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Cardiac Troponin Measurement in the Critically Ill: Potential for Guiding Clinical Management

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Background: Elevated cardiac troponin (cTn) in the absence of acute coronary syndromes (ACS) is associated with increased mortality in critically ill patients. There are no evidence-based interventions that reduce mortality in this group.

Objectives: We performed a retrospective investigation of the Veterans Administration Inpatient Evaluation Center database to determine whether drugs used in ACS (β -blockers, aspirin, and statins) are associated with reduced mortality in critically ill patients.

Methods: Thirty-day mortality was determined for non-ACS patients admitted to any Veterans Administration Intensive Care Unit between October 1, 2007, and September 30, 2008, adjusted for severity of illness. Troponin assay values were normalized across institutions.

Results: Multivariate analyses for 30-day mortality showed an odds ratio (OR) of 1.82 for patients with high cTn ($P < 0.0001$, cTn $> 10\%$ coefficient of variation) and 1.18 for intermediate cTn ($P = 0.0021$, cTn between lowest limit detectable and 10% coefficient of variation) compared with patients with no elevation, adjusting for severity of illness ($n = 19,979$). Logistic regression models showed that patients with no or intermediate elevations of cTn taking statins within 24 hours of cTn measurement had a lower mortality than patients not taking statins (OR, 0.66; 95% confidence interval [95% CI], 0.53–0.82; $P = 0.0003$), whereas patients with high cTn had a lower mortality if they were taking β -blockers or aspirin within 24 hours of cTn measurement compared to patients not taking β -blockers or aspirin (β -blockers: OR, 0.80; 95% CI, 0.68–0.94; $P = 0.0077$; aspirin: OR, 0.81; 95% CI, 0.69–0.96; $P = 0.0134$).

Conclusions: This retrospective study confirms an association between elevated troponin and outcomes in critically ill patients without ACS and identifies statins, β -blockers, and aspirin as potential outcome modifiers in a cTn-dependent manner.

Key Words: cardiac troponin, critically ill, β -blocker, aspirin, statin

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Critically ill patients who have elevated cardiac troponin (cTn) levels and no evidence of acute coronary syndrome (ACS) have a 2- to 4-fold increase in early mortality as compared with critically ill patients who do not have elevated cTn.^{1–5} Increased short-term mortality in patients with elevated cTn is observed in patients with sepsis,^{2,5,6} head trauma,⁷ and other critical injuries.⁷ This trend is also observed in critically ill children.⁸ The mechanism of cTn release in critically ill patients is poorly understood.⁹ The major concern when cTn is found to be elevated is whether rupture of a coronary arterial plaque has occurred and if the patient requires urgent coronary angiography and revascularization. Despite the prevalence of elevated cTn in this setting, obstructive coronary disease is infrequently found in the adult¹ and is not thought to be a likely occurrence in pediatric patients. Furthermore, comorbidities found in these cohorts (hypotension, renal failure, sepsis) often preclude invasive testing. In fact, there are no evidence-based treatments known to reduce mortality in this high risk population.¹⁰ Current practice is to medically manage patients with elevated cTn using medications known to reduce mortality in ischemic myocardial injury— β -blockers, statins, aspirin—however, there is little evidence to support this practice.¹⁰ The aim of this investigation was to determine 30-day mortality in critically ill patients with elevated cTn. We hypothesized that treatment with β -blockers, aspirin, or statins would be associated with reduced 30-day mortality.

METHODS

Patients and Study Design

The protocol for this investigation was approved by the University of Cincinnati Investigational Review Board. This retrospective cohort study utilized the Veterans Affairs Inpatient Evaluation Center (IPEC) centralized database. The IPEC is a national quality management program developed to measure risk adjusted mortality and evidence-based processes in inpatient care. The cohort consisted of patients admitted to any VA medical or surgical intensive care unit (ICU) between October 1, 2007, and September 30, 2008. Patients were excluded if they were admitted for an ACS, or if they had any prior ICU admission for an ACS within 1 year of ICU admission (Fig. 1). The ACS patients were excluded because mortality is known to be improved by the use of β -blockers, aspirin, or statins in this group. When patients had multiple ICU stays within the year, only the first ICU stay was analyzed. The overall severity of illness was determined for this cohort as described below ($n = 61,863$). Patients who had other known cardiac disease were excluded from the mortality analysis as treatment often involves β -blockers, aspirin, or statins ($n = 7734$). Lastly, we focused on patients who had cTn measured and for whom cTn ranges could be determined ($n = 19,979$) for the mortality analysis. Surgical procedures were assessed as part of the severity of illness. Barcode Medication Administration files were used to create indicators to determine if a patient was given β -blockers, aspirin, and/or statins within 24 hours of troponin elevation. Appendix 1 provides a list of the medications identified

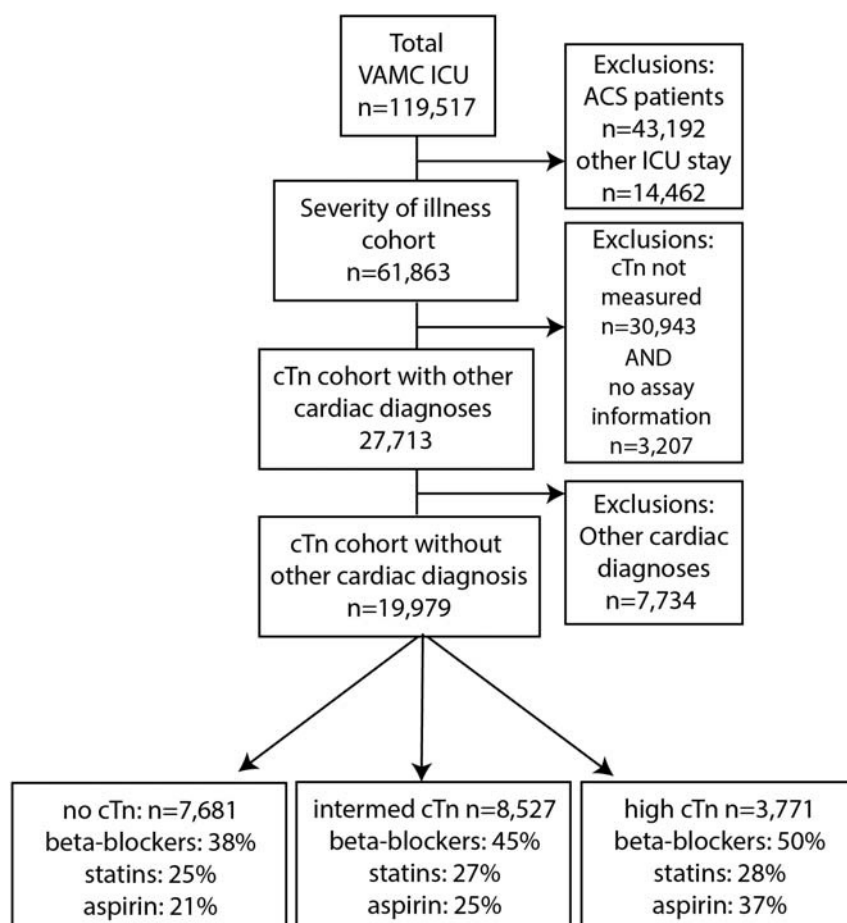


FIGURE 1. Branching algorithm describing VA-IPEC ICU patient population October 2007 to September 2008.

by this method. The backward elimination logistic regression model was used to statistically adjust for drug interactions among these 3 classes of medication.

Troponin Measurements

Both cTn T (cTnT) and cTn I (cTnI), components of a protein complex required for myocardial contraction, are well documented indicators of myocardial damage.¹¹ The cTnT assays are standardized across institutions, however, cTnI assays are not. We therefore normalized values for cTnI using the appropriate manufacturer assay information. Appendix 2 provides a detailed list of the assays used and their reference ranges ($\mu\text{g/L}$). The peak cTnI or cTnT measured within 24 hours from admission to discharge from the ICU was extracted. Facilities were sent a survey asking them to identify which troponin assay they were using. Standard ranges for each assay were used across all facilities. If this information was not available, these patients were excluded from the mortality analysis ($n = 3207$). The cTnI below the lowest limit detectable was classified as *no elevation*, levels between the lowest limit detectable and the 10% coefficient of variation (CV) were classified as *intermediate elevation*, and levels greater than the 10% CV were classified as *high elevation*. This classification was used to normalize cTn values across a range of values familiar to practitioners, not to classify patients according to the presence or absence of myocardial infarction. The 10% CV for cTnI is the lowest concentration of cTnI in which the assay has a CV of 10% or less. The CV is a measure of variability that adjusts for the magnitude of the mean between

assays, as compared to the standard deviation. The 10% CV is determined by assay manufacturers using samples of cTnI of known concentrations and is a measure of precision of the assay. For assays contemporary with this investigation, cTnI concentrations at the 99th percentile (accepted definition of myocardial infarction)¹¹ were within the “intermediate range” (see Appendix 2), and the cTnI concentration for the 10% CV was greater than the 99th percentile cTnI concentration.

A large group of patients (30,943) did not have cTn measured and were not included in the mortality analysis; ramifications of this are discussed in the Limitations section. Appendices 3 and 4 present baseline characteristics between those with nondefinable troponin ($n = 34,150$) and those with definable troponin ($n = 27,713$).

Study Outcomes

The primary outcome was 30-day mortality with respect to aspirin, β -blocker, or statin use in each group: patients with negative cTn, intermediate cTn, or high cTn.

Severity of Illness Model

The IPEC uses a statistical model to predict the probability of mortality. This ICU risk-adjusted model was modeled after APACHE III and has been validated against and used with VA data since 1995.^{12–15} This model includes laboratory values primarily, but also the primary diagnosis/procedure on admission to the ICU, comorbid diagnoses, use of immunosuppressant drugs,

TABLE 1. Variables Used to Determine the Severity of Illness in the IPEC Model

Serum albumin, g/L
Serum bilirubin, mg/dL
Blood-urea-nitrogen, mg/dL
Hematocrit, %
Blood glucose, mg/dL
Serum sodium, mEq/L
White blood cell count ($\times 10^3$ mm ³)
Partial pressure of arterial oxygen (PaO ₂)
Partial pressure of arterial carbon dioxide (PaCO ₂)
pH
Glomerular filtration rate (age, sex, and race adjusted)
Primary diagnosis/procedure on admission to the ICU
Age
Comorbid diagnoses
Use of immunosuppressant drugs
Source of admission into the ICU
Length of stay before the ICU stay

source of admission into the ICU, and length of stay before the ICU stay (Table 1). For each laboratory value, the most abnormal value within 48 hours of admission (24 hours before or after) was used. If there was no laboratory value within this time frame, the mean of the normal range for that value was substituted.¹³

Statistical Methods

Severity of Illness

A logistic regression model, similar to the IPEC model described above, was run on the cohort to obtain the predicted mortality for each patient at 30 days. This model was run because it was important to get an estimate of the severity of illness in a population without ACS. The logistic regression model assigns the probability of mortality to each patient (between 0 and 1). The linear predictor from this 30-day mortality was used as a covariate in subsequent troponin analyses to adjust for severity of illness. Laboratory values were modeled using restricted cubic splines, with the exception of PaCO₂ and pH which were combined and included as a categorical variable. Age was calculated at the time of hospital admission and modeled using a restricted cubic spline. Admission source was modeled as a categorical variable and

included the following sources: outpatient clinic, emergency department, nursing home, operating room, other hospital ward, acute care ward, and other. Immunosuppressant drug use was modeled as a dichotomous variable (1, on immunosuppressants; 0, not on immunosuppressants) using a list of immunosuppressant drugs in the pharmacy package. The length of stay before ICU was calculated as the amount of time between hospital admission and ICU admission and was modeled using restricted cubic splines. A dichotomous variable was created to indicate whether or not the patient had an International Classification of Diseases-9 procedure code within the 24 hours surrounding ICU admission.

Descriptive statistics (frequencies and means) were generated for each variable included in the risk adjustment model. Appendices 3 and 4 contain the categorical and continuous variables used to perform the risk adjustments, respectively. The average probability of mortality, referred to as severity of illness in these analyses, was calculated among troponin elevation categories and by medication use.

Troponin Levels and Mortality

Univariate analyses examining the relationship between 30-day mortality and troponin elevation were conducted using χ^2 tests. To determine if troponin is associated with mortality at 30 days after adjusting for severity of illness, a logistic regression model was run. This model included the linear predictor for each patient from the initial severity of illness calculation and troponin. Odds ratios (OR) and 95% confidence interval (95% CI) were calculated to compare mortality in patients with no troponin elevation to patients with high troponin or intermediate troponin elevation.

Medication Use and Outcome

Univariate analyses examining the relationship between mortality and medication use among each troponin elevation category were conducted using χ^2 tests. Initial multivariate analyses suggested a significant interaction between cTn level and medication use (likelihood ratio test based on χ^2 statistic of 9.837 with 3 degrees of freedom; $P = 0.0200$). Therefore, to determine if the use of β -blockers and/or statins and/or aspirin is associated with mortality reduction after adjusting for severity of illness, stratified logistic regression models were run. These models included the linear predictor for each patient from the initial severity of illness calculation, indicators for β -blockers, statins, and aspirin, and the interaction between β -blockers, statins, and/or aspirin. Stratified analyses were conducted among (1) those with no troponin elevation, (2) those with intermediate troponin elevation, and (3) those

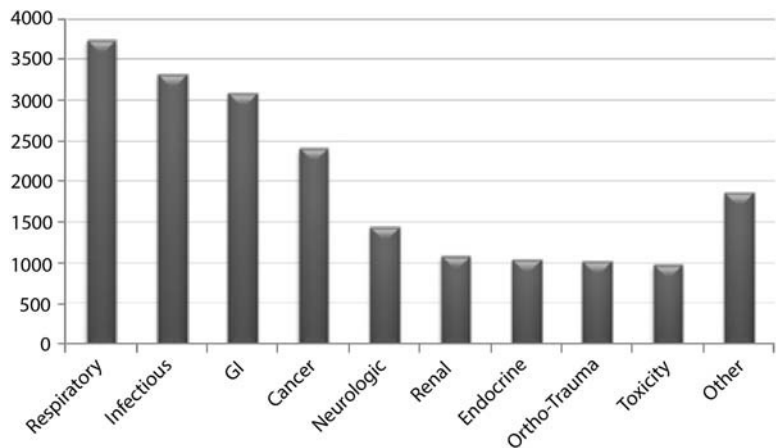


FIGURE 2. Distribution of primary diagnoses admitted to VA ICUs October 2007 to September 2008 (n = 19,979).

TABLE 2. Descriptive Statistics Age and Severity of Illness Increase With cTn, Whereas GFR Decreases With cTn

	cTn Negative (n = 7681)	cTn Intermediate (n = 8527)	cTn High (n = 3771)
Age (SD), y	63.7 (12.5)	66.8 (12.4)	69.0 (12.2)
Severity of illness (SD)	0.09 (0.15)	0.14 (0.18)	0.22 (0.24)
GFR (SD)	73.9 (42.5)	64.8 (39.6)	53.5 (43.8)

with high troponin elevation. All 3 logistic regression models were run for 30-day mortality. Both main effects and backward elimination models were conducted. The backward elimination model differs from the main effects model in that it includes not only the main effects of β -blockers, statins, and aspirin but also the interactions among these medications. In addition, terms are forced out one by one because they no longer achieve statistical significance at the 0.05 level. The inclusion significance level for the final backward elimination models was 0.05. Odds ratios and 95% CI were calculated for all covariates remaining in the backward elimination model. The final significance level used for all inferences was adjusted to account for the multiple tests as a result of stratification. Analyses were adjusted for severity of illness. SAS Version 9.3 was used for all analyses.

RESULTS

Characteristics of Patients

The cohort was predominantly men (96.6%) and white (68.7%). The majority of the patients were admitted for a respiratory diagnosis (18.7%), followed by infection (16.6%), gastrointestinal (15.5%), and cancer (12%) (Fig. 2). Patients who were cTn-negative were younger and had better renal function (Table 2), factors that were adjusted for in the mortality analysis.

Troponin and Mortality

Figure 3 shows the univariate analysis illustrating troponin level and 30-day mortality, Table 3 shows the tabulated results from each group. About 10.7% of the patients died at 30 days if they had no cTn elevation, whereas 16.6% of patients died if they had intermediate cTn, and 31.7% died at 30 days if they had high

cTn ($P < 0.0001$). After adjusting for severity of illness, mortality among critically ill patients was proportional to cTn levels and statistically significant. Multivariate analysis showed that the odds of death were 1.82 times greater for patients with high cTn compared with no cTn elevation, and 1.18 times greater for patients with intermediate cTn compared with no cTn elevation ($P < 0.0001$), after adjusting for severity of illness. The differences between the intermediate and high cTn groups were also statistically significant. Patients who did not have a cTn level measured had a lower severity of illness than those patients who had the blood test: cTn not measured, mean severity of illness, 0.07 (SD 0.14, $n = 34,150$), cTn measured, mean severity of illness 0.13 (SD 0.19, $n = 27,713$).

Medication Use and Outcome

Univariate analyses showed an increase in mortality proportional to cTn level within each medication class. Thirty-day mortality in those with no cTn elevation, intermediate cTn, and high cTn was 31.3%, 38.5%, and 40.6%, respectively, for patients on β -blockers ($P < 0.0001$); 14.9%, 19.0%, and 22.7%, respectively, for patients on statins ($P < 0.0001$); and 15.9%, 21.4%, and 31.2%, respectively, for patients on aspirin ($P < 0.0009$).

The results from the main effects model indicate that among those with no cTn elevation, statins were associated with lower mortality at 30 days (Table 4; OR, 0.68; 95% CI, 0.54–0.86). In those with an intermediate cTn elevation, statins were again associated with lower mortality at 30 days (OR, 0.69; 95% CI, 0.59–0.81). No relationships between mortality and β -blockers or aspirin were statistically significant for patients with negative or intermediate cTn. In contrast, among those with a high cTn elevation, statins were no longer statistically significant at 30 days, however, β -blockers (OR, 0.81; 95% CI, 0.68–0.95) and aspirin (OR, 0.81; 95% CI, 0.69–0.96) were associated with lower mortality. All analyses were adjusted for severity of illness. Inferences from the stratified models held up to multiple testing criteria.

The results from the backward elimination model were similar to the results from the main effects model (Table 5). In patients with no cTn or intermediate cTn elevation, statins alone were associated with lower mortality at 30 days (negative cTn: OR, 0.66; 95% CI, 0.53–0.82; intermediate cTn: OR, 0.69; 95% CI, 0.58–0.80). No relationships between mortality and β -blockers or aspirin were statistically significant for these groups. Patients with high cTn elevation, however, demonstrated an associated reduction in mortality with β -blockers (OR, 0.80; 95% CI, 0.68–0.94) and

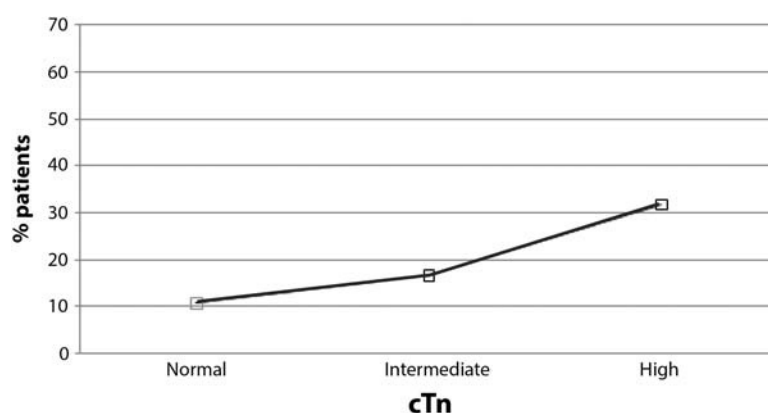
**FIGURE 3.** Relationship between 30-day mortality and troponin level: no cTn elevation, 10.7% 30-day mortality; intermediate cTn, 16.6% 30-day mortality; high cTn, 31.7% 30-day mortality.

TABLE 3. Effect of Troponin Level on Mortality: cTn Cohorts With Other Cardiac Diagnoses (27,713) and Without Other Cardiac Diagnoses (19,979)

Mortality, d	Effect	cTn Cohort With Other Cardiac Diagnoses		cTn cohort Without Other Cardiac Diagnoses	
		(n=27,713)		(n = 19,979)	
		Odds Ratio (CI)	P	Odds Ratio (CI)	P
30	Troponin—intermediate vs none	1.18 (1.07–1.30)	0.0012	1.18 (1.06–1.31)	0.0021
	Troponin—high vs none	1.79 (1.61–2.00)	<0.0001	1.82 (1.62–2.04)	<0.0001
	Troponin—high vs intermediate	1.53 (1.39–1.68)	<0.0001	1.54 (1.39–1.71)	<0.0001

Odds ratios adjusted for severity of illness.

Severity of illness was statistically significant in all models ($P < 0.0001$).

aspirin (OR, 0.81; 95% CI, 0.69–0.96) at 30 days, with no relationship between mortality and statin use.

DISCUSSION

The purpose of this investigation was to determine whether treatments administered to patients with cTn elevation in the setting of critical illness are associated with improved short-term mortality. This large retrospective study demonstrated 2 major findings: first, that cTn is associated with 30-day mortality in critically ill patients in the absence of ACS, and second, that statins, aspirin, and β -blockers were associated with decreased mortality in a cTn-dependent manner.

Troponin as an Independent Marker of Mortality

The relationship between cTn and mortality in critically ill patients is not new,^{1,3,4,8,16–19} but not all published work is in agreement.²⁰ A limitation of studies using cTnI as a marker is that there are many cTnI assay manufacturers. The decision to use the 99th percentile of cTnI or T to consider a sample suggestive of myocardial infarction has been set by consensus,¹¹ but the actual value of cTnI can vary based on the assay used. Cardiac troponin T values, however, are uniform because the assay is produced by a single manufacturer. One strength of this analysis is that cTnI values were normalized based on established cutoffs determined

by the assay used, and this provided a large sample of patients within each cTnI range. This work and the majority of published studies support the finding that cTn elevation in critically ill patients is associated with increased mortality. Increasing mortality risk from no cTn elevation to intermediate to high cTn elevation also suggests that cTn may be a continuous marker of mortality risk, as it is in patients with an ACS.²¹

The timing of cTn assessment may impact the observed result. This analysis used the highest cTn level that occurred on admission to the ICU. In a recent prospective observational study of septic patients, cTnI was highest on the first day of admission and decreased thereafter.²² It is likely that our analysis included the earliest, therefore highest, cTn measurement.

The role of high sensitivity cTnT testing (hs-cTnT) in identifying ICU patients at increased mortality risk is not clear. The high sensitivity assays can detect as little as 10 ng/L of troponin in the serum as opposed to a lower limit of detection of 32 ng/L in contemporary cTnI testing.²³ It is expected that high sensitivity testing would identify a larger group of at-risk patients who may benefit from the addition of aspirin, β -blocker, and statin. In a recent prospective observational study, hs-cTnT was measured in 144 consecutive ICU patients.²⁴ Sixty-two patients (43%) had evidence of hs-cTnT elevation without findings of an ACS, a larger group than was found in a previous study using the contemporary assays (19%).¹⁶ The non-ACS patients identified with hs-cTnT testing had a

TABLE 4. Effect of Medication Use on Mortality According to cTn Level—Main Effects Model

Troponin Elevation	Mortality, d	Effect	cTn Cohort With Other Cardiac Diagnoses		cTn Cohort Without Other Cardiac Diagnoses	
			(n = 27,713)		(n = 19,979)	
			Odds Ratio (95% CI)	P	Odds Ratio (95% CI)	P
None	30	On β -blocker	0.89 (0.75–1.06)	0.1899	0.93 (0.77–1.12)	0.4310
		On statin	0.66 (0.53–0.82)	0.0001	0.68 (0.54–0.86)	0.0011
		On aspirin	0.87 (0.71–1.07)	0.1999	0.91 (0.72–1.15)	0.4287
Intermediate	30	On β -blocker	0.94 (0.83–1.06)	0.2994	0.98 (0.85–1.12)	0.7440
		On statin	0.70 (0.61–0.81)	<0.0001	0.69 (0.59–0.81)	<0.0001
		On aspirin	0.92 (0.80–1.06)	0.2480	0.99 (0.84–1.16)	0.8507
High	30	On β -blocker	0.82 (0.70–0.94)	0.0063	0.81 (0.68–0.95)	0.0090
		On statin	0.99 (0.84–1.17)	0.9095	0.99 (0.82–1.19)	0.9105
		On aspirin	0.77 (0.66–0.90)	0.0007	0.81 (0.69–0.96)	0.0165

cTn cohorts with other cardiac diagnoses (27,713) and cTn cohort without other cardiac diagnoses (19,979). Odds ratio adjusted for severity of illness. Severity of illness was statistically significant in all models ($P < 0.0001$).

TABLE 5. Effect of Medication Use on Mortality According to cTn Level—Backward Elimination Model

Troponin Elevation	Mortality, d	Effect	cTn Cohort With Other Cardiac Diagnoses		cTn Cohort Without Other Cardiac Diagnoses	
			(n = 27,713)		(n = 19,979)	
			Odds Ratio (95% CI)	P	Odds Ratio (95% CI)	P
None	30	On statin	0.63 (0.51–0.77)	<0.0001	0.66 (0.53–0.82)	0.0003
Intermediate	30	On statin	0.68 (0.59–0.78)	<0.0001	0.69 (0.58–0.80)	<0.0001
High	30	On β -blocker	0.81 (0.70–0.94)	0.0053	0.80 (0.68–0.94)	0.0077
		On aspirin	0.77 (0.67–0.89)	0.0005	0.81 (0.69–0.96)	0.0134

cTn cohorts with other cardiac diagnoses (27,713) and cTn cohorts without other cardiac diagnoses (19,979). Odds ratios adjusted for severity of illness. Severity of illness was statistically significant in all models ($P < 0.0001$).

*Only terms significant at the 0.05 level are displayed.

23% risk of dying in the ICU as compared to patients with no hs-cTnT elevation (23 patients, 4.4% risk of ICU mortality). The hs-cTnT investigation, however, did not adjust for severity of illness (cTn levels correlate with severity of illness as found in this investigation) nor exclude for prior cardiac disease (16% of patients with hs-cTnT elevation had known ischemic heart disease). Patients with known ischemic heart disease should already have aspirin, β -blockers, and statin prescribed, as tolerated. It is not clear, therefore, that high sensitivity troponin levels would increase the number of at risk patients identified, or add further refinement to this question. Prospective analyses that exclude for prior ischemic heart disease and adjust for severity of illness will be needed to address whether hs-cTnT adds additional information to identify at-risk patients in the ICU.

Outcome Modification Associated With Medical Therapy

This analysis suggests that measurement of cTn in critically ill patients identifies a treatment pathway associated with reduced 30-day mortality (Fig. 4), although a prospective analysis including all at-risk patients and excluding all potentially confounding cardiac diseases is necessary to confirm this result. Critically ill patients with either no elevation of cTn or those with an intermediate value who are treated with statins had a significantly reduced mortality risk than those patients not treated with statins. In addition, patients with high cTn had lower associated 30-day mortality if they were treated with aspirin, β -blockers, or both, than cTn high patients who were not treated with aspirin or β -blockers. This suggests that cTn measurement not only identifies a mortality risk, but also indicates a potential treatment plan that is associated with reduced 30-day mortality. To our knowledge, this has not been previously reported.

Our findings are supported indirectly in the literature. A prospective, randomized, open-label study of 154 patients in septic shock showed that the β -blocker esmolol, titrated to keep heart rate between 80 and 94 beats per minute, resulted in an overall survival benefit (HR, 0.392; 95% CI, 0.261–0.590; $P < 0.001$), after adjustment for severity of illness.²⁵ Although cTn levels were not measured in this study, the Simplified Acute Physiology Score II was 52 in the treatment group, a value that predicts 50% mortality.²⁶ In our investigation, the degree of troponin elevation correlated with severity of illness. The high cTn cohort had a mortality of 31.7%, a severity of illness that was similar, if not more favorable, than the septic shock group of the Morelli et al investigation. Aspirin is also associated with reduced mortality in 2 large retrospective investigations of patients presenting with severe sepsis.^{27,28} Had

cTn levels been measured in these subjects, it is likely that the majority would have fallen into the high cTn group.

A meta-analysis of randomized controlled clinical trials of statins in septic patients,²⁹ and a systematic review that included observational studies,³⁰ provide conflicting results. Observational studies suggest a benefit from statin use, whereas randomized controlled clinical trials show no benefit in severe sepsis. It is possible that patients should first be assessed for cTn, as our data suggests. In a randomized, double blind, placebo controlled study of statin use in septic patients on the ward,³¹ 40 mg of atorvastatin prevented conversion of sepsis to severe sepsis (4% vs 24%, $P = 0.007$). The mean APACHE II score for this group correlated to an anticipated mortality rate of 15%. This was similar to the cTn none and cTn intermediate groups in our investigation (mortality rates 10.7% and 16.6%, respectively). Statins may therefore benefit a lower-risk group.

That statins may benefit patients who are less severely ill, whereas β -blockers and aspirin may benefit patients who are more severely ill, suggesting different mechanisms of disease process intervention in each group. Unpublished studies from this group suggest microvascular obstruction as a cause for end-organ damage in extremely ill patients. Thus, benefit from aspirin would make sense in that the antiplatelet effect may mitigate microvascular obstruction. The benefits from β -blockers may be largely cardioprotective by reducing cardiac work and myocardial oxygen demand, as is observed in patients presenting with a myocardial infarction.^{32,33} Significant cTn release indicates myocardial demand in excess of oxygen supply. β -blockers probably assist in reducing myocardial oxygen demand, thus may be cardioprotective. Benefit from statins in less severely ill patients points to intervention in an inflammatory process, before significant end organ damage has occurred.

Lastly, our investigation indicates that medications should be considered in all studies of mortality in critically ill patients,

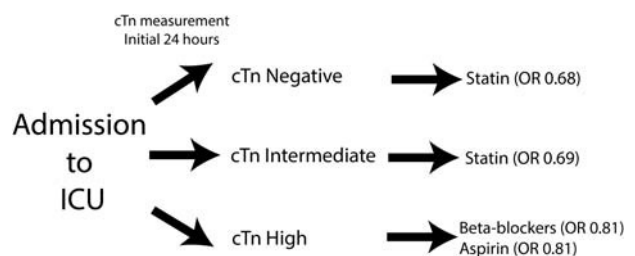


FIGURE 4. Measurement of cTn in critically ill patients may help guide potentially life-saving therapy.

as our data clearly show that medications are associated with outcome modification.

CONCLUSIONS

This large retrospective observational cohort study suggests that elevated cTn in critically ill patients without an ACS is an independent risk factor of 30-day mortality. Using 2 statistical models, our data further show lower 30-day mortality in both cTn-positive or -negative groups, depending on pharmacologic treatment. Patients with no or intermediate cTn elevation had a significantly lower associated risk of death if treated with statins, whereas cTn positive patients had a significantly lower associated risk of death if treated with aspirin and/or β -blockers. The presence of cTn in the blood of critically ill patients may identify populations of patients who benefit from different treatments, and in addition suggests a different pathophysiologic mechanism of increased mortality in each group. This work supports the need for prospective translational studies that investigate the mechanisms of improved mortality in each group.

LIMITATIONS

There are several important limitations to this investigation. This is a retrospective investigation of all patients who happened to get a cTn level drawn during admission. It is possible that some of the patients who had the test drawn may have had some cardiovascular complaint or sign reflecting a higher degree of illness. This is supported by our data. Patients who did not have cTnI measured were less severely ill than patients who had cTnI measured. Addition of patients with a lower severity of illness might impact our finding that statins improved the outcome in patients with no cTn; however, this remains to be shown in a prospective clinical trial. We were only able to assay medication use during the hospital stay. We do not know whether the medications were being taken before admission, or for how long the medications were administered after the cTn assay. We cannot conclude from our work that cTn is a continuous variable. A large prospective clinical trial could clarify this and may also provide cutoff values. Lastly, high sensitivity assays were not available for wide use during the period of data collection. As discussed, hs-cTnT assays may improve the resolution at the lower cTn levels, but this remains to be investigated.

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Appendix 1: Medications

Medication Type	Medication Class	Medication Name
β-Blockers	CV100	Atenolol
		Bisoprolol Fumarate
		C-Metoprolol
		Carvedilol
		Esmolol
		Laβlol
		Metoprolol
		Nadolol
		Pindolol
		Propranolol
Statins	CV350	Sotalol
		Atorvastatin
		Ezetimibe
		Fluvastatin
		Lovastatin
		Pravastatin
		Rosuvastatin
Aspirin	CN103	Simvastatin
		ASA/Butalbital/Caffeine
		Aspirin
		Choline/Trisalicylate

Appendix 2: Troponin Assays

Manufacturer	Assay	LLD (μg/L)	LLD-10%CV (μg/L)	10% CV (μg/L)	99% Cutoff/10% CV (μg/L)*
Abbott	AxSYM	<0.14	0.14–1.3	>1.3	0.3/1.22
Abbott	Architect	0.009	0.009–0.212	0.212	—
Bayer	ACS:180	<0.03	0.03–0.45	>0.45	0.1/0.37
Bayer	ACS Centaur	<0.02	0.02–0.45	>0.45	0.1/0.33
Beckman Coulter	Access/Access 2	<0.01	0.01–0.1	>0.1	0.04/0.09
Biosite	Triage	<0.19	0.19–0.69	>0.69	—
Dade Behring	Dimension RxL	<0.04	0.04–0.21	>0.21	0.07/0.26
Dade Behring	Stratus CS	<0.03	0.03–0.13	>0.13	—
DPC	Immulite	<0.2	0.2–0.8	>0.8	0.2/0.32
Ortho	Vitros	<0.038	0.038–0.2	>0.2	0.1/0.44
Roche	Elecsys	<0.01	0.01–0.04	>0.04	0.01/0.04
Abbott	ISTAT	<0.02	0.02–0.10	>0.10	—

*From Wu, Alan H. B. Chapter 2, *Contemporary Cardiology: Cardiovascular Biomarkers: Pathophysiology and Disease Management*. Ed. Morrow, D. A. 2006.

LLD indicates lowest limit detectable.

Appendix 3: Categorical Variables Used in Risk Adjustment

Characteristic	cTn Cohort With Other Cardiac Diagnoses (n = 27,713)		cTn Not Measured + No cTn Assay Information (n = 30,943 + 3207 = 34,150)		cTn Cohort Without Other Cardiac Disease (n = 19,979)	
	N	% of Study Cohort	N	% of Study Cohort	N	% of Study Cohort
Male*	26,677	96.3	34,150	94.8	19,304	96.6
Race*						
African American	5641	20.4	6436	18.9	3932	19.7
American Indian/Alaskan native	139	0.5	151	0.4	101	0.5
Asian/native Hawaiian/other	240	0.9	339	0.9	157	0.8
Pacific						
White	18,836	68.0	23,089	68	13,723	68.7
Unknown	2857	10.3	4134	12.1	2066	10.3
Source of Admission into ICU						
Ward	6769	24.4	5478	16.0	5274	26.4
Operating room	4793	17.3	16,087	47.1	3875	19.4
Other hospital	488	1.8	511	1.5	311	1.6
Clinic	6490	23.4	4409	12.9	4293	21.5
Nursing home	698	2.5	551	1.6	547	2.7
Psychiatric ward	190	0.7	161	0.5	156	0.8
Rehabilitation	94	0.3	104	0.3	76	0.4
Emergency department	8185	29.5	6837	20.0	5442	27.2
Unknown	6	0.0	12	0.0	5	0.0
On immunosuppressant	3916	14.1	3036	8.9	3139	15.7
Procedure within 24 h of ICU admission	5268	19.01	16,828	49.3	4285	21.5
Diagnosis						
Behavioral, mental, and psychiatric disorders	175	0.6	236	0.7	175	0.9
Cancers	2402	8.7	7079	20.7	2402	12.0
Cardiac/circulation	7734	27.9	4790	14.0	N/A	N/A
Complications	808	2.9	1468	4.3	808	4.0
Endocrine/nutritional diseases	1036	3.7	1182	3.46	1036	5.2
Gastrointestinal diseases	3086	11.1	4,437	13	3086	15.5
Genitourinary disease	212	0.8	691	2.0	212	1.1
Hematologic	282	1.0	294	0.9	282	1.4
Infectious	3310	11.9	2642	7.7	3310	16.6
Neurological	1441	5.2	2798	8.2	1441	7.2
Orthopedic and trauma	1029	3.7	2760	8.1	1029	5.2
Renal	1084	3.9	896	2.6	1084	5.4
Respiratory diseases	3742	13.5	2630	7.7	3742	18.7
Rheumatologic/connective	121	0.4	130	0.4	121	0.6
Tissue						
Substance abuse/toxicity	983	3.6	1426	4.2	983	4.9
Other	268	1.1	691	2.0	268	1.3
Comorbidities						
CHF	3527	12.7	1769	5.2	2515	12.6
Valvular disease	1277	4.6	725	2.1	648	3.2
Pulmonary circulation disorders	1022	3.7	479	1.4	659	3.3
Peripheral vascular disorders	1347	4.9	1504	4.4	953	4.8
Hypertension	14,565	52.6	16,246	47.6	10,219	51.2
Paralysis	931	3.4	937	2.7	788	3.9
Other neurological disorders	1691	6.1	1533	4.5	1410	7.1
Chronic pulmonary disease	5873	21.2	5501	16.1	4209	21.1
Diabetes, uncomplicated	6183	22.3	6280	18.4	4200	21.0
Diabetes, complicated	1437	5.2	1102	3.2	987	4.9

(Continued on next page)

Appendix 3: (Continued)

Characteristic	cTn Cohort With Other Cardiac Diagnoses (n = 27,713)		cTn Not Measured + No cTn Assay Information (n = 30,943 + 3207 = 34,150)		cTn Cohort Without Other Cardiac Disease (n = 19,979)	
	N	% of Study Cohort	N	% of Study Cohort	N	% of Study Cohort
Hypothyroidism	1437	5.2	1387	4.1	990	5.0
Renal failure	3202	11.6	2201	6.5	2287	11.5
Liver disease	1651	6.0	2177	6.4	1384	6.9
Peptic ulcer disease excluding bleeding	32	0.1	32	0.1	27	0.1
AIDS	164	0.6	252	0.7	134	0.7
Lymphoma	278	1.0	227	0.7	218	1.1
Metastatic cancer	1060	3.8	1454	4.3	943	4.7
Solid tumor without metastasis	1772	6.4	1857	5.4	1404	7.0
Rheumatoid arthritis/collvascular diseases	304	1.1	288	0.8	220	1.1
Coagulopathy	1377	5.0	1222	3.6	1145	5.7
Obesity	1296	4.7	1288	3.8	826	4.1
Weight loss	1044	3.8	926	2.7	892	4.5
Fluid and electrolyte disorders	6128	22.1	4540	13.3	5004	25.1
Blood loss anemia	289	1.0	346	1.0	256	1.3
Deficiency anemias	4154	15.0	3696	10.8	3228	16.2
Alcohol abuse	1980	7.1	2151	6.3	1541	7.7
Drug abuse	986	3.6	902	2.6	699	3.5
Psychoses	1617	5.8	1748	5.1	1229	6.2
Depression	1705	6.2	1924	5.6	1244	6.2

*Not used in risk adjustment.

Appendix 4: Continuous Variables Used in Risk Adjustment

Characteristics	Mean (SD) cTn Cohort With Other Cardiac Diagnoses (n = 27,713)	Mean (SD) cTn Not Measured + No cTn Assay Information (n = 30,943 + 3207) n = 34,150	Mean (SD) cTn Cohort Without Other Cardiac Disease (n = 19,979)
Age, y	66.0 (12.7)	62.7 (12.5)	66.1 (12.6)
Length of stay, d*	3.6 (5.8)	2.7 (4.0)	4.0 (6.4)
Length of stay before ICU, d	1.1 (2.7)	1.0 (2.4)	1.3 (2.9)
Physiologic variables			
White blood cell count, $\times 10^3 \text{ mm}^3$	11.3 (11.3)	10.5 (8.3)	12.1 (12.7)
Serum albumin, g/L	3.5 (0.9)	3.7 (0.8)	3.4 (0.9)
Serum bilirubin, mg/dL	1.1 (1.9)	1.1 (2.2)	1.2 (2.2)
Blood urea nitrogen, mg/dL	29.2 (24.8)	22.4 (20)	30.4 (26.3)
Serum creatinine, mg/dL	1.8 (1.8)	1.4 (1.6)	1.8 (1.9)
Blood glucose, mg/dL	163.8 (129.2)	147.2 (103.5)	171.8 (142.9)
Hematocrit, %	36.6 (7.1)	36.6 (6.7)	35.9 (7.3)
Serum sodium, mEq/L	136.7 (5.8)	137 (5.2)	136.4 (6.3)
Glomerular filtration rate	65.5 (39.4)	77.6 (38)	66.2 (42.2)
PaO ₂	93.5 (53.5)	103.7 (62.3)	93.4 (57.1)
PaCO ₂	41.4 (13.5)	40.1 (9.0)	42.0 (15.1)
pH	7.4 (0.1)	7.4 (0.1)	7.4 (0.1)

*Not used in risk adjustment.