Effects of non-surgical treatment of chronic periodontitis on insulin resistance and glucose tolerance in subjects without diabetes (PARODIA 2 study)

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ABSTRACT
This study aimed to assess the effects of non-surgical periodontal treatment (NSPT) of chronic periodontitis on insulin sensitivity, glucose tolerance, and serum C reactive protein (CRP) level in individuals without diabetes. Twenty individuals without diabetes with chronic periodontitis underwent NSPT, which consisted of complete scaling, polishing, root planing, and irrigation of the periodontal pockets with a 10% povidone-iodine solution. Periodontal indices (plaque index, gingival bleeding index, pocket depth, and clinical attachment loss), insulin sensitivity using the Short Insulin Tolerance Test index (KITT), glucose tolerance derived from oral glucose tolerance test, and serum CRP level were measured before and 3 months after the intervention. This study was conducted at the National Obesity Center of Yaoundé Central Hospital, Cameroon. After 3 months, we observed significant improvement in periodontal parameters (all p<0.001) and insulin sensitivity (3.72 (2.99–4.17) %/min before treatment vs 4.04 (3.67–4.78) %/min after treatment, p=0.001) and significant decrease in serum CRP level (2.35 (1.46–4.18) mg/L before vs 1.53 (1.03–2.12) mg/L after, p=0.033). There was a trend toward improvement in glucose tolerance, although not statistically significant after the intervention. This study suggests that NSPT of chronic periodontitis in individuals without diabetes is associated with increased insulin sensitivity and decreased serum CRP levels.

Trial registration number NCT02830113.

INTRODUCTION
Diabetes is one of the most common metabolic disorders globally, and type 2 diabetes mellitus (T2DM) is considered an epidemic and accounts for approximately 90% of total cases. Insulin resistance plays a crucial part in the pathogenesis of T2DM and prospective studies have demonstrated that it is the best predictor of diabetes. Chronic inflammation is one of the factors that can induce insulin resistance. Proinflammatory cytokines have been demonstrated to induce insulin resistance. Insulin resistance can therefore contribute to the development of T2DM.

On the other hand, reducing proinflammatory cytokines has been shown to improve insulin resistance. Chronic periodontitis is responsible for a long-lasting inflammation of the tissue surrounding the tooth. It brings forth a systemic inflammation even though the inflammation, in this case, is localized to the periodontal tissues. Treatment of periodontitis results in a reduction of the inflammation not only at the periodontal level but also at the systemic level, and this is marked by a reduction in serum inflammatory markers. C reactive protein (CRP) has been considered a possible mediator of the relationship between periodontitis and systemic conditions. CRP levels increase in direct proportion to severity of periodontitis, and a decrease in periodontal inflammation results in the same proportional reduction in CRP levels.

A possible reduction in severity of systemic diseases following periodontal therapy due to a decrease in inflammatory burden has been reported. We hypothesized that non-surgical periodontal treatment (NSPT), including scaling, root planing, and subgingival irrigation with povidone-iodine, would improve insulin sensitivity by reducing inflammation. Therefore, this study aimed to evaluate the effects of a non-surgical treatment of chronic periodontitis on inflammation (by CRP serum level measurements), insulin sensitivity, and glucose tolerance in an adult Cameroonian population without diabetes.

MATERIALS AND METHODS
All subjects provided written informed consent.

Setting and study population
This study was conducted at the National Obesity Center of Yaoundé Central Hospital, Cameroon. This was a quasi-experimental, single-arm study where each subject was his/her own control (pre and post treatment) that was conducted at the National Obesity Center of Yaoundé Central Hospital, Cameroon. This study was carried out at the National Obesity Center of Yaoundé Central Hospital, Cameroon. After 3 months, we observed significant improvement in periodontal parameters (all p<0.001) and insulin sensitivity (3.72 (2.99–4.17) %/min before treatment vs 4.04 (3.67–4.78) %/min after treatment, p=0.001) and significant decrease in serum CRP level (2.35 (1.46–4.18) mg/L before vs 1.53 (1.03–2.12) mg/L after, p=0.033). There was a trend toward improvement in glucose tolerance, although not statistically significant after the intervention. This study suggests that NSPT of chronic periodontitis in individuals without diabetes is associated with increased insulin sensitivity and decreased serum CRP levels.

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own control. We enrolled 24 adult subjects with mild to severe chronic periodontitis according to the 2012 Centers for Disease Control and Prevention in partnership with the American Academy of Periodontology (CDC-AAP) classification. Exclusion criteria were a condition capable of influencing any of the study parameters (diabetes, hypertension, pregnancy, arthritis), any clinically diagnosed condition which is contraindicated for any of the procedures undertaken in the study (allergies, cardiopathies), patients who underwent any periodontal treatment within 6 months preceding the study, and patients with less than 20 vital teeth. Alcohol users, smokers, and subjects taking medications that may affect insulin resistance/sensitivity and/or inflammation before and during the study period were also excluded.

Data collection
We performed an interview and a complete physical examination on all subjects. We measured anthropometric parameters, insulin sensitivity, glucose tolerance, and serum CRP level before and 3 months after NSPT. Periodontal parameters, including O’Leary’s plaque index (PI), Ainamo and Bay’s gingival bleeding index (GBI), pocket depth (PD), and clinical attachment loss (CAL), were also measured before and 3 months after NSPT. Our primary endpoint measurement was insulin sensitivity and glucose tolerance, and our secondary outcome was improvement in the periodontal parameters.

Anthropometric measurements
We measured height to the nearest 0.5 cm and weight to the nearest 0.1 kg and calculated the body mass index (BMI) using Quetelet index in all subjects. We measured waist and hip circumference to the nearest 0.5 cm and calculated the waist to hip ratio. Per cent body fat was measured by bio-impedancemetry (TANITA BC 420 MA, TANITA Corporation, Tabashi-ku, Tokyo, Japan).

Periodontal examination
PD and CAL using a periodontal probe (PCP UNC 12, Hu-Friedy Manufacturing, Chicago, USA) were made on six sites (distobuccal, buccal, mesiobuccal, mesiolingual, lingual, and distolingual) of all teeth except the third molars. PD was measured as the distance from the free gingival margin (GM) to the bottom of the periodontal pocket.

CAL was measured as the distance from the cementoenamel junction (CEJ) to the bottom of the sulcus or the periodontal pocket: $\text{CAL} = \text{PD} - (\text{CEJ} - \text{GM})$, where GM was subject to recession and CEJ was exposed; the distance from the CEJ to the GM was given a negative value, and a positive value when the GM covered the CEJ.

O’Leary’s PI was used to evaluate the quantity of dental plaque and Ainamo and Bay’s GBI to evaluate gingival inflammation. Four sites (mesiobuccal, buccal, distobuccal, and lingual) were evaluated for each tooth except the third molars, and PI and GBI were expressed as the percentage of positive sites of the total number of examined sites.

Metabolic assessments
Insulin sensitivity measurement
Insulin sensitivity was evaluated using a short insulin tolerance test (SITT). For SITT, a fixed bolus of regular insulin dose of 0.15 IU/kg body weight was intravenously injected into the fasting subjects. Plasma glucose levels were measured every 3 min for 15 min. The Short Insulin Tolerance Test Index (KITT) which represents the slope of the linear decline in plasma glucose was then calculated by dividing 0.693 by the plasma glucose half-life (50% from baseline), denoted as $t^{1/2}$.

Glucose tolerance measurement
Glucose tolerance was measured using the oral glucose tolerance test (OGTT). A 75 g OGTT was performed over 120 min after a 10-hour overnight fast. Blood samples were collected from an antecubital vein before and 120 min after the 75 g oral glucose load for glucose determination. Impaired fasting glucose and impaired glucose tolerance were defined according to the definition of the American Diabetes Association.

Biochemical measurements
Immunoturbidimetric method with agglutination to latex was used for CRP dosing using a CRP turbidimetric CROMATEST dosage kit (Linear Chemicals, Barcelona, Spain). Plasma glucose was measured by the hexokinase method (Roche Diagnostics, Mannheim, Germany).

Periodontal treatment
Mechanical therapy was completed at a single unlimited time appointment. Patients were trained and given instructions on oral hygiene practices and received oral hygiene kits, after scaling, polishing, and root planing of all teeth were done. The scaling was performed with a magnetostrictive ultrasonic device (Denjoy DUS-1A, Changsha, China), followed by polishing of all teeth with a polishing paste. Root planing and periodontal pockets curettage under local anesthesia were done using Gracey curettes, followed by subgingival irrigation with a 10% povidone-iodine solution. Povidone-iodine has rapid anti-bacterial activity and resists deactivation by salivary or blood proteins. these are the properties that justify its use. After mechanical therapy, each individual received 0.12% chlorhexidine solution for daily mouthwash for 7 days, twice a day. Six weeks after the intervention, a complete periodontal reassessment was carried out with bacterial plaque control procedures, including scaling and polishing, and guidance on oral hygiene was repeated.

Statistical analysis
Statistical analysis was performed using Stata V.12.0 software. Continuous variables were expressed as median (IQR) and categorical variables as count (percentage). The Wilcoxon matched-pairs signed-rank test was used to compare variables before and 3 months after NSPT. A $p$ value $\leq 0.05$ was considered statistically significant.
RESULTS

Twenty-four subjects were enrolled in this study. A total of 20 subjects completed the study after excluding four patients during the 3 months of follow-up. The study flow diagram is shown in figure 1.

Subjects characteristics

Women represented 25% of all patients (5 of 20). The age of the subjects ranged from 18 to 59 years, with a median of 37 (29–45) years. BMI ranged from 18.9 to 32.9 kg/m², with a median BMI of 23.4 (21.35–26.57) kg/m². According to the 2012 CDC-AAP classification, 8 subjects had severe periodontitis, 11 had moderate periodontitis, and 1 subject had mild periodontitis.

Anthropometric parameters before and 3 months after NSPT

BMI, waist and hip circumference, waist to hip ratio, and fat mass were not statistically different before and 3 months after NSPT (table 1).

Periodontal parameters before and 3 months after treatment

Three months after NSPT, we observed a statistically significant reduction in all periodontal parameters after 6 and 12 weeks (PI, GBI, PD, and CAL; all p<0.001; table 1).

Table 1

<table>
<thead>
<tr>
<th>Variables</th>
<th>Before NSPT</th>
<th>3 months after NSPT</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Anthropometric parameters</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>23.4 (21.35–26.57)</td>
<td>23.43 (21.64–27.05)</td>
<td>0.579</td>
</tr>
<tr>
<td>Hip circumference (cm)</td>
<td>97 (92–100)</td>
<td>98 (94–110)</td>
<td>0.321</td>
</tr>
<tr>
<td>Waist circumference (cm)</td>
<td>80 (76–94)</td>
<td>82 (77–94)</td>
<td>0.139</td>
</tr>
<tr>
<td>Waist to hip ratio</td>
<td>0.84 (0.79–0.87)</td>
<td>0.83 (0.79–0.87)</td>
<td>0.262</td>
</tr>
<tr>
<td>Fat mass percentage (%)</td>
<td>14.75 (10.77–28.12)</td>
<td>17.9 (12.12–27.32)</td>
<td>0.118</td>
</tr>
<tr>
<td><strong>Periodontal parameters</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Plaque index (%)</td>
<td>91.07 (67.05–100)</td>
<td>16.96 (6.25–23.21)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Gingival bleeding index (%)</td>
<td>62.8 (40.44–78.1)</td>
<td>1.08 (0.00–4.50)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Pocket depth (mm)</td>
<td>3.12 (2.98–3.45)</td>
<td>1.6 (1.52–1.81)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Attachment loss (mm)</td>
<td>3.35 (3.08–4.0)</td>
<td>1.7 (1.57–2.05)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td><strong>Metabolic and inflammatory parameters</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>KITT (%/min)</td>
<td>3.72 (2.99–4.17)</td>
<td>4.04 (3.67–4.78)</td>
<td>0.001</td>
</tr>
<tr>
<td>FBG (mg/dL)</td>
<td>88 (83–94)</td>
<td>85 (79–89)</td>
<td>0.023</td>
</tr>
<tr>
<td>2-hour post-OGTT glycemia (mg/dL)</td>
<td>116 (107–127)</td>
<td>112 (99–126)</td>
<td>0.050</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>2.35 (1.46–4.18)</td>
<td>1.53 (1.03–2.12)</td>
<td>0.033</td>
</tr>
</tbody>
</table>

Data are median (IQR).
Significant p values are in bold.
BMI, body mass index; CRP, C reactive protein; FBG, fasting blood glucose; KITT, glucose disappearance rate; NSPT, non-surgical periodontal treatment; OGTT, oral glucose tolerance test.
DISCUSSION
In this pilot study, we investigated changes in metabolic, inflammatory, and periodontal parameters 3 months after NSPT with subgingival irrigation using povidone-iodine and mouth washing using chlorhexidine in individuals without diabetes with chronic periodontitis. The results showed that, 3 months after NSPT, there was a significant improvement in all studied periodontal parameters (PI, GBI, PD, and CAL), insulin sensitivity, and CRP level. Glucose tolerance improved by 7.44%, but did not reach statistical significance. No significant changes in body fat percentage and BMI were observed.

Accumulating evidence has shown that treatment of periodontitis is associated with subsequent changes in the level of serum inflammatory markers. Our findings showed that treatment of periodontitis might contribute to improvement of insulin resistance likely via a reduction in inflammation. This study shows a trend toward improvement in glucose tolerance at 3 months post-NSPT as assessed by OGTT-derived parameter(s). We measured insulin sensitivity by SITT, a simple, easy, quick, low-cost, reproducible, and reliable method to evaluate insulin sensitivity in clinical practice and epidemiological studies. SITT is safe and has been shown to closely correlate with the hyperinsulinemic euglycemic clamp (gold standard) for insulin sensitivity measurement. This shift in insulin sensitivity may be associated with a reduction in inflammation as shown by a decrease in CRP, although the median CRP value was less than 5 mg/mL at the beginning of the study. The small sample size, the lack of a reference group, and the heterogeneous profile of the subjects remain the main limitations of this study. Nonetheless, body fat percentage, BMI, and waist to hip circumference ratio, which have known effects on insulin resistance, did not significantly change. Since each subject was his/her own control, the effect on insulin resistance of these parameters has therefore remained constant. Many studies suggest that periodontal disease, diabetes, and pre-diabetes share a common denominator, which is chronic inflammation characterized by an increased expression of common proinflammatory cytokines. However, the mechanisms between inflammation and these diseases are not yet fully understood. Since chronic periodontitis is an inflammatory disease, further randomized controlled trials controlling for confounders are needed to confirm its effect on blood glucose at the metabolic level.

CONCLUSION
To the best of our knowledge, this was the first study to evaluate the impact of non-surgical periodontal therapy on insulin resistance and glucose tolerance. Our results indicate that treating chronic periodontitis with NSPT reduces systemic inflammation and improves insulin sensitivity despite a marginal effect on glucose tolerance assessed using 2-hour postload glucose during OGTT in individuals without diabetes. Further, randomized controlled trials are needed to demonstrate the impact of NSPT on glucose tolerance and confirm its effect on insulin resistance.

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Contributors ARNN, ARN, and ES designed the research. ARNN, ND, ET, and ES performed the research. ARNN, NL, NFTT, CBM, and ES contributed new reagents/analytic tools. ARNN, NL, ET, AT, JCK, and ES analyzed the data. ARNN, NL, NFTT, JCK, AT, and ES wrote the paper. ARNN, NL, NFTT, JCK, CBM, AT, EB, MD, and ES critically revised and adopted the manuscript. ES is the guarantor of this work. All authors approved the final version of the manuscript.

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Competing interests None declared.

Patient consent for publication Not required.

Ethics approval All procedures performed in studies involving human subjects were following the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards. The study received approval from the institutional and research ethical committee of the Faculty of Medicine and Biomedical Sciences and the institutional review board of the Yaoundé Central Hospital of Cameroon (no: AR/DHCY/CM/AD_V/L_26/01/2015).

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