).

Conclusions Our preliminary findings indicate that PCT and hsCRP may have better diagnostic value in patients for LOS in comparison to EOS. Previous studies suggest a natural rise in levels on serial measurements in the first 72 hours in all neonates influencing its diagnostic value and presence of partum and maternal risk factors can also influence levels after birth in the absence of infection.

Neonatology – perinatal biology II
Concurrent session
8:00 AM
Saturday, January 22, 2022

#404 MULTIOMICS LONGITUDINAL MODELING OF PREECLAMPTIC PREGNANCIES

C Espinosa, I Maric, K Contrepois, M Moufarenj, IS Stelzer, D Feyaerts, X Han, A Tang, RJ Wong, GL Darmstadt, JD Winn, GM Shaw, DA Relman, SR Quake, MS Angst, M Snyder, DK Stevenson, B Gaudilliére, N Aghaeepour. Stanford University School of Medicine, Stanford, CA; Stanford University, Stanford, CA; University of the Pacific – San Francisco Campus, San Francisco, CA.

Purpose of Study Preeclampsia is a complex disease of pregnancy of unclear pathophysiology. Specific complex changes in a woman’s physiology precede diagnosis of preeclampsia. Understanding these changes at different levels of biology can be enabled by simultaneous application of multiple assays. We developed prediction models for preeclampsia risk by analyzing six omics datasets from a longitudinal cohort of pregnant women.

Methods Used We performed a multiomics analysis of the transcriptome, proteome, metabolome, lipidome, and microbiome from blood, urine and vaginal samples collected longitudinally during pregnancy at the Lucile Packard Children’s Hospital at Stanford University. Our discovery and validation cohort included 33 and 16 women, respectively. Prediction models were built for each omics set using the elastic net machine learning method and their integration was performed using stacked regression. Immune system mass cytometry features and available clinical variables were subsequently integrated with the model.

Summary of Results The multiomics model predicting which women will develop preeclampsia had high accuracy (area under the receiver operating characteristics curve (AUC) of 0.94; 95% confidence interval (CI): [0.90, 0.99]). A prediction model using only ten urine metabolites provided an accuracy equivalent to that of the model using the complete metabolomic dataset and was validated using an independent cohort (AUC=0.87; 95% CI: [0.76, 0.99]). Univariate analysis further confirmed statistical significance of proteins and metabolites chosen by the prediction model. Several proteins identified as biomarkers in our model have previously been associated with preeclampsia (e.g., LEP, VEGF-A) further validating our findings. Integration with clinical variables further improved prediction accuracy of the urine metabolome model (AUC=0.90; 95% CI: [0.80, 0.99], validated). Several biological pathways were identified to be associated with preeclampsia. Integration with the immune system data confirmed known pathological alterations associated with preeclampsia and suggested novel associations between the immune and proteomic dynamics.

Conclusions While further validation in larger populations is necessary, these encouraging results will serve as a basis for a simple, early diagnostic test for preeclampsia.
DISCOVERY AND VERIFICATION OF EXTRACELLULAR miRNA BIOMARKERS FOR NON-INVASIVE PREDICTION OF PREECLAMPSIA IN ASYMPTOMATIC WOMEN

N Do*. University of California San Diego, La Jolla, CA

Purpose of Study Preeclampsia is a pregnancy complication characterized by new-onset high blood pressure and protein in the urine. It is the most common cause of preterm birth and the second most common cause of maternal mortality. For patients with risk factors such as chronic hypertension, diabetes, or prior affected pregnancy, daily low dose aspirin administered between 12-28 weeks of gestation has been shown to decrease preeclampsia risk. However, many patients that develop preeclampsia do not have known risk factors. Thus, effective and non-invasive screening for high-risk pregnancies may be clinically useful. This study explores extracellular miRNAs (exRNAs) in blood and other biofluids as identifiers of preeclampsia risk. In a previous study, we identified candidate miRNA biomarkers for the prediction of preeclampsia. Our aim is to validate these biomarkers of interest in an independent set of patients in a larger cohort.

Methods Used To perform this validation, we isolated the exRNA from maternal serum from a new cohort of subjects using the Plasma/Serum Circulating and Exosomal RNA Purification Kit (Shurry format, Norgen Biotek). We then used small RNA sequencing to quantify the RNA in those samples. We will also use qPCR to specifically quantify the two miRNAs comprising our top bivariate candidate biomarker, hsa-miR-155-5p/hsa-miR-153-5p, which we showed in our original study to be correlated with preeclampsia risk.

Summary of Results exRNA of adequate quality was extracted from the maternal serum samples. The acquired sequencing data showed adequate quality for the large majority of exRNA samples, as measured by several metrics, including: total sequencing reads; percentage of miRNAs as compared to all RNA biotypes in the sample; and miRNA complexity, which indicates the number of detected distinct miRNA sequences in each sample.

Conclusions Going forward, we will perform data analysis to determine whether the candidate miRNAs from the original cohort are confirmed in the validation cohort. If the performance of the biomarkers of interest is validated, this will support their further development as clinical biomarkers for prediction of preeclampsia.

BREASTFEEDING DIFFICULTY INCREASES DEPRESSIVE SYMPTOMS AND IMPAIRS MATERNAL-INFANT BONDING: A CONTENT ANALYSIS OF MATERNAL BREASTFEEDING NARRATIVES

1EA Wright*, 1A Mehta, 1AL Nelson, 2H Stohl, 3M Economidis. 1Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA; 2Harbor-UCLA Medical Center, Torrance, CA

Purpose of Study Protection against postpartum depression (PPD) is a commonly cited maternal benefit of breastfeeding1. However, one study found that negative breastfeeding experiences may actually increase depressive symptoms instead of reducing them. The purpose of this study is to evaluate women’s self-reported associations between breastfeeding and PPD.

Methods Used An electronic search was conducted on Reddit including original user posts with keywords ‘breastfeeding’ and ‘depression’. Relevant posts and corresponding comments (n=584) were extracted from February 2013 to July 2021 and subject to qualitative content analysis.

Summary of Results Narratives most frequently described difficulty breastfeeding as the cause of new onset mental health symptoms (33%); however, mothers breastfeeding without difficulty were also susceptible (20%). Among mothers reporting exacerbated pre-existing mental health symptoms, a similar greater proportion described difficulty breastfeeding (17%), but still others experienced no difficulty (6%). Common words included in these posts were ‘guilt’ (17%), ‘fail’ (14%), ‘pressure’ (7%), ‘shame’ (6%), and ‘alone’ (6%), highlighting the emotional impact of breastfeeding difficulty. Only 3% of posts described breastfeeding as improving mental health symptoms. Other themes yielded from this study included weaning-induced mental health symptoms (17%) and decreased milk supply due to mental health symptoms (2%). A concerning large proportion of posts described a lack of knowledge (23%). Finally, among posts mentioning bonding (n=99), 58% reported that breastfeeding inhibited infant bonding, versus only 1% enhanced bonding—contradicting the prevailing idea that breastfeeding stimulates bonding2. After switching to formula, 63% reported increased bonding, 8% no change, and 6% decreased bonding.

Conclusions Mothers struggling to breastfeed may not experience the classically described benefits of breastfeeding, instead exhibiting increased depressive and other mental health symptoms and impaired maternal-infant bonding. These findings highlight the need for enhanced education on breastfeeding and mental health relationships, and adjustment of practices to meet the diversity of maternal experiences.

MULTIOMICS MODELING OF PRETERM BIRTH IN LOW- AND MIDDLE-INCOME COUNTRIES

CE Espinosa Bernal*, J Maric, DK Stevenson, N Aghaeepour. Stanford University School of Medicine, Stanford, CA

Purpose of Study Preterm birth (PTB) is the leading cause of death in children under five years of age across the globe.
Abstract #408 Table 1  Study demographics

<table>
<thead>
<tr>
<th>Race, Race group</th>
<th>GDM group (n, % of total GDM group)</th>
<th>Control group (n, % of total control group)</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>White, non-Hispanic</td>
<td>9 (41%)</td>
<td>81 (48%)</td>
<td>90</td>
</tr>
<tr>
<td>Hispanic</td>
<td>4 (18%)</td>
<td>33 (20%)</td>
<td>37</td>
</tr>
<tr>
<td>Asian</td>
<td>9 (41%)</td>
<td>42 (25%)</td>
<td>51</td>
</tr>
<tr>
<td>Black or African American</td>
<td>0</td>
<td>12 (7%)</td>
<td>12</td>
</tr>
<tr>
<td>American Indian or Alaska</td>
<td>0</td>
<td>1 (1%)</td>
<td>1</td>
</tr>
<tr>
<td>Native</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>22</td>
<td>169</td>
<td>191</td>
</tr>
</tbody>
</table>

**Abstract**

**Purpose of Study** Gestational diabetes (GDM) is a risk factor for the development of cardiovascular and metabolic disease in both mothers and their offspring, and central obesity is associated with the development of GDM. Detection of GDM in early gestation may improve health outcomes of at-risk mothers and babies. Our group and others have developed in-vivo magnetic resonance imaging (MRI) modalities suitable for monitoring pregnancies during early gestation. We hypothesized that early gestation MRI will predict the subsequent development of GDM.

**Methods**

200 women were recruited in the first trimester of pregnancy. We recorded the pregnancy course through four visits (at 11–14 weeks, 19–22 weeks, 36 weeks, and at delivery), and conducted chart reviews to gather clinical outcomes on the mother-baby dyad. Two MRIs were prospectively performed on each subject, at 14–16 weeks and 19–22 weeks. Using imaging software, placental volume was obtained by tracing placental regions of interest in three axes. Fat content was measured from the L1 through L5 vertebrae. Subcutaneous fat area ratio (SFAR) was derived by assessing the subcutaneous fat area normalized to the whole trunk area; visceral fat area ratio (VFAR) was derived by assessing the visceral fat area normalized to the whole visceral area.

**Summary of Results** An epidemiological model for the prediction of PTB achieved moderate accuracy (AUROC=0.70), highlighting the difficulty of the predictive task. Machine learning-based multiomic models had strong performance for the prediction of multiple relevant objectives, including time to delivery (Pearson’s R=0.64) and maternal covariates such as age (R=0.57), gravidity (R=0.53), and body mass index (BMI, R=0.80). The biological signature for time to delivery included fetal-associated proteins such as alpha-fetoprotein and immune proteins like PD-L1. Maternal age had a strong negative correlation with type IX collagen conserved across our multinational cohort. Gravidity had an age-independent association with levels of endothelial NOS and inflammatory chemokine CXCL13, an observation further validated on an independent Stanford cohort (n=17). BMI positively correlated with adipocyte hormone leptin and structural fatty acid protein FabP4.

**Conclusions** These results simultaneously identify novel signatures for maternal covariates impacting PTB and shed light on potential biological and socioeconomic interventions that generalize across multiple populations.

**#408 EARLY PREDICTION OF GESTATIONAL DIABETES MELLITUS BY IN-VIVO MAGNETIC RESONANCE IMAGING**

B Lee*, K Sung, C Jianzen, S Vangala, SU Devaskar. University of California Los Angeles, Los Angeles, CA

10.1136/jim-2022-WRMC.404

**Case report II**

**Concurrent session**

10:15 AM

**Saturday, January 22, 2022**

**#409 PERIANAL PSEUDOVVERRUCOUS PAPULES AND NODULES PRESENTING AS SUSPECTED CHILD ABUSE**

JP Wang*, K Shea. LAC+USC Medical Center/USC, Los Angeles, CA

10.1136/jim-2022-WRMC.405
Case Report Due to their sensitive location, perianal lesions often alert clinicians to potential sexual abuse. Perianal pseudoverrucous papules and nodules (PPPN), first described in patients with urostomies in 1979 under the name chronic papillomatous dermatitis, are often mistaken for condyloma acuminata due to their wart-like appearance. Early recognition and treatment of PPPN would lessen parental distress and avoid unnecessary workup for children who suffer from this illness.

Case Description The subject is a four-year-old Hispanic male who was admitted to the pediatric ward for perianal lesions, rectal pain, and itching. The patient’s lesions first appeared 8 months prior to this admission, gradually becoming larger and more numerous. The lesions were preceded by a history of constipation, minimally responsive to Miralax, and subsequent frequent and loose stools with stool-holding behaviors.

The subject was then prescribed topical mupirocin, bacitracin, lidocaine, and oral Keflex at multiple outside ED visits.

The patient was referred for inpatient management of his lesions by DCFS. He was diagnosed with severe constipation and encopresis. Dermatology clinically diagnosed PPPN without a biopsy. He underwent bowel disimpaction and was placed on an aggressive bowel regimen.

Patient was discharged on hydrocortisone, ketoconazole, and zinc oxide for four months with significant improvement of lesions. Patient was toilet trained six months after discharge with complete resolution of lesions.

Conclusion PPPN present with multiple well-demarcated and dome-shaped papules due to constant irritants. They have smooth surfaces and are about 2–10 mm in diameter. Their friable textures have a tendency to ulcerate and are prone to secondary infections.

Definitive diagnosis is made by skin biopsy, useful in cases with challenging clinical course that mimic other conditions. However, most cases are diagnosed clinically.

Abstract #409 Figure 1 Skin exam showed multiple 3–8 mm papules and nodules at the perianal area

The mainstay of treatment targets the inciting factors, including diaper use, encopresis, and colostomy bag. Therapy should be aimed to restore the skin barrier, eliminate any source of irritations, and treat superimposed infections.

#410 LATE PRETERM INFANT WITH POSTNATAL DIAGNOSIS OF RENAL TUBULAR DYSGENESIS

S Gaffar*, P Aora, R Ramarathnam. 1Los Angeles County University of Southern California Medical Center, Los Angeles, CA; 2Children’s Hospital of Los Angeles, Los Angeles, CA

10.1136/jim-2022-WRMC.406

Case Report A 2025 gram male infant was born to a 23-year-old multiparous woman by cesarean section for breech presentation at 34 gestational weeks. Rupture of membranes occurred 65 hours prior, with meconium-stained fluid. Pregnancy was complicated by limited prenatal care and polysubstance abuse (tetrahydrocannabinol and methamphetamine). Serologic testing was unremarkable except for unknown gonorrhea, chlamydia, and group B streptococcus.

He was intubated and given surfactant soon after delivery for labored breathing. Chest radiographs revealed pneumothoraces requiring bilateral chest tube placement. Empiric antibiotics were started as his clinical condition deteriorated with escalating ventilator requirements and hypotension requiring multiple inotropes. He became oliguric with hyponatremia (126 mEq/L), elevated creatinine (3.78 mg/dL), and non-cystic parenchymal disease on renal ultrasound. Renal failure was managed with strict fluid balance per nephrology, 1 week later, peritoneal dialysis commenced. Subsequent whole exome sequencing identified two ACE gene mutations, confirming renal tubular dysgenesis (RTD).

Since the patient was not considered a transplant candidate, a supportive care plan was formulated to optimize future transplant candidacy. Presently, he continues to receive peritoneal dialysis, gastrostomy tube feeds, and developmental therapy.

Discussion RTD is acquired in an autosomal recessive manner or from exposure to substances causing renal hyperperfusion. Immunohistochemistry detects epithelial membrane antigen but not CD-10, illustrating absence of proximal tubules. This adds to known glomerular and ductal hypoplasia. By these mechanisms, antenatal reduction of glomerular filtration manifests as intrauterine oligohydramnios.

Another hallmark of RTD is refractory hypotension arising from renin-angiotensin dysfunction. Fresh frozen plasma, intravenous hydrocortisone or fludrocortisone, and vasopressin infusion provide transient circulatory support. Glucocorticoids stimulate hepatic transcription of the renal tubular growth factor angiotensinogen. This degree of postnatal hypotension supports the postulation that antenatal hydrocortisone may improve neonatal outcomes.

RTD is a rare disease that starts in fetal life with defective tubule development. Perinatal oligohydramnios evolves into acute renal insufficiency, respiratory failure, and refractory hypotension in the neonatal period. Prenatal molecular genetic testing means that RTD is no longer relegated to exclusive postmortem diagnosis and may have a higher incidence than we think.

Abstract #411 A RARE CASE OF COFFIN-SIRIS SYNDROME CAUSED BY SOX4 MUTATION WITH NEW PHENOTYPIC FEATURES

1,2AE Lee*, 1,2J Friedman, 1,3J Lenberg, 1,2K Wigby. 1Rady Children’s Hospital San Diego, San Diego, CA; 2University of California San Diego, La Jolla, CA; 3Rady Children’s Institute for Genomic Medicine, San Diego, CA

10.1136/jim-2022-WRMC.407
Case Report

De novo missense variants involving the HMG domain of SOX4 were recently reported to cause a Coffin-Siris like syndrome characterized by global developmental delays, distinctive craniofacial features, hypotonia and 5th finger clinodactyly (OMIM # 184430). To date, only four cases have been reported in the literature (Zawerton et al, 2019). We report on a case of a young girl with global developmental delay, hypotonia, distinctive facies, cardiac anomalies, sensorineural hearing loss secondary to hypoplastic cochlear nerves and striking stereotypes, who was identified on whole genome sequencing to have a novel heterozygous de novo variant in SOX4 (c. 199A>G, p.Met67Val). The proband was the product of a 30 2/7 week uncomplicated pregnancy. At birth she was noted to have a peri-membranous VSD. At eight months of age she was markedly hypotonic and poorly responsive to sound, and was found to have underdeveloped cochlear nerves. She presented to our genetics service at 19 months of age where she was noted to have pre-auricular pits, small stature, and severe developmental delay. Family history was non-contributory. Chromosomal microarray showed only 24 Mb of homozygosity consistent with distant consanguinity. A congenital hypotonia panel was non-diagnostic. Duol whole genome sequencing with maternal sample identified the SOX4 variant, later confirmed to be de novo. Coffin-Siris syndrome is a rare pattern of malformation characterized by growth deficiency, microcephaly, coarse facial features, intellectual disability and hypoplastic fifth finger and toenails. Of those, the majority are caused by ARID1B, which is a gene encoding the subunits of the SWI/SNF complex which is essential in chromatin remodeling. Interestingly, SOX11 is another known cause of Coffin-Siris. SOX11, together with SOX4 and SOX12 and forms the SOXC complex, with highly conserved identical DNA-binding domains, giving insight to the mechanism of pathology which we elaborate on. Individuals with SOX4 variants appear to have relatively mild dysmorphology including anteverted nares, wide mouth with a cupid bow, and posteriorly rotated ears. Our proband extends this phenotype with unique features of hypoplastic cochlear nerves and severe stereotypes which we hypothesize are related to her severe hearing loss.

#412 PERITONEAL TUBERCULOSIS MIMICKING OVARIAN CANCER

1A Cox, 2C D’Assumpcao*, 3A Froush, 1A Heidari, 2AA Ramzan. 1Ross University School of Medicine, Miramar, FL; 2Kem Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.408

Case Report

Diagnosis of peritoneal tuberculosis may be challenging. This form of extrapulmonary infection may present as abdominal and/or pelvic masses with pain, bloating, and discomfort. The clinical manifestations are similar to ovarian cancer, including elevation in tumor marker CA-125, peritoneal seeding and lymphadenopathy. Multiple diagnostic procedures are sometimes needed to make a definitive diagnosis.

Methods Used

Retrospective case review following IRB approval.

Summary of Results

A 50-year-old Hispanic woman presented with generalized abdominal discomfort, bloating, fatigue, unintentional weight loss and night sweats. CT imaging revealed an 11 x 9 x 9 cm cystic pelvic mass associated with large volume loculated ascites, diffuse lymphadenopathy and peritoneal implants. She had an elevated CA-125 of 583 units/mL. Paracentesis was non-diagnostic. Intrapерitoneal biopsies revealed diffuse granulomatous inflammation with foreign body-like multinucleated cells and no evidence of malignancy. Acid-fast stain and acid-fast bacilli (AFB) smear and culture were negative. Chest x-ray showed bilateral hilar adenopathy and a granuloma in left upper lung, with subsequent positive QuantiFERON gold. Antituberculosis therapy was initiated with Isoniazid, rifampin, ethambutol, and pyrazinamide (RIFPE). After 5 months of RIFE therapy, the CA-125 normalized and there was dramatic radiographic improvement in lymphadenopathy and peritoneal disease burden. The pelvic mass persisted and she underwent bilateral salpingo-oophorectomy. Histopathology resulted as a serous cystoadenoma with necrotizing granulomatous inflammation. The AFB smear and mycobacterium tuberculosis complex PCR were positive. The patient recovered post-operatively and was re-initiated on RIFE therapy. Mycobacterial culture and sensitivity results are pending.

Conclusion

Peritoneal tuberculosis can be mistaken for ovarian cancer and thorough diagnostic evaluation is essential to ensure appropriate management and treatment.

#413 CLOSTRIDIUM PARAPUTRIFICUM IN 46-YEAR-OLD MALE WITH LIVER DISEASE

1C D’Assumpcao*, 1K Dao, 1A Heidari. 1Kem Medical Center, Bakersfield, CA; 2Ross University School of Medicine, Miramar, FL

10.1136/jim-2022-WRMC.409

Case Report

Clostridium paraputrificum is an anaerobic spore forming gram positive bacilli that is part of normal human gastrointestinal flora. Less than 1% of all clostridium infections in the literature are C. paraputrificum. We report a 46 year old male with alcoholic liver disease and lower gastrointestinal hemorrhage who was found to have C. paraputrificum bacteremia.

Method

Retrospective case report.

Case Presentation

A 46-year-old male with heavy alcohol abuse, hypertension and congestive heart failure presented with two weeks of worsening abdominal pain that progressed to bright red blood per rectum. He was found to be in septic shock at presentation to the emergency department. He received aggressive fluid resuscitation and vasopressor support. He was diagnosed with liver cirrhosis complicated by splenomegaly and ascites shortly after admission. Ascitic fluid analysis supported diagnosis of spontaneous bacterial peritonitis. He was started in ceftriaxone. He eventually developed acute oliguric kidney injury due to shock and hepatorenal syndrome. He was placed on intermittent dialysis. Admission blood cultures grew C. paraputrificum. Metronidazole was added. Upper and lower endoscopy found large internal hemorrhoids vessels associated with rectal varices requiring banding. He continued to requiring blood transfusions daily. He was a poor candidate for colorectal surgery and outpatient dialysis. Patient and family ultimately agreed with hospice care.

Conclusion

C. paraputrificum is a rarely reported cause of anaerobic septicemia. Translocation from the gastrointestinal tract is the usual path and mucosal damage should be investigated. Cirrhosis portends a poorer prognosis.
THE NEUROSYPHILIS AND COCCIDIOIDOMYCOSIS CONUNDRUM


10.1136/jim-2022-WRMC.410

Case Report Neurosyphilis is an infection of the central nervous system caused by Treponema pallidum that can occur following initial infection. Early forms of neurosyphilis affect cerebrospinal fluid, meninges, and vasculature while latter forms affect the brain and spinal cord parenchyma. We describe a case of neurosyphilis complicated by fluconazole toxicity during treatment of pulmonary coccidioidomycosis. A 26-year-old man with poorly controlled type 1 diabetes mellitus, miliary pulmonary coccidioidomycosis, heart failure with reduced ejection fraction, left eye cranial nerve III palsy and right cranial nerve VII palsy presented to the emergency department for weakness and syncope. Physical exam was remarkable for equal pupils unreactive to light or accommodation bilaterally, point tenderness of bilateral iliac crest, and left CNIII and right CNVII palsies. MRI brain with and without contrast showed slightly diffuse increased enhancement. CXR showed left peri hilar and right upper lobe infiltrates. Transesophageal echocardiogram showed an ejection fraction of 30%. Lumbar puncture (LP) performed showed WBC 5, RBC 0, glucose 48, protein 254 with cocci IgG WR CF 1:4 and an opening pressure of 14cm. Due to suprapharmacologic fluconazole level in the setting of disseminated cocci, fluconazole was discontinued and Cremesene was started with noted improvement of weakness. Neurology recommended repeat lumbar puncture with flow cytometry and cytology for carcinomatosis meningitis, oligoclonal bands and IgG synthesis rate in addition to RPR, VLDR and FTA-ABS. Repeat LP showed WBC 3, RBC 2, glucose 78 and protein 135.9 with opening pressure 30cm of H2O. Labs showed nonreactive RPR and VDRL with reactive FTA-ABS and positive syphilis antibodies. Furthermore, cerebrospinal fluid studies showed albuminocytologic dissociation consistent with neurosyphilis. A diagnosis of late latent syphilis with bilateral posterior subcapsular cataract, and CNIII and CNVII palsy was made and the patient was then started on Penicillin G for treatment of neurosyphilis. Neurosyphilis caused by T. pallidum is rare due to spontaneous resolution in cases without an inflammatory response. This case demonstrates a unique case of late latent syphilis consistent with neurosyphilis due to history of bilateral posterior subcapsular cataract at age 18 with cranial nerves III and VII palsy.

A CASE OF RAOUTTELLA PLANTICOLA BACTEREMIA IN AN IMMUNOCOMPROMISED MALE

1. Garcia *, 1J Obemdorf, 1J Tsai, 1N Mangat, 1A Heidari, 5S Mishra, 5K Radicic. Rosal University School of Medicine, Miramar, FL; 1American University of the Caribbean School of Medicine BV, Cupecoy, Sint Maarten (Dutch part); 1Kern Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.411

Case Report Raoutella Planticola formerly called Klebsiella planticola is a gram-negative aerobic rod that belongs to the Enterobacteriaceae family and is often established in aquatic habitat and soil. The published cases infected with R. planticola are scarce. It rarely causes infection in an immunocompetent host. In contrast, in immunocompromised patients, inoculation of R. planticola can surpass innate and adaptive host response leading to bacteremia. Here we describe a case of R. planticola bacteremia in a patient on chemotherapy who presented with fever with a recent dog bite to his right lower extremity as a potential source.

Methods A single patient case report was conducted after IRB approval.

Case Presentation A 41-year-old male with right knee osteosarcoma who was receiving infusions of Cyclophosphamide and Mesna presented to the emergency department (ED) with fever and chills for 3 days. He was bitten by his friend’s dog on the medial aspect of his right posterior calf resulting in a bleeding wound one month prior while working in the garden. One day later, he developed regional swelling and pain associated with fever which progressively worsened for almost two weeks, followed by complete self-resolution. While in ED, he was tachycardic and febrile to 39.4 °C. Imaging was remarkable. The site of the dog bite was well healed and had no tenderness, fluctuance, erythema, or warmth. Blood cultures were drawn from peripheral and his chemotherapy port and he was started on broad-spectrum antibiotics with vancomycin and cefepime. After two days, the blood culture from peripheral grew gram negative rods resembling enterics and cefepime was replaced by meropenem. The patient became afebrile. Blood cultures identified as R. planticola, sensitive to ceftriaxone. Antibiotic therapy was switched to ceftriaxone on hospital day 4. Repeat blood cultures were negative and the patient remained afebrile for greater than 24 hours. He was discharged with at-home infusions with Ceftriaxone for 14 days.

Conclusion R. planticola is an emerging infection, particularly in oncology patients. Clinicians should be aware and include it in their differential diagnosis of any unidentified gram negative infection in immunocompromised host as failure to treat in a timely manner could result in a fatal outcome.

Healthcare delivery research IV

Concurrent session

10:15 AM

Saturday, January 22, 2022

APPLICATION OF EXPLAINABLE MACHINE LEARNING IN ANALYZING SAFETY AND QUALITY REPORTS FROM A CLINICAL UNIT


10.1136/jim-2022-WRMC.412

Purpose of Study Accurate assessment of safety-related reports in the healthcare setting is essential for improving care quality, optimizing workflow, and preventing future safety incidents. At present such systems rely on a manual review process, which can be very time-consuming and difficult to standardize. The purpose of this study is to develop an explainable machine learning (ML) model for automatically identifying the severity of incident reports and streamlining quality improvement processes. We hypothesized that high severity incident reports can be automatically identified using ML methods.
Methods Used This study utilizes 7,095 operational quality reports collected in the radiation oncology department at the University of Washington from 2012–2021. Each report was assigned a near-miss risk index score (0–4) as part of ongoing clinical practice, rather than a separate annotation process. For this experimentation, the reports were assigned binary labels of high-severity (score of 0–2) and low-severity (score of 3–4), and the data was split into a training, validation, and test set with a 70–15–15 ratio. We chose Random Forest as the pilot model because this model has historically performed well in text classification tasks. The input features to the Random Forest model were term frequency-inverse document frequency vectors, where each dimension denotes the importance of a word. Model performance was evaluated using the area under the receiver operating characteristic curve (AUC), which integrates the diagnostic ability of a binary classifier at all operating points, and F1, which is the harmonic mean of precision and recall. We then used the popular SHapley Additive exPlanations (SHAP) package to understand which words in the safety reports are most indicative of report severity.

Summary of Results Our Random Forest model achieved an AUC of 0.66 and F1 of 0.42 on the withheld test set. The dataset was annotated over a 9-year period by different clinicians as a part of clinical workflow, resulting in increased variance in scoring that negatively impacts training and prediction performance. Our results compare favorably to the manual rating process where the interrater agreement of this dataset was 37.6% (Mullen et al, 2015). The SHAP interpretability analysis indicates that the top 10 most predictive words are ‘field’, ‘shift’, ‘use’, ‘treatment’, ‘therapist’, ‘schedule’, ‘isocent’, ‘catch’, ‘wrong’, and ‘cm’, in descending order of importance.

Conclusions This study shows that ML is a viable method to analyze and score operational quality reports in the clinical healthcare setting. We present a pilot prediction model with an AUC of 0.66 and F1 of 0.42, proving that a suitable detection alert tool to triage safety-related reports for timely review. To our knowledge, this approach has not been attempted to classify reports’ severity and fills a critical unmet need as a quality improvement tool in healthcare.

#418 DIAGNOSING MELANOMA AND METASTASES USING DEEP NEURAL NETWORKS, ARTIFICIAL INTELLIGENCE, AND MEDICAL IMAGING

JH Miao*. Cornell University, New York, NY
10.1136/jim-2022-WRMC.414

Purpose of Study Cancer is one of the most common causes of death globally, impacting millions of patients worldwide. Melanomas cause a large number of deaths because of metastases. Early and accurate cancer screening is important to improve patient outcomes and potentially save lives. Detecting melanoma and metastases with high accuracy and precision is thus essential. Computer-aided detection (CAD) systems have been developed around the world to help aid healthcare professionals. In this research, a computer-aided diagnosis using artificial intelligence and medical imaging is applied to create a machine learning model to evaluate the presence of melanoma and metastases in patients.

Methods Used In this research, a deep convolutional neural network model was designed using artificial intelligence and developed to help detect the presence of melanoma in imaging scans. The machine learning model was applied to a clinical patient dataset containing over 2000 patients. Data from 60% of the patients was used to train the machine learning model, and the remaining 40% of the patient data was used for testing performance.

Summary of Results The deep learning model achieved an 83.8% sensitivity and 84.3% specificity in detecting melanoma cancer and metastases in patients.

Conclusions Therefore, computer-aided diagnoses using artificial intelligence can help detect melanoma cancer and metastases in radiologic imaging to aid patient outcomes. Especially in areas where there are fewer healthcare resources and medical professionals present, the machine learning model can help aid melanoma cancer diagnosis in underserved populations.

#419 FINANCIAL ASSESSMENT OF A HYBRID HUMAN AND ARTIFICIAL INTELLIGENCE WORKFLOW FOR FOLLOW-UP RECOMMENDATIONS FROM RADIOLOGY

10.1136/jim-2022-WRMC.415

Purpose of Study The purpose of this study was to assess the financial costs associated with a hybrid human and artificial intelligence (AI) workflow for follow-up recommendations from radiology and pathology and to determine the cost savings potential provided by AI as an additional layer of review. This study compares the costs of different work processes used in our small community hospital and evaluates the cost implications of a hybrid human and AI workflow for follow-up recommendations from radiology and pathology.

Methods Used We performed a cost analysis to compare the costs of different work processes used in our small community hospital and evaluate the cost implications of a hybrid human and AI workflow for follow-up recommendations from radiology and pathology. The study was conducted at a community hospital in California, USA. The hospital's radiology department performs a large number of imaging procedures, including CT scans, MRIs, and X-rays. The hospital's pathology department performs a wide range of diagnostic and research procedures, including histology, cytology, and molecular testing. The study involved a comprehensive review of the hospital's financial records and a comparative analysis of the costs of different work processes. The analysis included a comparison of the costs of different work processes, such as traditional human review, AI review, and a hybrid human and AI workflow. The study also involved a review of the hospital's financial policies and a comparison of the costs of different work processes, such as traditional human review, AI review, and a hybrid human and AI workflow. The study was conducted at a community hospital in California, USA. The hospital's radiology department performs a large number of imaging procedures, including CT scans, MRIs, and X-rays. The hospital's pathology department performs a wide range of diagnostic and research procedures, including histology, cytology, and molecular testing. The study involved a comprehensive review of the hospital's financial records and a comparative analysis of the costs of different work processes. The analysis included a comparison of the costs of different work processes, such as traditional human review, AI review, and a hybrid human and AI workflow. The study also involved a review of the hospital's financial policies and a comparison of the costs of different work processes, such as traditional human review, AI review, and a hybrid human and AI workflow.
Purpose of Study Many radiology interpretations indicate the need for follow-up imaging, and adherence to these recommendations is often incomplete. Lack of follow up on these recommendations may result in poor patient outcomes. At our institution we deploy a hybrid workflow utilizing a combination of natural language processing (NLP) software to tag radiology exams and a quality and safety nurse to handle subsequent communication and documentation with referring medical providers or patients. In this study, we assess the financial costs and benefits of our institution’s hybrid workflow to determine the overall financial feasibility of this system.

Methods Used After Institutional Review Board approval, the total number of radiology scans with follow-up imaging recommendations between February 1, 2020 and January 1, 2021 were pulled for inclusion in this study. Mammography patients were excluded due to a separate, predefined screening protocol. Reports flagged as ‘overdue’ by the mPower Follow-Up Recommendation Algorithm (Nuance Communications Inc., Burlington, MA) and/or the quality and safety nurse were included to determine revenue. Revenue was calculated based on the Medicare 2020 and 2021 values for patients who had overdue clinically indicated follow-up exams, whose ordering providers were notified by the nurse, and who then completed the necessary imaging. Personnel cost of a quality and safety nurse was calculated by determining the percentage of time the nurse spent on follow-ups and multiplying this number by the average national salary of a quality and safety nurse.

Summary of Results A total of 3,011 patients were flagged as overdue. After careful review of the charts, 327 were missing orders for follow up recommended imaging. Of these 84 patients completed follow up outside of our institution or declined follow up. Quality nurse interventions led to 244 patients completing their follow-up imaging at our institution. The follow-up imaging of these patients accounted for $35,106.90 of revenue, which represents a conservative, generalizable value based on the 2020 and 2021 Medicare national rates. The quality and safety nurse spent about 60% of her time on follow-ups and therefore, personnel cost was estimated at $26,326.80.

Conclusions Our cost and revenue estimates are based off of national averages and as such will vary significantly by location. However, our analysis suggests that a hybrid human and NLP workflow for follow-up recommendations from radiology would most likely not cause financial losses for institutions. Additionally, this workflow acts as a safety net to ensure patients receive adequate follow up imaging and markedly mitigates risks from litigation.
Lauderdale and the Broward County Housing Authority, housing authority residents, in Broward County, the 17th largest county in the US by population, residents were surveyed regarding their preferences of accessing care through telemedicine (phones, tablets, or computers) or through face-to-face interactions. We performed secondary data analysis of survey data collected by the YMCA of South Florida. This analysis involved descriptive statistics for describing the respondents and their preferences, Chi-square for detecting significant differences between males and females for preferring to utilize telemedicine rather than face-to-face exchanges with respondents’ health care providers, and logistic regression to determine the odds of males’ or females’ likelihood of preferring accessing care via telemedicine.

Summary of Results Twenty-five percent (25.70%, n=140) of respondents preferred to visit their physician through the use of telemedicine rather than attending in-person. A significantly higher percentage of males (37.5%, n=40) reported a preference for utilizing telemedicine than females (21.0%, n=100) ($X^2[1, n=140] = 4.07, p = .04$). Males were found to have had two times the odds of preferring telemedicine (aOR 2.26, 95% CI 1.01–5.03) as compared to females.

Conclusions These findings could shape the way clinical healthcare providers offer telemedicine services to their patients. Since diverse housing authority resident males have a preference towards telemedicine, it is worthwhile that health care providers serving this vulnerable population during COVID-19 offer options for patients to access care through telemedicine. Future research can be done to assess the reason for this gender difference in the preference for telemedicine, as well as to understand if there are specific platforms patients prefer when accessing telemedicine (phone, computer, etc.).

#422 A REVIEW OF THE LEVEL OF EVIDENCE IN POINT OF CARE ULTRASOUND WITHIN THE PICU

A Walls*, A Willyerd, E Su, Phoenix Children’s Hospital, Phoenix, AZ; The University of Texas Health Science Center at Houston John P and Katherine G McGovern Medical School, Houston, TX

10.1136/jim-2022-WRMC.418

Purpose of Study To review the last decade of point of care ultrasound research and determine the level of evidence provided as it pertains to the pediatric intensive care unit.

Methods Used A retrospective review of the current literature was performed utilizing the following publicly available databases: Pubmed, Scopus and Cochrane Databases regarding bedside point of care ultrasound in the pediatric population. Each data source was queried for the following inclusion criteria terms: ‘pediatric’, ‘ultrasound’, ‘bedside’, ‘intensive care’ and ‘critical care’. Data was collected from 2011 to 2020. Each article was subcategorized into: Case Report, Case Series, Cohort Study, Systematic Review, Meta-Analysis and Randomized Controlled Trial and provided a level of evidence (LOE) based on the respective study design. Exclusion criteria included studies performed in the neonatal intensive care unit, studies that included adults and anesthetics regional pain investigatipns that involved point of care ultrasound. Statistical analysis was performed on Minitab 5.0 (State College, PA).

Summary of Results A total of 122 publications met our inclusion criteria and 31 met exclusion criteria. The overall LOE in the pediatric ultrasound literature did not improve significantly from 2011 to 2020 (P=0.321) as per ANOVA calculations. However, the overall year over year number of pediatric publications increased each year except for 2015 and 2017 (0%, -35%), respectively. The average LOE increased each year except for 2013, 2017 and 2018. The most frequent publication type was found to be the retrospective cohort study with a frequency of 31 out of the 121 (26%), reported studies. There were 11 Randomized Control Trials, 14 Systematic Reviews and Meta-Analysis which accounted for Level 1 and Level 2 evidence, respectively. The year over year change in Level 1 and Level 2 evidence did not significantly change over the study period (P=0.643).

Conclusions The literature regarding the use of point of care ultrasound in the critical care setting remains limited. Our investigation reveals no statistically significant improvement in the LOE over the past decade. Despite this trend, there is obvious increased interest in point of care ultrasound, as the sheer volume of articles tended to increase year over year during the study period. Our community should continue with investigations which fall into higher LOE categories in order to aid in clinical decision making at bedside and assist in the creation of practice guidelines.

Hematology and oncology III

Concurrent session

10:15 AM

Saturday, January 22, 2022

#423 SICKLE CELL PAIN CRISIS COMPLICATED BY OPIOID INDUCED HYPERALGESIA, TREATED WITH LOW DOSE KETAMINE INFUSION

F Venter*, R Dunn, J Bhandohal, S Mishra, E Cobos. Kiem Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.419

Case Report PURPOSE OF STUDY: Sickle Cell Disease (SCD) can present with Acute Vaso-occlusive Crisis (VOC) and neuropathic pain, involving central sensitization, peripheral injury, and hyperalgesia. Management of VOC is challenging and requires IV fluids, oxygen and significant doses of intravenous (IV) opioids. However, this can result in opioid induced hyperalgesia (OIH). Few papers have explored the use of low dose Ketamine to manage pain during VOC. It could be considered as an adjunct analgesic agent on the use of low-dose ketamine to manage pain during the critical care setting. We present here a case management of a 36-year-old female with SCD in pain, worsened with increasing doses of IV opioids. OIH was suspected and treated with a low dose ketamine infusion resulting in a dramatic resolution of her pain.

Methods This is a single case review after IRB approval.

Summary of Results A 36-year-old African American female with lifelong SS genotype SCD was admitted for VOC worsening over 4 days and unalleviated by her prescribed breakthrough pain medications. Treatment was started with fluids and increasing doses of IV opioids. By day 5, the pain became unbearable, suggesting OIH in the setting of concomitant VOC. She was treated with low dose Ketamine infusion at 0.1 mg/kg/hr, increased to 0.2 mg/kg/hr with dramatic resolution of her pain. Once IV ketamine and opioids were discontinued, patient’s pain was comfortably managed only on oral maintenance MS Contin for the following 48 hours.
Conclusion OIH is postulated to develop when the central glutamate transporter system is inhibited, increasing the amount of glutamate available to excitatory N-Methyl-D-aspartic acid (NMDA) receptors. This allows for cross talk of the neural mechanisms for pain and tolerance. Morphine targets NMDA receptors. When administered, it is neurotoxic and causes apoptotic cell death of the dorsal horn, thus sensitizing the neurons. Ketamine is a very potent inhibitor of NMDA receptors and can block the neurotoxic properties of morphine. After ketamine is discontinued, the patient does not revert back to their OIH state, thus allowing for the safe return to longer-acting opioids. Few publications have reported on the use of low-dose ketamine to manage pain during VOC. It could be considered as an adjunct analgesic agent during VOC episodes in patients that endorse persistent severe pain despite receiving high-dose opioid therapy.

#424 IMMUNE THROMBOCYTOPENIA FOLLOWING COVID-19 VACCINE
S Prasad*, M Adebayo, R Jariwal, G Petersen, E Cobos. Kern Medical Center, Bakersfield, CA
10.1136/jim-2022-WRMC.420

Purpose of Study Several vaccines have been developed and are being administered against severe acute respiratory syndrome coronavirus 2. Common side effects include fever, chills, headache, myalgia, and soreness at the injection site. However, some rare adverse effects have also been reported. Cases of thrombocytopenia following COVID-19 vaccination have been reported since the initiation of mass vaccinations. We present a case of thrombocytopenia presenting with petechiae and mucosal bleeding which developed as an adverse response after first-dose administration of the Moderna COVID-19 vaccine and was refractory to first-line therapy. Treatment options for refractory thrombocytopenia are discussed.

Methods Used Retrospective case study.

Summary of Results A 58-year-old Hispanic male presented to our hospital for acute onset of mucosal bleeding, petechiae and easy bruising. He denied any previous history of spontaneous bleeding or easy bruising. Patient had received his first dose of the Moderna COVID-19 vaccine three weeks prior to the development of spontaneous bleeding. Physical examination was notable for diffuse petechiae along the arms, legs, and abdomen along with numerous oral lesions and gingival bleeding. He was noted to have a platelet count of 3 x 10^9/L with all other cell lines within normal limits. After a comprehensive and exhaustive workup of all well-known precipitants of immune thrombocytopenic purpura returned negative, a diagnosis of idiopathic thrombocytopenic purpura was made. Patient was treated with first-line therapy of platelet transfusion, high dose steroids, and intravenous immunoglobulin (IVIG) multiple times but platelet count remained refractory. We then treated the patient with romiplostim—a thrombopoietin receptor agonist—and fostamatinib—an inhibitor of spleen tyrosine kinase. The platelet count improved and his symptoms were resolved.

Conclusions Since the mechanism of COVID-19 vaccine induced thrombocytopenia is unclear, it is important to acknowledge that some patients may not respond to the standard accepted treatment with steroids and IVIG. Cases similar to ours may benefit from therapy with romiplostim and fostamatinib when first-line therapy is unsuccessful.

#425 CASE SERIES OF DIFFUSE LARGE B CELL LYMPHOMA PRESENTING AS PELVIC MASS
R Sharma*, A Heidari, E Cobos, G Petersen, AA Ramzan, R Polineri. Kern Medical Center, Bakersfield, CA
10.1136/jim-2022-WRMC.421

Purpose of Study Primary diffuse large B-cell lymphoma (DLBCL) of the pelvis is rare with a non-specific clinical presentation. Although management of this tumor is standardized with high curative rates, the prognosis depends on timely and accurate diagnosis and therapy. Presenting with non-specific clinical symptoms, a multidisciplinary approach is recommended to diagnose and treat this disease. In this case series we present two patients that presented to our institution with primary DLBCL of the pelvis initially suspected as gynecologic malignancy.

Methods Used This study was approved by the Institutional Review Board of Kern Medical. A retrospective review of both the patient’s record was performed. Literature search was conducted on PubMed and Google Scholar. The following search terms were applied: diffuse large B-cell lymphoma of the pelvis, pelvic tumors, R-CHOP.

Summary of Results A 27-year-old woman presented with intractable abdominal pain, distention and unintentional weight loss. She had large bilateral ovarian masses, peritoneal carcinomatosis and metastatic adenopathy. A CT guided biopsy of a peritoneal implant was performed. A 58-year-old woman presented with a large pre-sacral mass and inguinal adenopathy. An inguinal lymphadenectomy was performed. Pathology for both patients revealed large B-cells positive for CD45+, CD20+, TCL1+, CD4+, consistent with diffuse large b cell lymphoma (DLBCL). Both patients were treated with systemic chemoimmunotherapy with monoclonal antibody against CD20 (rituximab) in combination with Cyclophosphamide, Doxorubicin, Vincristine, Prednisone (R-CHOP). One patient completed therapy and currently has no evidence of disease and the other patient has been receiving therapy.

Conclusions Even though the incidence of DLBCL is on the rise, pelvis as the primary location remains rare. Given a high rate of cure and entirely different approach to treatment for lymphoma as compared to gynecological malignancies it is important to remember this condition in the differential diagnosis. A multidisciplinary approach between gynecology oncology, medical oncology and pathology would result in early diagnosis treatment and cure.

#426 EXTREMEDULLARY MYELOMA OF LIVER MASQUERAADING AS GALLBLADDER CARCINOMA
S Kolagatla*, F Smith, M Julapelly, J Fiercy, N Moka. Appalachian Regional Healthcare, Lexington, KY; 2Harley Medical Center, Flint, MI
10.1136/jim-2022-WRMC.422

Case Report Multiple myeloma (MM) is a malignant clonal proliferation of plasma cells in the bone marrow. Extramedullary myeloma (EMM) refers to soft tissue or visceral proliferation of clonal plasma cells can occur approximately in 15% of
patients with MM. Organs involved by EMM may include skin, central nervous system, and viscera. Our case is EMM initially diagnosed as a gallbladder carcinoma.

A 76-year-old male found to have IgG Kappa MM with normal cytogenetics and 40% involvement of bone marrow. Offered transplant after CR from bortezomib (V), dexamethasone (D) and thalidomide but patient declined. Subsequent relapse treated with carfilzomib and later daratumumab regimen.

Kappa light chains (K) increased but bone marrow negative. PET and MRI of liver with contrast confirmed liver mass and retroperitoneal lymphadenopathy but the gall bladder not visualized. The liver mass was suspected to be a gallbladder carcinoma.

CT guided biopsy of the liver mass showed proliferation of plasma cells. Given the rapid progression of K received V, D, cyclophosphamide and etoposide without cisplatin.

Initial regression of the liver lesions and K but immediate worsening. Developed rapidly evolving non-painful, firm cutaneous nodules over his anterior upper abdomen and chest (figure 1B). Skin lesion biopsy showed plasma cells. He is currently on elotuzumab V and D, but his kappa light chains continue to rise.

EMM suspected in patients with MM who have localized to a particular organ or Light chain escape. EMM can mimic a primary malignancy such as gallbladder carcinoma as in this case. EMM is aggressive with <6 month survival. Number or prior lines of therapy are associated with higher risk of extramedullary relapse. Lenolidamide and prior allogenic transplant lowered the risk of EMM. No prospective studies defining an optimal treatment strategy for extramedullary myeloma. Vemurafenib for BRAF mutation or immune therapies with CAR-T cells have shown promising results in a limited number of EMM patients.
and procedures are helpful. MRI and CT scans provide information regarding possible etiologies and, therefore, direct the approach to a tissue diagnosis. Tissue evaluation with immunohistochemistry remains the mainstay in the diagnosis of lymphomas. Our patient’s immunohistochemistry was positive for tumor markers discovered in only a subset of tumors, directing towards treatment with combination chemotherapy instead of local measures (e.g., radiation therapy, percutaneous vascular procedures).

Therapeutic maneuvers for clinically stable patients such as head elevation can reduce hydrostatic pressure and relieve edema. For symptomatic patients, radiation therapy and intravascular stent placement are recommended. Nonetheless, initial management should be led by the severity of symptoms and the underlying malignancy. The primary therapy for T-cell lymphoblastic lymphoma is hyper-CVAD chemotherapy, which our patient received.

While the severity of symptoms differs widely, SVC syndrome may be fatal. Given the sporadic presentation and aggressive nature of T-cell lymphoblastic lymphoma, early recognition and identifying when to intervene are critical factors for positive patient outcome.

### Abstracts

#### #428 VINBLASTINE INDUCED PARALYTIC ILEUS IN A YOUNG FEMALE WITH HODGKIN LYMPHOMA

G Malolot, P Chan, H Aboaid, N Raza, R Polineri. Ross University School of Medicine, Miramar, FL; Kern Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.424

Purpose of Study A rare presentation of paralytic ileus induced from Vinblastine treatment.

Methods Used Retrospective Study

Summary of Results 31-year-old female with Hodgkin lymphoma stage II presented with progressive diffuse non-radiating abdominal pain, nausea, and vomiting. Patient’s symptoms started a few days after initiation of second cycle of ABVD (Adriamycin, Bleomycin, Vinblastine, Dacarbazine) therapy. Laboratory studies were remarkable for anemia and leukopenia. Abdominal X-ray was significant for gaseous distention mostly prominent in the colon just beneath the left hemidiaphragm. Further imaging studies with Computed tomography (CT) abdomen and pelvis revealed colonic gaseous distention with moderate residual fecal material and moderate small bowel fluid-filled distention. Given high suspicion for paralytic ileus induced by Vinblastine, the patient was treated with prokinetic agent metoclopramide, a stimulant laxative, and vinblastine was continued. Patients symptoms resolved with conservative management.

Conclusions Paralytic ileus is a common clinical condition that is associated with autonomic neuropathy subsequently leading to decreased bowel peristalsis. Most are often caused postoperatively but a small niche of the population undergoing chemotherapy treatment with vinca alkaloids vincristine and vinblastine are susceptible with this condition. Toxicity profile of Vinblastine affecting the gastrointestinal system seems to be dose-related.

Metabolism of Vinblastine is processed by the hepatic cytochrome P450 3A. Majority of the pharmacokinetic interactions are due to previously administered drugs that are metabolized same as Vinblastine. After coadministration effects can lead to potent enzyme inducers or inhibitors. There are abundant drugs out in the market that can interfere with Vinblastine metabolism which can potentially aggregate increasing drug-related toxicity.

The importance of this case illustrates that early recognition of Vinblastine induced paralytic ileus can lead to better outcomes and shorter hospital courses as treatment as mentioned above is different based on the cause of paralytic ileus.

#### #429 CARCINOSARCOMA OF DUODENUM AND PANCREASES SUCCESSFULLY TREATED BY WHIPPLE’S PROCEDURE

1P Chan, G Malolot, N Raza, R Polineri. Ross University School of Medicine, Miramar, FL; Kern Medical Center, Bakersfield, CA

10.1136/jim-2022-WRMC.425

Purpose of Study A rare presentation of carcinosarcoma of duodenum and pancreas treated with Whipple procedure and adjuvant chemotherapy.

Methods Used Retrospective Study

Summary of Results 53-year-old female incidentally found to be anemic with hemoglobin of 7.7 g/dL and liver function test which showed alkaline phosphatase of 679 unit/L, aspartate transaminase of 129 unit/L, alanine transaminase of 220 unit/L, and bilirubin of 1.6 mg/dL on regular lab workup ordered by surgeon prior to elective lipoma removal. Due to anemia, transaminitis, and hyperbilirubinemia, computed tomography (CT) scan of abdomen and pelvis were performed which was remarkable for a 5 centimeter ulcerative mass in the second part of duodenum without evidence of metastasis. An Esophagogastroduodenoscopy (EGD) was performed which revealed non-obstructive circumferential ulcerated mass with biopsy finding may represent a sarcomatoid carcinoma or carcinosarcoma. Malignant cells stain positive with AE-1/AE-3, CD10, CA 19-9, and CK7. Patient acutely developed worsening abdominal pain and was hospitalized with gastric outlet obstruction and underwent Whipple’s procedure. Patient had a carcinosarcoma with 60% sarcomatous component on 40% adenocarcinoma component in the primary tumor. However, there were 8 out of 43 lymph nodes involved predominantly with adenocarcinoma. Final pathology was indicative for staged III carcinosarcoma of the duodenum and pancreas requiring adjuvant chemotherapy of Folfirinox for six months. Patients’ follow up imaging were unremarkable and the patient has no evidence of disease.

Conclusions Carcinosarcomas are a rare malignant tumors that consists of a mixture of two components, the carcinomatous and sarcomatous elements. Commonest sites include uterus, head and neck and are extremely rare in the duodenum. Localized tumors located in the region that can potentially obstruct gastric outlet flow such as in this case ought to be resected. Whipple’s procedure has shown an efficacious role in the treatment plan. Although, given that there are no direct guidelines on treatment for such a rare carcinosarcoma, this case highlights the importance of timely diagnosis, resection, and adjuvant therapy considering there is a high risk for recurrence.
Immuno-logy and rheumatology II
Concurrent session
10:15 AM
Saturday, January 22, 2022

#430 B-CELL INTERFERON-β CORRELATES WITH LUPUS NEPHRITIS IN SYSTEMIC LUPUS ERYTHEMATOSUS

1FK Alduraiabi*, 2H Fatima, 1W Chatham, 1H Hu, 1,2J Mountz. The University of Alabama at Birmingham, Birmingham, AL; 1VA Clinic Birmingham, Birmingham, AL.

10.1136/jim-2022-WRMC.426

Purpose of Study Early diagnosis of lupus nephritis (LN) can be challenging since some patients do not exhibit overt clinical manifestations until advanced stages. B cell interferon-beta (IFNβ) correlates with development of B cell autoimmune phenotype. The objective of the present study is to determine if elevated IFNβ in circulating B cells can be a useful indicator for the development of more severe histopathologic features of LN.

Methods Used Flow cytometry was used to quantitate intracellular IFNβ in naïve (IgD+CD27−) CD19+ B-cells in the peripheral blood mononuclear cells (PBMCs) of a cross-sectional cohort (N=80) of patients with systemic lupus erythematosus (SLE), 33 of whom had lupus nephritis. Serologic and clinical manifestations of LN included anti-DNA, anti-Sm, C3, C4, and urine protein/creatinine ratio were determined. The correlation of B-cell IFNβ with lupus nephritis classification and histopathological findings, light, electron microscopy, and immunofluorescence (IF) for deposition of IgM, IgG, IgA, C1q, and C3 was determined in 23 of the 33 patients for whom renal biopsy data was available.

Summary of Results LN was identified in 41% of our cohort of 80 SLE patients. Naïve B-cell IFNβ was positively associated with the development of LN but not cutaneous disease. Higher levels of B-cell IFNβ also correlated with higher levels of circulating anti-dsDNA, anti-Sm, and the urinary protein/creatinine ratio. Biopsy examination revealed that proliferative LN lesions (Class III, IV with or without V) characterized by significantly elevated endocapillary hypercellularity, fibrous crescent, and fibrocellular crescent were significantly associated with high B-cell IFNβ. Surprisingly, IgG, IgA, IgM, C3, and C1q deposition in the kidney was not correlated with B-cell IFNβ.

Conclusions Our results suggest that B-cell IFNβ can be used in combination with other clinical diagnostic markers to assist in identifying patients who are at high risk of developing advanced LN.

#431 IDENTIFICATION OF MESENTERIC LYMPH NODE COMMENSALS INVOLVED IN NEONATAL CD4 T CELL ACTIVATION

1RE Temme*, 2J Schwensen, 3M Shenoy, 4M Koch. 1University of Washington School of Medicine, Seattle, WA; 2Fred Hutchinson Cancer Research Center, Seattle, WA

10.1136/jim-2022-WRMC.427

Purpose of Study Maternal antibodies have an important role in neonatal immune development, particularly in regard to gut-associated lymphoid tissue and the microbiota. Neonatal mice born to antibody-deficient (uMT−/−) mothers show both an elevation in CD4 T-cell activation and increase in microbial growth in mesenteric lymph nodes (mLN) compared to pups born to wild type mothers. These results are supported by experimental data from germ-free mice that show significantly lower CD4 activation. It is therefore hypothesized the microbiota is responsible for stimulating this observed immune response. The goal of this project was to identify differences in commensals colonizing mLN of pups born to uMT−/− mothers versus antibody-sufficient mothers (uMT+) in order to find a candidate microbe that could be eliciting the observed CD4 T cell expansion phenotype. This is important because understanding the mechanism in which adaptive immune responses work in response to microbiota can be used to develop more effective immune-modulating therapeutic interventions.

Methods Used We compared neonatal mice born to uMT−/− dams and uMT± dams; all pups are able to produce antibodies but the uMT−/− pups did not receive antibodies via breast milk from their mothers. At day 21 of age, the mLN were harvested, homogenized, and placed in an anaerobic chamber. Each mLN was divided equally and plated on two selective growth media. When colonies formed, they were individually restreaked on the same media, then underwent 16S PCR amplification, DNA purification, and sent for sequencing. The resulting sequences were matched to bacterial species via 16S sequence identity using BLAST and the bacterial specimens were frozen for future use.

Summary of Results Sequencing revealed 60 possible bacterial species in both groups. Further broken down, 34 (57%) of these species were shared between uMT−/− and uMT±, while 21 (34%) were unique to uMT−/− and 5 (8%) were unique to uMT±. Bacterial species in the shared group fell largely into the Staphylococcus and Lactobacillus genera with a few in the Parabacteroides genus. In the uMT−/− unique group, Bifidobacterium, Escherichia, Muribaculum/Duncaniella, Bacteroides, and Streptococcus genera were identified. The uMT± unique bacteria were in the Enterococcus genus.

Conclusions Results from this project along with previous data support that receiving antibodies during the neonatal interval does affect microbial diversity, with the majority of unique bacterial species growing from uMT−/− mLN. It’s possible one of these unique species is responsible for the CD4 T cell expansion phenotype observed in past experiments but further investigation is required to determine the mechanism behind how the microbiota is eliciting this T cell response.

#432 RHEUMATOID ARTHRITIS AMONG MINERS IN NEW MEXICO, UTAH, AND COLORADO


10.1136/jim-2022-WRMC.428

Purpose of Study Our previous population-based studies have shown that coal miners in Appalachia have a 2–3-fold increased odds of rheumatoid arthritis (RA), consistent with the association between RA and silica reported in other settings. We wished to ascertain if RA risk was similarly elevated in mineral extraction workers in the western US with likely
Abstract 

#432 Table 1  
Rheumatoid arthritis (RA) associated with silica exposure: multivariate logistic regression including smoking, race/ethnicity, and age

<table>
<thead>
<tr>
<th>RA Definition (models exclude RA by other definition only)</th>
<th>Underground hard rock mining</th>
<th>Underground soft rock mining (including coal)</th>
<th>Surface mining/ore processing</th>
<th>Silica from non-mining sources</th>
</tr>
</thead>
<tbody>
<tr>
<td>RA + glucocorticoids (model n = 1165)</td>
<td>2.97 (1.40, 5.12)</td>
<td>8.80 (3.55, 21.83)</td>
<td>4.34 (2.47, 5.97)</td>
<td>3.15 (1.73, 5.76)</td>
</tr>
<tr>
<td>RA + DMARDs (model n=1170)</td>
<td>6.33 (3.01, 13.39)</td>
<td>4.47 (1.59, 12.57)</td>
<td>2.46 (1.24, 4.93)</td>
<td>2.91 (1.73, 4.88)</td>
</tr>
</tbody>
</table>

Cells are odds ratios (95% confidence intervals). Referent=no silica exposure. DMARD = disease modifying anti-rheumatic drug. 

#433 LIPOSOMAL FORMULATION OF REDUCED GLUTATHIONE DECREASES BACTERIAL LOAD AND ALTERS THE IMMUNE RESPONSE IN MYCOBACTERIUM TUBERCULOSIS INFECTION

1N Kachour, 1A Beever, 1J Owens, 2R Cao, 3A Kalloli, 8R Kumar, 1K Sasania, 1C Vaughn, 1M Singh, 2E Truong, 2C Khattachourian, 1C Sidjian, 1K Zakery, 1W Khamas, 2S Subbian, 1, 2Venketaraman. 1Western University of Health Sciences, Graduate College of Biomedical Sciences, Pomona, CA; 2Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA; 3Rutgers University Public Health Research Center at New Jersey Medical School, Newark, NJ; 4Western University of Health Sciences College of Veterinary Medicine, Pomona, CA.

Purpose of Study Reduced form of Glutathione (GSH) has direct antimycobacterial activity at physiological concentrations and inhibits the growth of Mycobacterium tuberculosis (Mt) within monocytes. This experiment aims to elucidate the effects of a liposomal formulation of GSH (L-GSH) on the levels of free radicals, inflammatory cytokines, and granuloma formation.

Methods Used C57BL6 (WT) mice (11 males and 11 females) were infected with approximately 1000 CFU of H37Rv strain of Mt. Treated groups included mice treated with either 40mM L-GSH (40LGSH) or 80mM L-GSH (80LGSH) administered via drinking water. Control groups were administered mock treatment in the form of plain drinking water.

Summary of Results Mt infection increased free radical production as measured by Malondialdehyde (MDA) and diminished the levels of reduced and total forms of GSH in the lungs. Treatment with 40LGSH resulted in a significant increase in the levels of both reduced form of glutathione at 2 weeks, 4 weeks and 8 weeks post-infection and total glutathione at 4 weeks and 8 weeks post-infection in lung lysates. 40LGSH treatment showed significant increase in the levels of reduced and total forms of glutathione in the plasma at 4 weeks post-infection. 80LGSH treatment significantly increased the levels of reduced and total forms of glutathione in the lung lysates at 8 weeks post-infection. Both treatments resulted in a reduction of MDA levels in lung tissue and plasma. L-GSH treatments reduced the lung area involved in granuloma formation as well as granuloma size and complexity. 40LGSH treatment decreased the bacterial burden in the lungs significantly at 4 weeks post-infection, in the spleen at 2, 4 and 8 weeks post-infection and in the liver at 4-weeks post-infection. Both treatments significantly decreased the levels of IL-6 at 4 weeks post-infection and IL-10 at 8 weeks post-infection, and significantly increased the levels of IL-2 at 2- and 8-weeks post-infection in lung lysates. 40LGSH treatment significantly increased the levels of Interferon gamma at 2-, 4- and 8-weeks post-infection in lung lysates.

Conclusions 40LGSH was more effective in increasing the levels of reduced and total forms of GSH in the lung and plasma. L-GSH treatment altered cytokine profile and reduced bacterial load, granulomatous response, and free radical levels in mice. Thus, L-GSH may be implicated as an adjuvant to standard therapy in those infected with Mt.
Abstract #434 Table 1  Patient population, N=23

<table>
<thead>
<tr>
<th>Sex</th>
<th>Female 19 (83%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Race</td>
<td>Hispanic 11 (48%)</td>
</tr>
<tr>
<td>Asian/Pacific Islander</td>
<td>1 (4%)</td>
</tr>
<tr>
<td>Black</td>
<td>3 (13%)</td>
</tr>
<tr>
<td>White, Non-Hispanic</td>
<td>6 (26%)</td>
</tr>
<tr>
<td>Other</td>
<td>2 (9%)</td>
</tr>
<tr>
<td>Insurance</td>
<td>Medicaid 13 (56%)</td>
</tr>
<tr>
<td></td>
<td>Other 2 (9%)</td>
</tr>
<tr>
<td></td>
<td>Commercial 8 (35%)</td>
</tr>
<tr>
<td>Rheumatology Diagnosis</td>
<td>JIA 13 (56%)</td>
</tr>
<tr>
<td></td>
<td>Spondylarthrtis 3 (13%)</td>
</tr>
<tr>
<td></td>
<td>SLE 3 (13%)</td>
</tr>
<tr>
<td></td>
<td>Vasculitis 2 (9%)</td>
</tr>
<tr>
<td></td>
<td>JDM 1 (4%)</td>
</tr>
<tr>
<td></td>
<td>Other 3 (13%)</td>
</tr>
<tr>
<td>Rheumatology Medications</td>
<td>Biologics 11 (48%)</td>
</tr>
<tr>
<td></td>
<td>JAK 2 (9%)</td>
</tr>
<tr>
<td></td>
<td>MTX/LEF 11 (48%)</td>
</tr>
<tr>
<td></td>
<td>MMF/AZA 5 (22%)</td>
</tr>
<tr>
<td></td>
<td>HCQ 3 (13%)</td>
</tr>
<tr>
<td></td>
<td>Steroids 6 (26%)</td>
</tr>
</tbody>
</table>

Methods Used This is a descriptive cohort with a plan to transfer from pediatric to adult rheumatology within the University of Colorado. Collaborating pediatric and adult rheumatologists created a transfer log to streamline tracking of patients and information sharing. Patients were retrospectively identified for inclusion over a 14 month period. We captured demographics, rheumatologic disease, disease activity, medications, hospital use, insurance status, and compliance with care plans. Data was collected by chart review and managed using REDCap.

Summary of Results Twenty-three patients were identified with plans to transfer to adult rheumatology (table 1). 70% completed a first visit in adult rheumatology with an average of 137 days from the last pediatric rheumatology visit. 30% had their first adult visit within 4 months of their last pediatric visit. In subgroup analysis, patients who did not transfer within 4 months waited a mean of 346 days before the first adult visit. Five patients never established with adult rheumatology and had a longer interval between their last 2 pediatric rheumatology visits compared to those who transferred (313 ± 136 days vs. 170.3 ± 155 days, p=0.07). Disease complexity, number of medications, and disease activity were similar between groups.

Conclusions We examined the outcomes of 23 pediatric patients with a plan to transfer care to an adult rheumatology clinic. Interestingly, disease complexity and insurance status did not appear to predict time to transfer of care. Ongoing analysis will allow for the further development of metrics for defining a successful transfer, identification of risk factors for poor transfer outcomes, and process improvement interventions.

Neonatology general VI
Concurrent session
10:15 AM
Saturday, January 22, 2022

#435 CORRELATION OF MATERNAL AND CORD BLOOD SARS-COV-2 IMMUNOGLOBULIN LEVELS IN COVID-19 DURING PREGNANCY

G Rojas*, M Alam, C Yu. University of California Irvine, Irvine, CA
10.1136/jim-2022-WRMC.431

Purpose of Study COVID-19 caused by the SARS-CoV-2 virus has led to a worldwide pandemic but the impact of SARS-CoV-2 infection during pregnancy and delivery with subsequent neonatal effects still remains unclear. It is known that pregnant women are at higher risk of viral infections given an alteration in immune response, it is therefore likely that many would have had asymptomatic COVID-19 infection at some point during pregnancy. Studies have shown that maternal antibodies can be detected in umbilical cord blood as early as the first trimester and are a reliable source of identification of maternal past and recent infections at the time of delivery. As there is potential of in-utero transmission, it is crucial to determine the significance of a positive serology test at the time of delivery.

Objectives To evaluate the presence SARS-CoV-2 antibodies in cord blood of infants born to mothers with active COVID-19 infection. The goal of this case control study is to provide a better estimate of the incidence of COVID-19 infection in pregnant women admitted to UC Irvine Medical Center, to determine the significance of a positive serology test at the time of delivery, as there is potential of in-utero transmission, it is crucial to determine the significance of a positive serology test at the time of delivery.

Methods Used This is a descriptive cohort with a plan to transfer from pediatric to adult rheumatology within the University of Colorado. Collaborating pediatric and adult rheumatologists created a transfer log to streamline tracking of patients and information sharing. Patients were retrospectively identified for inclusion over a 14 month period. We captured demographics, rheumatologic disease, disease activity, medications, hospital use, insurance status, and compliance with care plans. Data was collected by chart review and managed using REDCap.

Summary of Results Twenty-three patients were identified with plans to transfer to adult rheumatology (table 1). 70% completed a first visit in adult rheumatology with an average of 137 days from the last pediatric rheumatology visit. 30% had their first adult visit within 4 months of their last pediatric visit. In subgroup analysis, patients who did not transfer within 4 months waited a mean of 346 days before the first adult visit. Five patients never established with adult rheumatology and had a longer interval between their last 2 pediatric rheumatology visits compared to those who transferred (313 ± 136 days vs. 170.3 ± 155 days, p=0.07). Disease complexity, number of medications, and disease activity were similar between groups.

Conclusions We examined the outcomes of 23 pediatric patients with a plan to transfer care to an adult rheumatology clinic. Interestingly, disease complexity and insurance status did not appear to predict time to transfer of care. Ongoing analysis will allow for the further development of metrics for defining a successful transfer, identification of risk factors for poor transfer outcomes, and process improvement interventions.

Conclusions SARS-CoV-2 immunoglobulins levels in cord blood correlate with maternal levels and cord blood can be used as a reliable non-risky source to detect maternal and neonatal Covid-19 infection.
## Abstracts

### #436  
**THE IMPACT OF CORONAVIRUS DISEASE 2019 ON PREGNANCY AND INFANT CHARACTERISTICS IN NEW MEXICO**  

*T Emery*, †KN Baillie, †J Maxwell, ‡H Ko.  
†University of New Mexico School of Medicine, Albuquerque, NM; ‡University of New Mexico Health Sciences Center, Albuquerque, NM  
10.1136/jim-2022-WRMC.432

**Purpose of Study** Severe Acute Respiratory Syndrome Coronavirus-1 (SARS CoV-1) and Middle East Respiratory Syndrome (MERS), two illnesses caused by a coronavirus, have been shown to affect maternal and neonatal morbidity and mortality. Coronavirus disease 2019 (COVID-19) has been shown to affect vasculature including placental changes such as microcalcifications and thrombi formation. Significant remodeling of the placenta occurs at the end of the first trimester and into the second trimester. Insults during this crucial period can affect placental size and functionality. These changes may contribute to intrauterine fetal growth restriction (IUGR). We explored the relationship between 1) antenatally-acquired maternal COVID-19 infection and fetal growth and 2) the timing of antenatal COVID-19 infection and neonatal birth weight.

**Methods** Used A retrospective chart review was completed using the University of New Mexico (UNM) Data Warehouse and the electronic medical record system. ICD-10 codes were used to identify those that had a positive pregnancy test and positive COVID screening test between 3/1/2020 and 3/24/2021 at UNM Hospital. Individuals who had a positive pregnancy test but a negative or undocumented COVID screening test without symptoms were included for comparison as a control group. Chi-square analysis was used to compare categorical data.

**Summary of Results** To date, 240 maternal charts have been reviewed with three excluded due to pregnancy termination and 237 neonatal charts have been reviewed. Thirty-four women were identified as COVID-19 positive during pregnancy, 41 women with negative testing, one woman who declined testing, and the remainder with no testing documented and no symptoms reported. Twenty-two women were identified as having COVID-19 infection in the third trimester of pregnancy, with 9% of these infants being small for gestational age. The infants born to women who were negative for COVID-19 were small for gestational age in 12% of the births. Interestingly, the incidence of delivery via cesarean section was 12% in the 41 deliveries to COVID-19 negative women. Of those positive, 27% had cesarean section deliveries, a nearly significant increase (p=0.06).

**Conclusions** The charts reviewed to date did not reveal any significant difference in infant growth parameters at birth in women who tested negative or positive for COVID-19 during pregnancy. However, there was an increased incidence of cesarean section deliveries in women with COVID-19 infection during pregnancy. Most women were not symptomatic, so this does not seem to be related to illness. Additional investigations are required to further delineate if this relationship persists.

### #437  
**DEBRIEFING PERFORMANCE IN A SIMULATION-BASED TRAINING PROGRAM IN NEONATAL RESUSCITATION: SIMULATING SUCCESS**  

†BN Liu*, †1H Lee, †L Bennett, †K Padua.  
†Stanford, Stanford, CA; †California Perinatal Quality Care Collaborative, Stanford, CA  
10.1136/jim-2022-WRMC.433

**Purpose of Study** Neonatal resuscitation is a complex procedure that requires the finesse of a multidisciplinary team. The California Perinatal Quality Care Collaborative (CPQCC) and the Center for Advanced Pediatric Perinatal Education (CAPE) implemented an on-site simulation-based training program called Simulation Success at 15 NICUs over 15-months. Simulation and debriefing present an opportunity for the team to reduce errors and identify latent safety threats. There is a gap in knowledge on real-life debriefing practice for neonatal simulation. This project analyzes the debriefing performance of the NICUs through the course of the program.

**Methods** Used The debrief videos were anonymized and scored according to CAPE Real-Time Debriefing Evaluation tool. The fifteen sites were randomly assigned into four groups (G1, G2, G3, G4). Performance was evaluated through the ratio of trainee responses to the sum of instructor questions and instructor statements (TR:IQ+S). A goal TR:IQ+S was set to be greater than 3:1. Least squares linear regression was applied to the ratios, yielding a line of best fit for each group.

**Summary of Results** 82 total videos were including in the analysis. There were 22, 28, 17, and 15 videos for G1, G2, G3, and G4, respectively. The ratios of G1, G2, and G4 reveal no substantial trends and lie within the same order of magnitude, approaching a slope of zero. However, G3 illustrates a significant rate of change that is one order of magnitude larger than the others. All linear regressions yield a line that lie under the target ratio.

**Conclusions** Variations in debriefer and trainee attendance at the sites may contribute to the lack of trends in G1, G2, and G4. The absence of significant declining debrief performance...
highlight the potential for sustaining debriefing in simulation-based training. G3's trend suggests increased engagement in trainees over time. Lack of ratios near 3:1 hint that this target may be difficult to achieve in practice. Moving forward, evaluation of the simulations in tandem with debriefs may be telling, especially in the context of qualitative briefing performance and correlation to patient outcomes.

**#438 TIME TO POSITIVITY IN BLOOD CULTURES IN A LEVEL IV NICU**

1Y Nishihara, 1C MacBrayne, 2A Priizi, 1I Zenge, 1T Grover, 1S Parker, 2Children's Hospital Colorado, Aurora, CO; 2University of Colorado, Denver, CO

10.1136/jim-2022-WRMC.A34

**Purpose of Study** In the Neonatal Intensive Care Unit (NICU) setting, subtle clinical deterioration of the infant, combined with the lack of specificity in clinical signs to identify true infection often triggers an evaluation for sepsis - where cultures are obtained, and empiric antibiotics are initiated.

Limiting the duration of antibiotic exposure has potential benefits in curtailing antimicrobial resistance and reducing unwanted adverse effects.

We aimed to determine the time to positivity (TTP) of blood cultures in a free-standing level IV NICU over a 6-year period, with the goal to reassess our antimicrobial practice in the NICU.

**Methods Used** Data were extracted from the Children's Hospital Colorado data warehouse for all patients admitted to the NICU, who had a positive blood culture between January 2013 to December 2018. These patient's charts were reviewed for both microbiologic and clinical data. TTP was calculated based on date and time culture was collected, compared to the date and time growth was first reported. Micro-organisms were categorized into absolute pathogens, potential pathogens (e.g., CoNS, other strep), common contaminants, yeast and other less frequently identified organisms.

**Summary of Results** A total of 314 positive blood cultures were identified from 270 individuals. The mean gestational age was 34.59 weeks with an average birthweight of 2356g. Of the 314 positive culture results, 299 (95%) were identified within 24h of blood culture collection. These findings highlight opportunities for antimicrobial stewardship to limit antibiotic exposure in the NICU. The high mortality within 4 weeks of blood culture positivity warrants further study.

**#439 MATERNAL AND PERIPARTUM RISK FACTORS THAT AFFECT C-REACTIVE PROTEIN AND PROCALCITONIN IN NON-INFECTED NEWBORNS DURING THE FIRST 72 HOURS OF LIFE**

A Vachhani*, A Chambless, M Durand, R Ramanathan, R Celayabab. LAC+USC Medical Center, Keck School of Medicine of USC, Los Angeles, CA

10.1136/jim-2022-WRMC.A35

**Purpose of Study** Early detection of infection is difficult as the first signs of infection are nonspecific. Providers often rely on biomarkers to help identify these infants including high sensitivity c-reactive protein (hsCRP) and procalcitonin (PCT). Maternal and peripartum risk factors may influence hsCRP and PCT making levels difficult to interpret. The purpose of this study was to compare the effect of maternal and peripartum risk factors on hsCRP and PCT levels in non-infected infants during the first 72 hours of life.

**Methods Used** Retrospective study of infants admitted to the neonatal intensive care unit from August 2018-February 2021 who were worked up for early onset sepsis. Infants with three paired PCT and hsCRP values and blood, urine or cerebrospinal fluid culture were included. Biomarker levels were drawn at 0, 1, 2 time points at 12-24 hours intervals from birth. Demographics and laboratory results were collected. Infants were classified as non-infected and included in the study if cultures were negative and prolonged antibiotics were not continued for presumed infection. Maternal risk factors included pre-eclampsia and diabetes mellitus (DM). Peripartum risk factors included vaginal delivery, need for positive pressure ventilation (PPV) in the delivery room, presence of meconium-stained amniotic fluid (MSAF) and diagnosis of chorioamnionitis. Data was analyzed with Wilcoxon-rank sum test.

**Summary of Results** A total of 275 infants were classified as non-infected. The median (IQR) gestational age was 34.9 (33.1, 37.6) weeks and median (IQR) birth weight was 2285 (1850, 3120) grams. One hundred one (37%) of infants were born by vaginal delivery. Serial labs were drawn at time points 0, 1, 2 corresponding to median (IQR) hour of life of 1 (1, 2), 13 (12, 20), and 32 (25, 40) hours respectively. Maternal diagnosis of DM did not affect hsCRP or PCT (data not shown). Peripartum risk factors such as MSAF, chorioamnionitis and PPV at birth affected hsCRP, but not PCT. Only maternal diagnosis of pre-eclampsia affected PCT on the second timepoint (table 1).

**Conclusions** Our preliminary findings suggest that maternal and peripartum risk factors affect hsCRP more than PCT, particularly chorioamnionitis and presence of MSAF. These risk factors contributed to rise in hsCRP levels after birth making it difficult to interpret levels in infants screened for infection. Our study showed a natural rise and fall of PCT in non-infected infants, regardless of exposure to risk factors within 72 hours of life. PCT may be a more specific biomarker to screen infants at risk for infection immediately after birth in comparison to hsCRP. However, further studies are necessary to establish time-based reference ranges for PCT to increase clinical utility.
Abstracts

### Abstract #439 Table 1 Risk factors affecting biomarkers

<table>
<thead>
<tr>
<th>Risk Factor</th>
<th>hsCRP (mg/L)</th>
<th>PCT (ng/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Risk Factor Present</td>
<td>Risk Factor Not Present</td>
</tr>
<tr>
<td>Meconium N=22</td>
<td>N=250</td>
<td>N=22</td>
</tr>
<tr>
<td>Stained Amniotic Fluid N=26</td>
<td>N=249</td>
<td>N=26</td>
</tr>
<tr>
<td>Time 0 Lab Draw</td>
<td>0.3</td>
<td>0.2</td>
</tr>
<tr>
<td>(0.2, 0.7)</td>
<td>(0.2, 0.3)</td>
<td>(0.14, 0.15, 0.32)</td>
</tr>
<tr>
<td>Time 1 Lab Draw</td>
<td>5.1</td>
<td>1.0</td>
</tr>
<tr>
<td>(0.7, 0.5, 2.4)</td>
<td>(0.99, 0.84, 7.86)</td>
<td>(10.7)</td>
</tr>
<tr>
<td>Time 2 Lab Draw</td>
<td>4.8</td>
<td>1.2</td>
</tr>
<tr>
<td>(0.9, 0.98)</td>
<td>(0.56, 0.83, 6.55)</td>
<td>(2.96)</td>
</tr>
<tr>
<td>Chorioamnionitis N=26</td>
<td>N=249</td>
<td>N=26</td>
</tr>
<tr>
<td>Time 0 Lab Draw</td>
<td>0.3</td>
<td>0.2</td>
</tr>
<tr>
<td>(0.3, 0.5)</td>
<td>(0.2, 0.3)</td>
<td>(0.15, 0.15, 0.33)</td>
</tr>
<tr>
<td>Time 1 Lab Draw</td>
<td>9.0</td>
<td>1.0</td>
</tr>
<tr>
<td>(0.9, 0.5, 2.1)</td>
<td>(1.67, 0.84, 7.53)</td>
<td>(20.8)</td>
</tr>
<tr>
<td>Time 2 Lab Draw</td>
<td>8.0</td>
<td>1.1</td>
</tr>
<tr>
<td>(1.7, 3.0)</td>
<td>(0.5, 3.9)</td>
<td>(0.56, 0.81, 6.22)</td>
</tr>
<tr>
<td>Pre-Eclampsia* N=71</td>
<td>N=203</td>
<td>N=71</td>
</tr>
<tr>
<td>Time 0 Lab Draw</td>
<td>0.2</td>
<td>0.2</td>
</tr>
<tr>
<td>(0.2, 0.3)</td>
<td>(0.2, 0.3)</td>
<td>(0.16, 0.14, 0.34)</td>
</tr>
<tr>
<td>Time 1 Lab Draw</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>(0.5, 3.7)</td>
<td>(0.5, 2.8)</td>
<td>(0.67, 0.92, 9.6)</td>
</tr>
<tr>
<td>Time 2 Lab Draw</td>
<td>1.9</td>
<td>1.3</td>
</tr>
<tr>
<td>(0.8, 5.8)</td>
<td>(0.5, 4.0)</td>
<td>(0.66, 0.84, 7.14)</td>
</tr>
<tr>
<td>Vaginal Delivery N=101</td>
<td>N=174</td>
<td>N=101</td>
</tr>
<tr>
<td>Time 0 Lab Draw</td>
<td>0.2</td>
<td>0.2</td>
</tr>
<tr>
<td>(0.2, 0.4)</td>
<td>(0.2, 0.3)</td>
<td>(0.15, 0.17)</td>
</tr>
<tr>
<td>Time 1 Lab Draw</td>
<td>1.3</td>
<td>1.0</td>
</tr>
<tr>
<td>(0.7, 0.5, 4)</td>
<td>(0.4, 1.6)</td>
<td>(1.25, 8.18)</td>
</tr>
<tr>
<td>Time 2 Lab Draw</td>
<td>1.8</td>
<td>1.1</td>
</tr>
<tr>
<td>(0.7, 0.5, 9)</td>
<td>(0.5, 3.7)</td>
<td>(0.87, 3.7)</td>
</tr>
<tr>
<td>Use of Positive Pressure at Birth N=162</td>
<td>N=113</td>
<td>N=162</td>
</tr>
<tr>
<td>Time 0 Lab Draw</td>
<td>0.2</td>
<td>0.3</td>
</tr>
<tr>
<td>(0.2, 0.3)</td>
<td>(0.2, 0.4)</td>
<td>(0.16, 0.14, 0.48)</td>
</tr>
<tr>
<td>Time 1 Lab Draw</td>
<td>1.0</td>
<td>1.3</td>
</tr>
<tr>
<td>(0.5, 1.6)</td>
<td>(0.6, 5.8)</td>
<td>(0.84, 1.01, 7.11)</td>
</tr>
<tr>
<td>Time 2 Lab Draw</td>
<td>1.2</td>
<td>1.7</td>
</tr>
<tr>
<td>(0.6, 4.1)</td>
<td>(0.5, 7.5)</td>
<td>(0.98, 0.59, 5.61)</td>
</tr>
</tbody>
</table>

Data are shown as median (IQR) ^ Some missing data

### Abstract #440 EXPERIENCE WITH PARENTAL LEAVE DURING PEDIATRIC FELLOWSHIP: THE FELLOW PERSPECTIVE

N Dyess*, B Weikel, J Barker, T Garrington, TA Parker. University of Colorado Denver School of Medicine, Aurora, CO

10.1136/jim-2022-WRMC.436

Purpose of Study To describe the experience of parental leave during pediatric fellowship training from the fellows’ perspective.

Methods Used A national web-survey was sent to fellows at ACGME-accredited pediatric fellowship programs, either directly or through their program.

Summary of Results Of the 1003 fellows who opened the survey, 726 (72%) provided information on their experience with parental leave during fellowship training. 43% reported having children, 26% planned on having another child during fellowship, and 16% are unsure. Of those with children, 59% have 1 child and 27% have 2 children. Approximately half of those with children reported having at least one child during fellowship. Of the 56% who were not planning on having a child during fellowship, only 14% stated their program’s parental leave policy (PLP) was a deterrent.

Of respondents who do not have children, 56% experienced colleagues who had or adopted a child during fellowship. 24% reported feeling unduly burdened by their colleague’s decision to have a child during fellowship and 13% felt their colleagues should have delayed childbearing.

Of these, 47% felt that an alternative PLP may have alleviated these feelings.

Of respondents who had children during fellowship, daycare and a fellow’s partner were the primary means of childcare while the fellow is at work. 81% of fellows breastfed their children during fellowship for an average of 9 months. 82% of respondents who had a child during fellowship stated they did not experience significant financial difficulties. 56% reported receiving adequate PLP information prior to their leave and 36% reported they did not. 15% of respondents reported feeling discrimination at work while pregnant. 33% of respondents who had children during fellowship are having to extend their fellowship end date due to their parental leaves, and 34% took unpaid weeks to construct their parental leaves (average of 4 weeks). Regardless, 87% of respondents do not wish they would have delayed childbearing until after fellowship.

Conclusions Many pediatric fellows have children or are planning on having children during fellowship. Areas where a trainee’s experience with parental leave during pediatric fellowship could improve include increased clarity and transparency of PLP to better guide a fellow’s family planning, allowance of parental leave without having to extend training, increased paid leave to minimize unpaid leave utilization, increased advocacy on the parent fellow’s behalf to eradicate discrimination, and increased protection of co-fellows not planning on having children in fellowship.
Purpose of Study AAP, ACOG, and NRP have recommended delayed cord clamping (DCC) for term and preterm deliveries for at least 30–60 seconds after birth. It has been established that DCC provides short term and long-term benefits. In preterm infants, DCC is associated with improved transitional circulation, higher hemoglobin, decreased need for blood transfusion, and lower incidence of necrotizing enterocolitis. The purpose of this study is to compare short-term and long-term outcomes in late preterm infants (LPT) exposed to delayed cord clamping vs. immediate cord clamping (ICC).

Methods Used Retrospective study of preterm infants born at 34.0/7–36.6/7 weeks of gestation and admitted to the neonatal intensive care unit (NICU) between 2018–2020. Demographics, cord clamping information, laboratory values, clinical and neurodevelopmental data were extracted from electronic medical records and the neonatal database. Age and Stages Questionnaire was used to evaluate neurodevelopment at 18 months chronologic age (CA). Data was analyzed with Chi-Square or Fisher Exact Test and Wilcoxon Rank Sum test where appropriate.

Summary of Results There were 156 infants admitted to the NICU during the study period. Preliminary analysis included 50 infants exposed to ICC and 50 infants exposed to DCC. There were 19/100 (19%) infants followed up for neurodevelopment at 18 months. Infants exposed to DCC were significantly smaller, immature and with lower APGAR scores at 1 and 5 minutes compared to infants in the ICC group. Infants exposed to DCC had significantly lower median temperature on admission to the NICU and a higher rate of hypothermia defined as temperature less than 36.5 °C. One infant in the DCC group had a temperature less than 36°C. Median hemoglobin levels at 18–36 hours of life and rate of phototherapy were not significantly different between groups. No infant exposed to DCC compared to 3 (6%) infants in ICC received packed red blood cell transfusion within the first 24 hours of life. Median hemoglobin values at 12 months CA and median ASQ scores at 18 months of CA were similar in both groups. (Table 1).

Conclusions Our preliminary data showed that DCC compared to ICC in LPT infants did not result in a higher hemoglobin at birth and at 1 year of age, and neurodevelopmental outcomes at 18 months CA was similar. DCC in immature and smaller infants can be a risk factor for hypothermia, therefore, it is important to follow recommendations by NRP to prevent this adverse effect. Data collection is continuing to increase sample size and verify these findings.

## Neonatology pulmonary III

Concurrent session

10:15 AM
Saturday, January 22, 2022

### #442 HIGHER WEIGHT FOR LENGTH IS ASSOCIATED WITH INCREASED SEVERITY IN BRONCHOPULMONARY DYSPLASIA IN PRETERM INFANTS

R. Kalkkot, K. Hanvey, K.C. Hart, P. Alu*. The University of Mississippi Medical Center, Jackson, MS

Purpose of Study Bronchopulmonary dysplasia (BPD) causes significant morbidity and mortality in infants. Obesity negatively impacts lung function in children and adolescents. There are no studies correlating weight for length (Wt/L) or the body mass index (BMI) with BPD outcomes in preterm infants with higher Respiratory Severity Scores (RSS) in BPD. Wt/L is accurate in predicting fat-free mass and fat mass in premature infants. Our aim was to explore with a prospective observational study if higher BMI correlates with higher RSS in preterm infants with BPD.

Methods Used Supported by NIH (Award # 1U54GM115428). Preterm infants born @<30 weeks requiring respiratory support @30–33 weeks for >2 days were enrolled. Weight, length, and head circumference were measured weekly. Data on Wt/L z-scores & BMI were collected. RSS was calculated as per the STOP-ROP trial. Data were analyzed using descriptive and inferential statistics. Linear regression analysis was used to assess the association between BMI and RSS. The results of this study will provide insights into the impact of obesity on the severity of BPD.

**Abstract #442 Table 1** Demographics, short- and long-term outcomes between the two groups

<table>
<thead>
<tr>
<th>DEMOGRAPHICS</th>
<th>Immediate Cord Clamping n=50</th>
<th>Immediate Cord Clamping n=50</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth Weight (g) *</td>
<td>2560 (2360, 2890)</td>
<td>2230 (2068, 2525)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Gestational Age (wks) *</td>
<td>35.8 (35.1, 36.3)</td>
<td>34.7 (34.1, 35.1)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Male gender, n (%)</td>
<td>29 (58)</td>
<td>25 (50)</td>
<td>0.42</td>
</tr>
<tr>
<td>Hispanic Race, n (%)</td>
<td>29 (55)</td>
<td>24 (48)</td>
<td>0.09</td>
</tr>
<tr>
<td>Cesarean Section, n (%)</td>
<td>38 (76)</td>
<td>33 (66)</td>
<td>0.27</td>
</tr>
<tr>
<td>Maternal Preeclampsia, n (%)</td>
<td>7 (14)</td>
<td>11 (22)</td>
<td>0.30</td>
</tr>
<tr>
<td>Maternal Gestational Diabetes, n (%)</td>
<td>11 (22)</td>
<td>13 (26)</td>
<td>0.64</td>
</tr>
<tr>
<td>APGAR score 1 min *</td>
<td>7 (3, 8)</td>
<td>8 (7, 8)</td>
<td>0.03</td>
</tr>
<tr>
<td>APGAR score 5 min *</td>
<td>8 (6, 9)</td>
<td>9 (8, 9)</td>
<td>0.03</td>
</tr>
</tbody>
</table>

**Summary of Results** There were 156 infants admitted to the NICU during the study period. Preliminary analysis included 50 infants exposed to ICC and 50 infants exposed to DCC. There were 19/100 (19%) infants followed up for neurodevelopment at 18 months. Infants exposed to DCC were significantly smaller, immature and with lower APGAR scores at 1 and 5 minutes compared to infants in the ICC group. Infants exposed to DCC had significantly lower median temperature on admission to the NICU and a higher rate of hypothermia defined as temperature less than 36.5 °C. One infant in the DCC group had a temperature less than 36°C. Median hemoglobin levels at 18–36 hours of life and rate of phototherapy were not significantly different between groups. No infant exposed to DCC compared to 3 (6%) infants in ICC received packed red blood cell transfusion within the first 24 hours of life. Median hemoglobin values at 12 months CA and median ASQ scores at 18 months of CA were similar in both groups. (Table 1).

**Conclusions** Our preliminary data showed that DCC compared to ICC in LPT infants did not result in a higher hemoglobin at birth and at 1 year of age, and neurodevelopmental outcomes at 18 months CA was similar. DCC in immature and smaller infants can be a risk factor for hypothermia, therefore, it is important to follow recommendations by NRP to prevent this adverse effect. Data collection is continuing to increase sample size and verify these findings.
mixed model regression was used to study the relationship between the continuous variables with potential covariates.

Summary of Results We enrolled 81 preterm infants. Maternal and neonatal demographics are in the figure. RSS was significantly higher in grade 3 BPD- 1.0 vs 0.23 in grade 1 BPD (p <0.0001) (figure 1). For every unit increase in Wt/L Z score, RSS increased by 0.069 (p <0.0001). RSS significantly correlated with Wt/L Z scores, percentiles, and BMI (P ≤ 0.0001) even after including postnatal steroids, gestational age, and sex in the regression model. The results remained significant after excluding infants with postnatal steroids. The mean RSS @ 40 weeks in Wt/L ≤ 50 was 0.09 ± 0.27 compared to 0.47 ± 0.37 in >50 percentile category (p=0.01).

Conclusions Our study is the first to show that higher BMI and W/L may adversely affect respiratory severity in BPD infants. Hence, an optimal W/L ratio should be maintained to avoid additional burdens in infants with BPD.

#443 NOVEL APPLICATION OF NEONATAL POINT-OF-CARE ULTRASOUND: ULTRASOUND-GUIDED LESS INVASIVE SURFACANT ADMINISTRATION

1J Velasquez*, 2R Ramanathan, 3A Yeh. 1LAC+USC Medical Center, LA, CA; 2Keck School of Medicine of USC, LA, CA

10.1136/jim-2022-WRMC.439

Purpose of Study Point-of-care ultrasound (POCUS) has become a very useful tool in guiding bedside clinical management. This study aims to describe a novel and innovative application of neonatal POCUS: Ultrasound-guided Less Invasive Surfactant Administration (LISA).

Methods Used This is a single-center, prospective observational study conducted in a level III NICU. In our unit, LISA is performed routinely in newborns on noninvasive respiratory support needing surfactant therapy. A 16 GA x 5.25" BD Angiocath is adapted to deliver the surfactant. The catheter insertion was performed as per standard unit protocol. As part of the study protocol, POCUS was used to identify and confirm catheter position inside the trachea, along with video laryngoscopy in all cases to ensure the catheter passing through the vocal cords.

POCUS was performed with the linear transducer (GE LOGIQ e) positioned transversely over the suprasternal notch. The trachea can be identified in the midline position as a hyperechoic air-mucosa interface with reverberation artifacts. A palpable anatomic landmark allows for a simpler, more feasible, and reproducible technique while ensuring, based on anatomy, the visualization of the catheter in the correct position.
IS EARLY LIFE ACETAMINOPHEN EXPOSURE RELATED TO AN INCREASED RISK OF BPD?

1 S McKenna*, 2 CJ Wright, 1 L Sherlock, 1 D Riebel, 1 E Jensen. 1University of Colorado – Anschutz Medical Campus, Aurora, CO; 2The Children’s Hospital of Philadelphia, Philadelphia, PA

10.1136/jim-2022-WRMC.440

Purpose of Study Acetaminophen(APAP) is commonly administered to preterm infants and is increasingly used to treat the patent duc tus arteriosus(PDA). Preclinical models demonstrate that the developing lung is susceptible to APAP-induced injury. Whether early life APAP exposure contributes to lung injury in preterm neonates is unknown.

Methods Used We performed a systematic literature search on PUBMED to identify randomized controlled trials evaluating APAP for prevention or treatment of a PDA. Relevant trial and clinical data including treatment group rates of BPD were abstracted. Random effects meta-analysis and meta-regression according to the mean gestational age(GA) of trial participants were performed.

Summary of Results Of 196 identified manuscripts, 12 trials with 1001 subjects were included in the final analyses (figure 1). Mean GA ranged from 25.3 to 33.6. The outcome of BPD, although variably defined, was reported for 930 subjects. Averaged across all trials, APAP exposure did not significantly increase the risk of BPD [figure 2A; RR 1.07 (0.83, 1.37)]. However, meta-regression demonstrated a possible inverse relationship between GA and BPD risk. For each week decrease in mean study GA, the risk difference for developing BPD with APAP exposure increased by 1.2% (range 3% to -0.5%; p=0.17; figure 2B). When limited to the 8 studies that compared oral APAP to oral ibuprofen, this weekly risk increased to 1.7%(range 4.0% to -0.6; p=.14; figure 2C). These results bordered on statistical significance, but the power to detect true differences at lower gestational age was limited by the low number of trial participants born <28 wks. Among all 12 reviewed trials, only 127 subjects born <28 wks were randomized to APAP. The 8 trials comparing oral APAP to ibuprofen randomized only 68 subjects <28 wks to APAP.

Conclusions Data from published RCTs show no significant effect of APAP on BPD risk in preterm infants. However, a possible and concerning inverse association between the risk of BPD with APAP exposure by GA suggests possible adverse effects among the least mature infants. More data are needed to determine whether early life APAP exposure in extremely premature infants is safe.
PREDICTORS AND OUTCOMES OF LATE-PRETERM NEONATES NEEDING RESPIRATORY SUPPORT IN THE DELIVERY ROOM

A Fikinos*, FB Wertheimer, R Ramanathan, M Biriwale. Los Angeles County University of Southern California Medical Center, Los Angeles, CA

Purpose of Study The etiology and management of respiratory distress in late preterm neonates is not as well described as in the very preterm population. This study identified factors associated with delivery room (DR) positive pressure ventilation (PPV) support in the late preterm population.

Methods Used Preterm neonates with a gestational age of 34–36 weeks who were admitted to LAC+USC from 2017–2020 were included in this retrospective review. DR ventilatory support, as well as maternal history, initial lab results, and comorbidity diagnoses were abstracted. Study variables were analyzed with Chi-squared and t-tests.

Summary of Results Approximately 37% [167/460] of the neonates required PPV in the DR and 92% (n=153/167) were placed on nasal cannula. Nasal continuous positive airway pressure (NCPAP) mode was most utilized (n=132), followed by high flow nasal cannula (HFNC) (n=89), nasal intermittent positive pressure ventilation (NIPPV) (n=105), and intubation (n=18). Face masks were only utilized in preparation for intubation.

Maternal factors associated with the requirement of PPV in the DR included pre-eclampsia (16.4% [45/230] vs 24.5% [39/159], p=0.044), placenta previa (2.2% [6/274] vs 9.3% [15/161], p=0.002), and placental abruption (1.1% [3/274] vs 4.9% [8/162], p=0.023). In our study, chorioamnionitis, prolonged rupture of membranes > 18 hours, substance abuse, and meconium aspiration were not significantly associated with the need for PPV. Neonatal factors are listed in table 1.

Air leaks on initial X-ray (0% [0/279] vs 2.4% [4/164], p=0.018, respiratory distress syndrome (RDS) (0.4% [1/279] vs 14.6% [24/164], p = 0.001), and sepsis (1.8% [5/278] vs 5.3% [9/164], p=0.047) were significant comorbidities associated with the use of DR PPV support in late preterm neonates.

Conclusions Late preterm neonates who required PPV support in the DR tended to be younger, with lower APGAR scores, and higher PCO₂. These infants were more likely to need invasive ventilatory support at 24 hours. Maternal conditions compromising fetal perfusion were associated with respiratory support requirements while maternal infections were not. Neonates who received PPV support in the DR were more likely to have air leaks on initial X-ray, RDS and sepsis.

SURVEY OF CONTINUOUS POSITIVE AIRWAY PRESSURE (CPAP) PRACTICES AMONG NEONATAL PROVIDERS IN THE UNITED STATES

R Mamidi*, K MacDonald, M Go, C McEvoy, Oregon Health and Science University, Portland, OR

Purpose of Study To evaluate practice variation in the initiation, management, and discontinuation of neonatal nasal CPAP (nCPAP) among the Neonatal-Perinatal section of the American Academy of Pediatrics (AAP). We hypothesized large variations in clinical practice particularly with regards to discontinuation of CPAP and use of chinstraps.

Methods Used A web-based IRB approved survey consisting of 25 questions was administered to the AAP Neonatal-Perinatal section. It inquired about the NICU size, level, and location, and evaluated approaches for initiation and management of CPAP, and methods and post-menstrual ages used to wean infants born at <32 weeks gestation off CPAP. It also assessed the use of chinstraps and rates of nipple feeding while on CPAP. Data were analyzed using descriptive statistics.

Summary of Results Overall, 857/3700 (23%) of providers (718 neonatologists and 98 neonatal fellows) consented to the survey. The majority (83%) worked in a NICU with >20 beds and 94% in a level 3 or higher. 822/830 (99%) used CPAP with 300 (36%) primarily using bubble CPAP in infants <32 weeks gestation; 229 (27%) using ventilator driven CPAP and 262 (31%) using a combination of these. In terms of the primary interface applied: 297/781 (38%) used short binasal prongs; 61 (8%) used long binasal prongs; 106 (14%) used a nasal mask; 295 (38%) used a combination of these interfaces. There was wide variation in: 1) the maximum CPAP setting applied to an infant <32 weeks gestation in the first 72 hours of life with 46/795 (6%) using 5 cmH₂O; 238 (32%) using 6 cmH₂O; 244 (31%) using 7 cm H₂O; 216 (27%) using 8 cm H₂O; 31 (4%) using ≥ 8 cm H₂O; 2) the typical minimum CPAP setting prior to stopping or weaning CPAP, responses were: 224/794 (28%) decreased to 4 cmH₂O; 493 (62%) to 0.4 cmH₂O; and 53 (7%) decreased to 0 cmH₂O; and 3) the use of chinstraps: 151 (19%) use a chinstrap at all times with 190 (24%) use it only during feeding.

Abstract #445 Table 1 Neonatal factors associated with the requirement of PPV in the DR

<table>
<thead>
<tr>
<th>Gestational age (weeks)</th>
<th>Birth weight (g)</th>
<th>5 min APGAR</th>
<th>First blood pH</th>
<th>First blood gas CO₂</th>
<th>Invasive ventilation at 24 hours (%)</th>
<th>Need for surfactant (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>No respiratory support in DR</td>
<td>35.5</td>
<td>2585</td>
<td>8.9</td>
<td>7.16</td>
<td>44.8</td>
<td>0</td>
</tr>
<tr>
<td>Respiratory support in DR</td>
<td>35.1</td>
<td>2716</td>
<td>7.6</td>
<td>7.27</td>
<td>52.4</td>
<td>6.7</td>
</tr>
<tr>
<td>P-value</td>
<td>&lt;0.001</td>
<td>0.020</td>
<td>&lt;0.001</td>
<td>0.464</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Abstract #446 Table 1 Percent of responses for preferred method of weaning off nCPAP

<table>
<thead>
<tr>
<th>Preferred Method to Wean off nCPAP</th>
<th>% of Responses Favoring this Method</th>
</tr>
</thead>
<tbody>
<tr>
<td>Directly to room air</td>
<td>37%</td>
</tr>
<tr>
<td>Wean to high or low flow nasal cannula and then room air</td>
<td>59%</td>
</tr>
<tr>
<td>Increased time per day off CPAP until off</td>
<td>5%</td>
</tr>
</tbody>
</table>

Abstract #446 Table 1 Percent of responses for preferred method of weaning off nCPAP

5 cmH2O and 44 (6%) to 6 cmH2O; 3) the methods to wean off CPAP (see table 1). When asked when CPAP is discontinued or weaned, 69% responded when specific stability criteria were achieved; 22% at a specific PMA (5% at >30 weeks; 85% at >32 weeks; 11% at >34 weeks), and 8% answered other. Greater than 65% of responses endorsed that chinstraps improve CPAP efficacy; however, less than 11% routinely applied a chinstrap. When asked regarding barriers to their use, 27% responded there were no barriers; 20% responded no sign of leak at the mouth; 15% reported CPAP bubbling is adequate; 10% did not have the equipment. 78% of responses (614/879) never allowed infants to nipple on CPAP. 54% of providers worked in a NICU without guidelines for CPAP initiation and 64% without guidelines for CPAP discontinuation. 

Conclusions There are significant practice variations across neonatology caregivers with regards to CPAP initiation, management, and discontinuation. Further evidence-based research is needed in relation to respiratory outcomes to optimize CPAP strategies.

Neuroscience III
Concurrent session
10:15 AM
Saturday, January 22, 2022

#447 INTRADISCAL CONCENTRATED BONE MARROW ASPIRATE INJECTIONS REDUCE PAIN IN PATIENTS WITH LUMBAR DEGENERATIVE DISC DISEASE

1 NM Davies*, 1 GB Forrmy, 2 J Hawley, 3 J Galloway, 1, 3 J McGirley, 1 University of Washington School of Medicine, Seattle, WA; 2 University of Idaho, Moscow, ID; 3 The McGirley Clinic, Casper, WY

Purpose of Study Low back pain, often caused by degenerative disc disease (DDD), is one of the most common reasons for primary care and orthopedic office visits. It’s also among the costliest medical issues affecting our healthcare system. Concentrated bone marrow aspirate (BMAC) injections, along with conservative management, represent a cost-effective and minimally invasive treatment option. Our hypothesis is that BMAC injections, along with conservative care, can provide short and long-term pain reduction in patients with DDD.

Methods Used A retrospective chart review was conducted for 17 patients with DDD who received a BMAC injection along with conservative care in our clinic over the past 6 years. Each patient underwent a procedure where 60 ml of bone marrow was aspirated from the posterior iliac crest, concentrated to 10 ml, and then injected into the degenerated disc. Patients were instructed to limit activity to non-weight bearing for 3 weeks post-procedure and partial weight-bearing with a 5 lb. lifting limit for an additional 3 weeks. All NSAIDs were held 10 days prior to and 3 months following the procedure. A 0–10 patient self-reported pain scale was used as the primary outcome. Secondary outcomes included adverse events and additional treatments. Pain scores were collected prior to the initial treatment and post-treatment at 3 weeks, 6 weeks, 3 months, 6 months, 12 months, 18 months, 24 months, and 36 months. Comparisons were analyzed using a paired-samples t-test.

Summary of Results 17 patients (11 male, 6 female) with ages ranging from 27-80 years old (49.7 ±16.4 years) were included in the study with an average follow-up time of 18.2 ± 16.3 months. Reported pain levels were significantly reduced at 3 months post-injection compared with baseline (13 patients; Δ-3.3 points; p = < 0.001). Additionally, pain continued to decrease up to 3 years post-injection (7 patients; Δ-4.5 points from baseline; p = 0.016). No adverse events were reported. No patients elected for surgery following treatment or received repeat BMAC injections during the 36-month follow-up period.

Conclusions Patients with DDD who were treated with BMAC, along with conservative care, experienced a significant improvement in long-term pain levels without major adverse events related to the treatment. This represents a viable minimally invasive treatment option in patients with painful DDD.
difference. Of the 26 cases, three cases fell outside of normal physiologic MCT interocular differences with values as follows: case 3247 (58.64µm, OS > OD), case 5721 (38.55µm, OS > OD) and case 7051 (74.46, OD > OS). The remaining 23 cases had MCT interocular differences within normal physiologic bounds. Each case was then matched to the patient diagnoses for further evaluation.

Conclusions The 3 cases with abnormal interocular MCT values presented with the following diagnoses: wet AMD (OS), subclinical MNV (OD), dry AMD with GA (OS) and dry AMD with drusen (OD), and a right sided orbital AV fistula. Given the few cases which displayed abnormal MCT values, any significant correlation between choroid thickness as a biomarker for AMD or other ophthalmic conditions would seem premature. As the sample population remains small, increased recruitment of participants will be pursued for further study.

#449 A CLOUD-BASED PIPELINE TO PROCESS VERY PRETERM INFANT DIFFUSION MRI AND AUTOMATED FIBER QUANTIFICATION TRACTOGRAPHY

1LE Rickerich*, 1D Sproul, 1L Bruckert, 1G Lema-Usabiaga, 2SE Dubner, 2KE Travis, 1Stanford University, Stanford, CA; 2Stanford University School of Medicine, Stanford, CA

10.1136/jim-2022-WRMC.445

Purpose of Study Preterm infants are at high risk for white matter (WM) injury detectable using advanced diffusion MRI (dMRI) methods. WM microstructural metrics derived from tractography of dMRI relate to neonatal illness and clinical outcomes. Preparing and processing dMRI data, identifying WM tracts, and extracting microstructural metrics is complex and data-intensive. Our goal was to implement a fully reproducible, cloud-based pipeline to process dMRI, perform tractography, and extract microstructural metrics in a database of infants born at <32 weeks and imaged at near-term gestational age.

Methods Used The processing pipeline utilizes Flywheel, a cloud-based data processing platform. Analyzes are implemented as Dockerized containers or gears. Aligned T1-weighted (T1w) anatomical images are segmented using Infant Freesurfer. Infant Freesurfer output and a template identifying anatomical regions of interest (ROIs) are fed together into a processing gear to create ROIs in each infant subject space. dMRI data is preprocessed and the anatomical T1w data, brain mask (distinguishing brain from non-brain), output markers, and dMRI data are used to generate individual tract profiles and extract WM metrics.

Summary of Results To test our pipeline, we pre-selected a cohort of infants born <28 weeks gestation (N = 55; mean gestational age = 26.5 (1.2) weeks; mean postmenstrual age at scan = 37.1 (1.8) weeks) with both T1w and dMRI scans, intended for analysis in a future study. We segmented and extracted fractional anisotropy (FA) and mean diffusivity (MD) values for 6six corpus callosum segments. T-tests assessed consistency of different approaches to segmenting tracts (whole-brain tractography, WBT, versus region of interest-to-region of interest, ROI-to-ROI).

31 subjects (56.4%) successfully ran through the pipeline. The remaining 24 subjects were likely unsuccessful due to computational resource allocation issues. Within each corpus callosum segment, values were generated for over 90% of subjects. Mean FA and MD are shown in the table 1. Results suggest variability for some tracts. Mean FA differed between WBT and ROI in the posterior CC. Our results were consistent with published studies with regard to both WBT and ROI-to-ROI tractography.

Conclusions We successfully implemented a reproducible pipeline to analyze preterm infant dMRI data. The next steps are to verify the anatomical accuracy of our ROI approaches, resolve resource allocation issues and process remaining subjects, and expand the pipeline to include other major cerebral and cerebellar tracts. Successful implementation of the pipeline will enable reproducible analyses to address questions relevant to clinicians and researchers surrounding neonatal brain development in the context of preterm birth.

#450 ASSESSING CONCUSSION EDUCATION IN ALLOPATHIC MEDICAL AND PHYSICAL THERAPY PROGRAMS

11E A Tangog*, 11W Cheung, 1L McKay, 1Alisky, 1Baron, 1Western University of Health Sciences, Pomona, CA; 1Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA; 1The University of Arizona College of Medicine Tucson, Tucson, AZ

Purpose of Study Our previous study assessed the state of concussion education and students’ interest in the subject across osteopathic medical schools in the country. The results from the first study suggested that osteopathic medical students expressed a desire to obtain a more formal concussion education in their respective programs. This study evaluates the same parameters regarding concussion symptoms, diagnosis and management, while also expanding the study population to include students from allopathic medical and physical therapy programs across the country.

Abstract #449 Table 1

<table>
<thead>
<tr>
<th>Tract</th>
<th>n</th>
<th>WBT: FA mean (SD)</th>
<th>ROI: FA mean (SD)</th>
<th>WBT: MD mean (SD)</th>
<th>ROI: MD mean (SD)</th>
<th>FA t-test statistic</th>
<th>FA t-test p-value</th>
<th>MD t-test statistic</th>
<th>MD t-test p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>CC-Oc.</td>
<td>31</td>
<td>0.13 (0.06)</td>
<td>0.17 (0.07)</td>
<td>1.46 (0.21)</td>
<td>1.52 (0.19)</td>
<td>-2.27</td>
<td>0.03</td>
<td>-1.19</td>
<td>0.23</td>
</tr>
<tr>
<td>CC-Post.</td>
<td>31</td>
<td>0.11 (0.04)</td>
<td>0.14 (0.05)</td>
<td>1.48 (0.17)</td>
<td>1.57 (0.25)</td>
<td>-2.12</td>
<td>0.04</td>
<td>-1.69</td>
<td>0.1</td>
</tr>
<tr>
<td>Parietal</td>
<td>28</td>
<td>0.12 (0.05)</td>
<td>0.15 (0.05)</td>
<td>1.56 (0.42)</td>
<td>1.56 (0.16)</td>
<td>-2.55</td>
<td>0.01</td>
<td>-0.88</td>
<td>0.38</td>
</tr>
<tr>
<td>CC-Sup. Parietal</td>
<td>28</td>
<td>0.14 (0.04)</td>
<td>0.15 (0.03)</td>
<td>1.51 (0.26)</td>
<td>1.56 (0.16)</td>
<td>-1.95</td>
<td>0.06</td>
<td>-0.86</td>
<td>0.39</td>
</tr>
<tr>
<td>CC-Motor</td>
<td>28</td>
<td>0.16 (0.03)</td>
<td>0.17 (0.04)</td>
<td>1.57 (0.28)</td>
<td>1.85 (0.22)</td>
<td>-1.32</td>
<td>0.19</td>
<td>-1.11</td>
<td>0.27</td>
</tr>
<tr>
<td>CC-Ant. Frontal</td>
<td>30</td>
<td>0.15 (0.04)</td>
<td>0.17 (0.04)</td>
<td>1.55 (0.24)</td>
<td>1.62 (0.20)</td>
<td>-1.63</td>
<td>0.11</td>
<td>-1.23</td>
<td>0.22</td>
</tr>
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</table>
Methods Used An electronic survey was distributed to current allopathic medical and physical therapy students from all years of training in the United States. The same survey from the first study was used with an addition of two questions, making the total 16 questions. Four questions corresponded to participant demographics, six questions evaluated knowledge in concussion diagnosis and treatment, and four questions gauged any prior experience with concussion education. The two new questions allowed participants to select their gender and differentiate which program they were in. Data were collected in aggregate.

Summary of Results Based on the preliminary data collection of 338 responses, about 50% of both allopathic medical and physical therapy students correctly identified the definition of a concussion. Furthermore, more than half of responders identified that they have not received any formal training or education on concussions and request to see official education on concussion pathophysiology, symptoms, and treatment integrated into their program’s curriculum.

Conclusions While data collection is still ongoing, the preliminary results of our study suggest that some allopathic medical and physical therapy programs lack the inclusion of this topic in their curriculums. Furthermore, participants identify that receiving concussion education in their preclinical years is essential, valuable, and applicable to any setting of medical care. Overall, our data support a want and need of a comprehensive concussion education in both programs across the country.

#451 ACUTE MONOCULAR VISUAL LOSS AFTER SINUS SURGERY

P Parkh*, OM Dumitracu. Mayo Clinic Arizona, Scottsdale, AZ

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Postoperative monocular vision loss is caused by ischemic optic neuropathy (ION) or central retinal artery occlusion (CRAO). While brain MR-DWI (diffusion-weighted imaging) is not routinely used for diagnosis, recent studies have shown restricted diffusion of the optic nerve (ON) or retina may indicate acute CRAO. Here, we present a case of acute CRAO to highlight this neuroimaging finding and further therapy options.

Case-Report A 71-year-old male without vascular risk factors underwent maxillary bilateral antrostomy and septoplasty for chronic sinusitis. 20–30 minutes upon awakening, he reported acute painless left eye vision loss. Visual acuity (VA) was bare light perception (left) and 20/25 (right). Facial CT was negative for pathology. Fundoscopy showed retinal whitening, cherry-red spot, narrowed arterioles with scattered segmentation, and no hemorrhage. Brain MRI showed left ON diffusion restriction and increased T2 signal (figure 1), concerning for posterior ION; however, proximal CRAO was more likely given the clinical exam. Patient received oral aspirin, intravenous verapamil and alteplase, IV acetazolamide and dexamethasone, and topical Alphagan. At 6-month follow-up, left eye VA was no light perception. Despite extensive workup, no other etiology was found.

Discussion We present a case of proximal CRAO post-sinus surgery in which brain MR-DWI helped with lesion localization. The outcome was dismal despite therapy. Both ION and CRAO can cause postoperative vision loss. PION has normal ophthalmoscopic findings, and ON DWI due to cytotoxic edema has 81–100% sensitivity and 86–100% specificity. CRAO has retinal edema, segmentation of arterioles, and a cherry-red spot. ON DWI sensitivity and specificity are 55%/70–100%, per a recent study. In our case, the ON diffusion restriction was caused by CRAO in its narrowest part, where it enters the ON. A similar reported case of post-sinus surgery acute CRAO underwent therapy with hyperbaric oxygen (HBO) with visual improvement. American Academy of Ophthalmology recommends emergent stroke workup and antiplatelet; there is no level I data to support HBO or other treatments for CRAO. In conclusion, we highlight the combination of retinal ischemic features on fundoscopy and proximal ON diffusion restriction on MR-DWI to be diagnostic of proximal CRAO. Future studies should evaluate the accuracy and utility of MR-DWI and HBO in acute CRAO.

Published only – not presented

#452 CLINICAL VIABILITY OF BIOMARKERS FOR TOXIC STRESS: AN INVESTIGATION

1,2 Pasumarti*. 1Nova Southeastern University, Fort Lauderdale, FL; 2Office of the California Surgeon General, Sacramento, CA

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Purpose of Study The toxic stress response is a biological process that may be caused by the prolonged activation of the stress response due to exposure to cumulative adversity. This process can disrupt biological systems through neuroendocrine, immunologic, and epigenetic mechanisms. The toxic stress
response has been correlated with acute and chronic illnesses, including some of the leading causes of death such as heart disease, suicide, and cancer. Early clinical identification of the toxic stress response and associated health conditions is imperative to limit potentially lifelong negative health effects. The use of clinical biomarkers stands to be a promising method of early identification of the toxic stress response. However, research around clinical biomarkers is still in its nascency and there has been little to assess the clinical viability of these biomarkers. The objective of this study is to determine the potential clinical viability of biomarkers for the toxic stress response.

Methods Used A literature review was conducted to aggregate biomarkers for the toxic stress response. Biomarkers were then scored based on invasiveness (non-invasive and invasive) and definitiveness in diagnosing the toxic stress response (not definitive, semi-definitive, and mostly definitive). Biomarkers with a low invasive score and a high definitive score were considered to have high clinical viability.

Summary of Results Biomarkers considered primary mediators of the toxic stress response, such as cortisol, epinephrine, and norepinephrine, were considered among the most clinically viable biomarkers with low invasive scores and fairly definite scores. Some promising biomarkers with currently low clinical viability include telomere shortening and infant eye-tracking.

Conclusions This research could indicate which biomarkers might be suitable for the development of clinical guidance and which might need further research before more widespread acceptance. Areas for future research include identifying novel biomarkers for the toxic stress response and expanding the body of literature and the strength of the evidence surrounding these biomarkers.

## #453 FULMINANT HEPATIC FAILURE SECONDARY TO WILSONS DISEASE

1VK Narang*, 1M Truong, 1K Grewal, 1L Mosavi, 1J Kalha, 1S Mishra, 2K Radicic, 3UCLA-Kern Medical, Bakersfield, CA; 2Ross University School of Medicine, Miramar, FL

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Case Report Wilson Disease (WD) is a rare genetic disorder of impaired copper excretion causing accumulation of copper in vital organs including the liver, brain, and cornea. Diagnosis usually occurs between the ages of 5 and 35 years. Patients typically present with cirrhosis, neurological symptoms, and Kayser-Fleischer rings. Rarely, WD can present as fulminant hepatic failure necessitating liver transplant. Here, we describe a 36-year-old male with alcohol abuse with acute liver failure initially deemed secondary to alcohol. However, biochemical findings were highly suggestive of WD.

Methods Retrospective single case review after IRB approval

Case Report A 36-year-old male with a history of alcohol abuse presented to the hospital with bilateral lower extremity edema. He had no known prior liver disease and had stopped alcohol 2 months ago. Initial workup showed acute liver failure with elevated liver function tests (LFTs), and imaging findings of decompensated cirrhosis. His mentation declined, and he developed tremors and hallucinations. He was treated with diuretics, thiamine, and Ativan for alcohol withdrawal. Despite one week of treatment, his LFTs remained elevated, and his mentation declined. Further workup revealed ceruloplasmin of 13 mg/dL. Initial 24-hour urinary copper excretion was unobtainable due to his mentation, but after penicillamine challenge it was 111 mcg/24 hr. He was noted to have possible Kayser-Fleischer rings but slit-lamp exam could not be performed due to accessibility. Patient had a Leipzig score of 4, suggestive of WD. Genetic testing for ATP7B mutation was negative. Brain MRI revealed significant cerebral and cerebellar atrophy without parenchymal involvement. Liver biopsy was unobtainable due to elevated INR. After penicillamine, he had significant improvement in cognition and motor function. He was accepted to a liver transplant center.

Conclusions A history of alcohol abuse in a patient with liver failure usually leads to the suspicion of alcohol as the etiology. This case highlights the importance of a full workup to avoid anchoring bias. Our patient had a low ceruloplasmin and a high 24 hr urinary copper excretion which may suggest WD. Liver biopsy would be needed for confirmation. Early recognition and liver transplant evaluation are imperative in these patients.

## #454 COVID-19 RELATED LIFESTYLE CHANGE IS MULTIDIMENSIONAL

1K Morris*, 1M Bouchonville, 1S Fanchiang, 1Charles Drew University of Medicine and Science, Los Angeles, CA; 2University of New Mexico School of Medicine, Albuquerque, NM

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Purpose of Study Both, negative and positive lifestyle changes have been reported during the COVID-19 pandemic and many focused on particular areas of lifestyle. However, few studies have implemented multidimensional lifestyle measures. The work of cultural neuroscience has suggested that through self-reflection, a general pattern of self-enhancement motivation will be elicited regardless of the cultural background. We proposed that implementing a multidimensional lifestyle measure (the Short Multidimensional Inventory Lifestyle Evaluation - Confinement) with additional dimensions: Lifelong Learning, Nature/Environmental Connection, and Sense of Safety to explore its potential for clinical use, would lead to self-reflection beyond conventional measures to facilitate positive lifestyle change for health promotion and chronic illness management.

Methods Used We conducted a cross-sectional, anonymous online 40-item multidimensional survey focusing on lifestyle using social-media recruitment as a pilot. People 18 years or older were target survey respondents.

Summary of Results Among participants (n=37), 75% were female; more than more than 50% were from suburban areas, and about a third of them lived in urban areas. The results indicated changes in six dimensions with positive changes in Physical Activity, Lifelong Learning, and Sense of Safety but worse changes in Social Support and Screen Time. Whereas about equal proportion of the respondents reported positive and negative changes in Nature/Environmental Connection dimension. More than 50% of respondents indicated that taking this lifestyle survey gave insight into their lifestyle and 64% of them would do periodic lifestyle surveys or something similar to this multidimensional lifestyle survey for self-reflection.

Conclusions All three additional dimensions of the multidimensional lifestyle survey reflected lifestyle changes due to the COVID-19 pandemic beyond Physical Activity, Social
Support, and Screen Time dimensions. The proposed multidimensional lifestyle evaluation may be useful for self-reflection and further studies will be necessary to determine if the expanded lifestyle evaluation is superior for activating behavioral change.

#455 ANALYSIS OF TREATMENT GUIDELINES FOR HYPEREMESIS GRavidARUM AND NAUSEA/VOMITING OF PREGNANCY
A Morrow*, R McCann, B Brooks. Rocky Vista University College of Osteopathic Medicine, Livins, UT

Purpose of Study Hyperemesis gravidarum (HG) is the most severe form of nausea and vomiting of pregnancy (NPV), or morning sickness. 2% of pregnancies in the United States are affected by HG. The condition is characterized by severe vomiting in pregnant women, especially during the first trimester, often leading to hypovolemia and weight loss. The standard of care for HG and NPV is commonly ineffective. We hypothesize that based on patient experience, the current treatment guidelines for HG are not clinically effective. Our objective is to identify the efficacy of the management protocol that is currently in place for NVP and HG.

Methods Used A survey was designed based on diagnostic criteria as well as standard demographic identifiers and the most common medications used in the treatment of HG and was distributed to NPV and HG user groups on social media.

Summary of Results In our study, the vast majority of women diagnosed with HG trialed at least five medications, most of which were ineffective or had severe side effects.

Conclusions The data presented in this research provides insight into the suffering that patients with these diagnoses face day-to-day. Establishing this gap in treatment can facilitate the development of effective treatments that will provide relief for thousands of women.

This work was presented in part at Rocky Vista University Appreciation Day (virtual) on October 15, 2021

#456 ACUTE PROMYELOCYTIC LEUKEMIA OF THE EYE
R Dunn*, F Venter, S Mishra. Kern Medical Center, Bakersfield, CA

Purpose of Study Acute promyelocytic leukemia (APML) often presents with systemic symptoms such as weakness, fatigue, infections, ecchymoses and bleeding. It is usually associated with the translocation of t(15;17) causing a defect in the Retinoic Acid Receptor (RAR). However, this case reviews a 37-year-old male previously treated for Acute Myeloid Leukemia (AML) who presented with APML of the eye, confirmed by biopsy. This case stresses the importance of including a fundoscopic eye exam for possible ocular involvement – which is not uncommon in APML and may easily be missed.

Methods IRB approval was obtained, and single case review was done.

Summary of Results A 37-year-old Hispanic male presented with progressively worsening generalized weakness, fatigue, shortness of breath and easy bruising for 1 week. Upon admission, patient was hemodynamically stable and afibrile. Labs were concerning for complete blood count with differential showing megalocytes at 0.2x10^3/mcL, promyelocytes of 0.1x10^3/mcL, and blasts of 0.1x10^3/mcL. D-Dimer was also elevated at greater than 5000 fibrinogen equivalent units/mL. Peripheral smear was obtained which showed blast cells with Auer rods, suggesting APML. Diagnosis was confirmed with flow cytometry showing CD117 positive atypical/immature myeloid cells. FISH analysis showed t(15;17) translocation. Patient was started on All-trans Retinoic acid (ATRA) 60mg BID on with remission of APML. However, the patient developed left orbital swelling and pain on treatment day 7. CT Maxillofacial w/Contrast was obtained showing a rim-enhancing fluid collection. Ophthalmology consult was attempted but could not be done due to accessibility. Patient underwent biopsy of the eye at higher center and was consistent with APML. Patient was re-admitted at our facility and restarted on ATRA and intrathecal chemotherapy with great response.

Conclusion Acute leukemias generally present with systemic presentations like fever, body aches, fatigue and bleeding in APML. Ocular involvement can be due to direct infiltration of the leukemia or secondary to some blood abnormality like anemia, thrombocytopenia, or leukocytosis. Leukemic retinopathies are present in around 35.4% of leukemia patient and may present manifest after induction of chemotherapy. Clinicians should be vigilant to look for ocular involvement to prevent blindness.

#457 REPARATION OF THE ION GRADIENTS: THE ROLE OF ION CHANNELS AND THEIR POTENTIAL TO GUIDE THE CLINICAL MANAGEMENT OF EXTERNALLY INDUCED SKIN WOUNDS
1K Ma*, 2M Zhao, 1Western University of Health Sciences, Pomona, Ca; 2University of California Davis, Sacramento, CA

Purpose of Study The skin holds unique ion gradients that play a vital role in the process of wound healing. While the physiology behind this mechanism remains unclear, recent innovations regarding cutaneous ion channels may provide a better understanding of this biophysiological process. This is paramount because while molecular channel immunohistology has been utilized in forensic pathology to differentiate wound age postmortem, the same has not yet been reviewed for ion channels in skin wounds. Thus, a deeper comprehension of ion channels is pertinent in not only understanding ion gradient associated wound healing but would also provide a clinical significance in its potential to differentiate wound stage and wound types.

Methods Used A comprehensive literature search was used to search for over 100 studies involving ion channels in the context of skin wounds. Journal articles were reviewed from PubMed, ScienceDirect, and Google Scholar. Inclusion criteria limited the review to ion channels studied in the context of ion gradients, wound stage determination, and types of skin wounds.

Summary of Results Calcium, potassium, water, sodium, and chloride gradients have been found in the skin allowing for the identification of nine channels/pumps matching our inclusion criteria. These include Kcnj8, Kcnh2, TRPV1, TRPV2,
TRVP3, TRVP4, Orai1, Na+/K+ ATPase, and AQP3 channels. Kcnh2 and Kcnj8 were found in association with the homeostatic phase of wound healing, AQP3 channels in association with the proliferative phase, and TRVP2 in association with the remodeling/maturation phase. Ion channels also have potential for differentiating different wound types. These include TRVP3 and AQP3’s association with burn wounds and Orai1’s association with UVB-induced wounds.

Conclusions Potential channels involved in the repair of ion gradients following externally induced skin wounds were identified based on their variability and timing of expression following the event. These characteristics provided further evidence to our proposed feedback relationships between the gradients and the notion that they may play a clinical role in differentiating wound stages and types. However, there has yet been a channel identified as a main contribution to the inflammatory stage. Instead, a combination of decreased Na+/K+, Kcnh2, and Kcnj8 expression may allow for differentiation of this stage. The ability to differentiate wounds would provide improved diagnostic accuracy in the clinical context and allow for the personalization of treatment options. Given the role of ion fluxes and resultant wound electric fields in wound healing, temporal and spatial expression of ion channels and pumps is expected to offer a significant understanding of the powerful mechanism of electric signaling in wound healing.

[Necrotizing Fasciitis: A Potential Nidus for Hepatorenal Syndrome

#458

NECROTIZING FASCIITIS: A POTENTIAL NIDUS FOR HEPATORENAL SYNDROME

1K Ma*, 2K Chen, 3R Li, 4P Flores. 1Western University of Health Sciences, Pomona, CA; 2Hemet Global Medical Center, Hemet, CA

10.1136/jim-2022-WRMC.A54

Case Report Studies proposed bacterial translocation from the GI tract as a potential source of hepatorenal syndrome (HRS). While this has been reported in a few cases, they occurred in the setting of liver trauma associated endogenous infections. Here we report a case of HRS following the onset of necrotizing fascitis (NF). Overall, suggesting that in the absence of liver trauma, spread from skin wounds may contribute to the onset of HRS.

A female patient presents with bleeding from NF extending from her bilateral flanks to her upper thighs and a history of alcoholic cirrhosis. Kidney functions were originally within normal limits, however, reached 4.40mg/dL by day 18.

Despite trials of albumin, midodrine and withholding diuretics, creatinine levels continually rose, suggesting type 1 HRS. NF was also identified through cultures isolating Enterococcus Faecalis, Staphylococcus Aureus and Klebsiella Oxytoca. CT scan further demonstrated skin interruptions, ulcers and anasarca affected regions. Routine wound care, surgical debridement, and antibiotic treatment with ceftriaxone and daptomycin was utilized.

Aside from NF, there were no other identifiable causes of HRS. The patient’s bleeding event on admission occurred closely to her onset of HRS. This suggests that hematogenous/contiguous spread of cutaneous infections or bacterial products may be responsible for the onset of HRS in those with preexisting liver diseases. We suggest early utilization of barrier therapies and antimicrobial dressings. This case further questions whether only skin wounds close to the kidneys contribute to HRS or if distal wounds may also be influential.

A RARE CASE OF ACUTE PANCREATITIS DUE TO EPSTEIN BARR VIRUS INFECTION

1M Tu*, 2V Marquez, 3H Ilpawatwe. 1Clinica Sierra Vista, Bakersfield, CA; 2Ross University School of Medicine – Barbados Campus, Bridgetown, Barbados

10.1136/jim-2022-WRMC.A55

Case Report A 17yo male with a history of fatty liver disease presented to our emergency room with a 1-day history of severe left upper and lower epigastric pain. The patient was normotensive and afibrile, with a soft nontendstened abdomen and LUQ, epigastric and LLQ tenderness with no guarding, rebound tenderness, no bruising noted. Laboratory data revealed a total white blood cell count of 13.1x103/mcl, hemoglobin of 15.0 g/dl, platelet count of 240 x10^3/ml, prothrombin time of 15.4 s, international normalized ratio (INR) of 1.23, aspartate aminotransferase of 46 U/l, alanine aminotransferase of 104 U/l, alkaline phosphatase of 148 U/l, total bilirubin of 1.2 mg/dl, and lipase of 2,431 U/l (normal range: 0—160 U/l). Computed tomography (CT) of the abdomen and pelvis revealed acute necrotizing pancreatitis with a small unorganized acute necrotic collection, edematous pancreas, and hepatomegaly. Magnetic resonance imaging (MRI) showed necrotizing pancreatitis with moderate peripancreatic, perisplenic, and peripancreatic inflammatory fluid with no drainable fluid collection identified. No biliary ductal dilatation. No evidence of cholelithiasis or choledocholithiasis. Hepatomegaly. Splenomegaly suggesting underlying portal hypertension. On hospital day 2 patient had a temperature max of 38.6C, tachycardic in the 120s, tachynpeic in the 30s. SIRS criteria were met and septic workout was initiated and the patient started on Meropenem, with pancreatic fluid as a possible source of infection. Patient was treated for pancreatitis with guideline direct medical therapy. Viral hepatitis profiles (A, B, C), HIV RNA, and monospot test were all negative. The diagnosis of EBV infection was made by the positive result.

Abstract #458 Figure 1
of EBV immunoglobulin G(IgG)(VCA, viral capsid antigen) 129.00 unit/mL. Labs additionally showed CMV antibody IgG 3.10 unit/mL.

Discussion EBV-induced acute pancreatitis is a rare condition, the sooner an etiology can be deduced, the more options that a patient has in terms of therapy. Our patient has EBV-induced acute pancreatitis, a condition often not seen in conjunction. He did not exhibit any symptoms of infectious mononucleosis at the time of presentation, which is commonly associated with EBV infection. He deviates from many common associations. Is. With this case, we wanted to emphasize the importance of being vigilant, have broad differentials and early intervention based on clinical suspicion. For the best possible patient outcomes.

TACKLING SEVERE INTRAVENTRICULAR HEMORRHAGE IN A SINGLE CENTER LEVEL IV NICU

E Squire*, J Reiss, L Bain. Stanford University School of Medicine, Stanford, CA

10.1136/jim-2022-WRMC.456

Purpose of Study Premature infants <32w gestational age are at increased risk of severe (grade 3, 4) intraventricular hemorrhage (IVH) due to immature cerebral vasculature and decreased autoregulation of cerebral blood flow. Severe IVH is associated with increased long-term morbidity and mortality in this population. There are numerous modifiable risk factors and protective factors in the antenatal, intrapartum, and postnatal time periods that are associated with decreased rates of severe IVH. This quality improvement study examines modifiable risk factors of severe IVH at a single center level IV NICU, which noted a recent increase in severe IVH rates following several years of sustained reduction after implementation of a ‘Brain Care Bundle’ targeted at IVH prevention. A retrospective chart review of all inborn babies <32w gestational age from 2010 to 2020 who were diagnosed with either grade 3 or grade 4 IVH during their NICU stay demonstrated that some IVH risk reduction strategies such prenatal steroid use and elimination of the use of sodium bicarbonate have been consistently used over time, however other practices such as maintaining midline head positioning for the first 3 days of life and delayed umbilical cord clamping showed variability among patients. Inconsistent use of best practices provides an opportunity to use quality improvement strategies to reenforce elements of our Brain Care Bundle and reduce severe IVH rates once again. Our goal is to reduce severe IVH rates in inborn infants <32 weeks GA by 50% from 10% to 5% over the next 2 years consistent with rates in this population across other California NICUs. We will do this by re-educating and re-enforcing our current Brain Care Bundle for NICU providers and identifying ways to improve adherence to IVH reduction strategies. We also plan to analyze additional data, specifically CO2 levels in the first week of life, in hopes of adding novel interventions to our Brain Care Bundle. We will then monitor provider compliance with our updated Brain Care Bundle as well as overall rates of severe IVH to determine the impact of our interventions.

Methods Used

Summary of Results As above

Conclusions As above

PROTECTIVE MECHANISM OF BERBERINE ON SPINAL CORD INJURY IN RATS

H Duan, C Hao, X Yang*. The First Hospital of Shanxi Medical University, Taiyuan, China; Shanxi Provincial Key Laboratory of Brain Science and Neuropsychiatric Diseases, Taiyuan, China

10.1136/jim-2022-WRMC.457

Purpose of Study It is an important cause of neurological dysfunction for secondary inflammation after spinal cord injury. At present, there are relatively few effective drugs as an inflammatory inhibitor of spinal cord injury. Berberine is a common inflammatory inhibitor of traditional Chinese medicine extract, so we tried to explore the therapeutic significance of this drug for spinal cord injury.

Methods Used

The SD rats were randomly divided into three groups: sham operation group (n=30), non-intervention group(n=30), and Berberine intervention group(n=30). The T10 SCI model was established by Allen’s percussion method.BBB score,SEP and MEP latency and amplitude were used to evaluate hind limb function. The water content of spinal cord was measured. The spinal cell structure was observed by HE and Nissl staining and electron microscopy. The motor function was detected by SEP(Somatosensory Evoked Potential) and MEP (motor evoked potential. BBB (Basso-Beattie-Bresnahan) score was used to evaluate the hind limb function of rats.

Summary of Results

The BBB score of berberine in SCI model was significantly lower than that in sham operation group from postoperative day 1 to 31. Hindlimb function of Berberine intervention group began to recover faster than the non-intervention group on day 10 (p < 0.05). The latency of SEP and MEP after spinal cord injury was significantly longer than that of sham operation group (p < 0.05), but there was no significant difference in the latency of SEP and MEP between the two groups (p > 0.05). The amplitude of SEP and MEP in rats after spinal cord injury was significantly lower than that in the sham operation group (P < 0.05). In addition, after berberine treatment, the degree of spinal cord edema in rats decreased to some extent (p < 0.05).

Conclusions Berberine can promote the recovery of injured spinal cord to some extent, which may be related to its inflammatory inhibition.