Glycemic control and adipokines after periodontal therapy in patients with Type 2 diabetes and chronic periodontitis

Abstract: The mechanism by which chronic periodontitis (CP) affects type 2 diabetes (T2DM) remains unclear. Therefore, the aim of this study is to evaluate the effects of periodontal therapy (PT) on the glycemic control and adipokines of patients with T2DM and CP with the purpose of elucidating the possible mechanisms by which CP influences T2DM. Forty-four patients with T2DM and CP were randomly divided into two groups according to whether they underwent PT. Periodontal status, blood glucose, and the levels of serum tumor necrosis factor α (TNF-α), interleukin-6 (IL-6), adiponectin (APN), and fibroblast growth factor-21 (FGF-21) were measured at baseline and after 3 months. The results revealed that the probing depth (PD) and attachment loss (AL) were significantly improved, the serum levels of TNF-α and IL-6 were significantly decreased, and APN and FGF-21 exhibited substantial increases in the intervention group after 3 months (p < 0.05), whereas no significant changes were observed in the control group. The glycated hemoglobin (HbA1c) levels in both groups decreased significantly after 3 months compared with baseline (p < 0.05), but the intervention group exhibited a significantly greater change (p < 0.05). In conclusion, PT may relieve periodontal inflammation, which causes a reduction of insulin-antagonizing adipokines and an increase in insulin-sensitizing adipokines, thereby eliciting an improvement in glycemic control.

Keywords: Adipokines; Diabetes Mellitus; Periodontitis.

Introduction

Diabetes mellitus (DM) is a metabolic disorder that is mainly characterized by hyperglycemia due to impaired insulin action. The most common form of DM is type 2 diabetes mellitus (T2DM), which accounts for approximately 90% of all DM patients.1-2 Due to social mobility and rapid urbanization, the global prevalence of DM has increased sharply in recent years. Globally, 415 million adults have diabetes, and by 2040, this number will rise to 642 million.2 China now has the largest epidemic worldwide, and a recent study demonstrated that the prevalence of DM is up to 11.6% (95%CI: 11.3% –11.8%) among Chinese adults.3

Chronic periodontitis (CP) is one of the most common chronic infectious diseases and is characterized by destruction of the supporting structures of the teeth. According to the estimated data by Paul IE, 46% of adults in...
the United States have periodontitis, and 8.9% have severe periodontitis. In China, periodontal diseases have become one of the major public health problems. The third national epidemiological investigation on oral diseases in Chinese people demonstrated that gingival bleeding and periodontal pocket depths (PDs) > 4 mm are present in most middle-aged people (77.3% and 41.0%, respectively) and many elderly people (68.0% and 52.3%, respectively). Evidence indicates a “two-way” relationship between T2DM and CP. A large number of epidemiological studies have demonstrated that T2DM may increase the risk of CP by two- to three-fold. Conversely, CP may be a risk factor for worsened glycemic control in diabetic individuals and may increase the risk of related complications. Currently, the mechanism by which T2DM influences CP is relatively clear, but the direct effect of CP on T2DM is not yet well understood. In recent years, the roles of adipokines in inflammatory responses and insulin resistance (IR) have attracted much attention from researchers. Many studies have found that adipose tissue is not only a passive energy reservoir but is also an active endocrine organ that can produce and release diverse secretory proteins called adipokines into the systemic circulation. Currently, increasing numbers of adipokines are being found to influence inflammatory responses and IR. According to their effects on IR, adipokines can be classified into two categories. The first is insulin-sensitizing adipokines, which can improve insulin sensitivity, including visfatin, adiponectin (APN), fibroblast growth factor-21 (FGF-21), etc. The second is insulin-antagonizing adipokines, which can promote insulin resistance including tumor necrosis factor α (TNF-α), interleukin-6 (IL-6), resistin, etc.

Studies suggest CP can lead to an elevated inflammatory state and affect the levels of serum adipokines and inflammatory mediators. In this study, we investigated the effects of periodontal therapy (PT) on periodontal status, glycemic control, and adipokines in patients with T2DM and CP. Our hypothesis is that CP increases the levels of systemic inflammatory factors, which can affect serum adipokine levels and act to aggravate insulin resistance and worsen glycemic control. Alternatively, PT and the removal of the cause of this local chronic inflammation may positively influence inflammatory responses and IR by decreasing the levels of insulin-sensitizing adipokines and the increasing the levels of insulin-sensitizing adipokines, which would then improve glycemic control in these patients. Therefore, the aim of this study is to evaluate the effects of PT on the adipokines and glycemic control of patients with T2DM and CP with the purpose of exploring the role of adipokines in the possible mechanism of the effect of CP on T2DM.

**Methodology**

**Subjects**

The process of subject selection and inclusion is illustrated in Figure 1. A total of 155 T2DM patients with stable statuses who were followed up at Xiamen Lianqian Community Health Center after the diagnoses of diabetes at a grade three hospital between June 2014 and December 2014 were screened via a questionnaire investigation and clinical periodontal examination. Forty-four subjects who fulfilled the following inclusion criteria were selected for further study: 1) patients with a diagnosis of T2DM for over one year by a grade three hospital who volunteered for the research; 2) 6.50 ≤ Hb Alc % ≤ 10.00; 3) chronic periodontitis, ≥15 remaining teeth, and more than 30% of teeth with probing depths (PDs) ≥ 5 mm and attachment loss (AL) > 4 mm, or more than 60% of teeth with a PD > 4 mm and an AL ≥ 3 mm; 4) a body mass index (BMI) < 30 kg/m²; 5) without periodontal treatment in the previous 6 months; 6) without antibiotic or non-steroidal anti-inflammatory drug administration in the 3 months; and 7) without serious systemic diseases or complications.

A group of 44 random numbers was generated using SPSS ver. 17.0 (SPSS, Inc., Chicago, USA). These numbers were used to recruit and blindly randomize 44 subjects to either a control group (n = 22 subjects who underwent no intervention) or an intervention group (n = 22 subject who underwent a periodontal intervention) with a 1:1 allocation ratio. Ultimately, a total of 39 of the 44 recruited patients finished the study. Three patients in the intervention group were lost to follow-up because they could not revisit on
time. One patient in the control group performed the periodontal treatment in the follow-up period, and one patient migrated to another place.

Informed consent was obtained from all patients. This study was approved by the Ethics Committee of Xiamen Dentistry Hospital. To isolate the effects of the periodontal intervention on glycemic control and adipokines, no changes in medications or diet were made for any subject during the study period.

Examiner training and calibration

To avoid the bias due to different examiners, a single examiner (Jingsong Liu) was involved in the study. The examiner is an attending periodontist and has been treating patients for 8 years in Xiamen Lianqian Community Health Center. Before participating in the study, he had been trained for one month in the Department of Periodontology of Xiamen Dentistry Hospital. To validate the reliability of the periodontal examinations, an intra-examiner reliability assessment was executed, and good agreement (complete agreement rate > 70% and less than 1 mm error >80% in all parameters) was required.

Clinical periodontal examination

The PDs and ALs of all of the remaining teeth were tested at baseline and after 3 months. PD and AL were examined at six sites (mesiobuccal, buccal, distobuccal, distolingual, lingual, and mesiolingual) per tooth, and the average was calculated. A Williams periodontal probe was used for the clinical periodontal measurements, and all measurements were performed by a single examiner (Jingsong Liu) who was masked to the systemic status of each patient.

Periodontal intervention

All subjects in the intervention group underwent a periodontal intervention that included oral hygiene (utilizing correct methods and soft-bristled toothbrushes, interdental brushes and dental floss), full-mouth scaling (supragingival and subgingival scaling), the extraction of hopeless teeth, and the restoration of balanced occlusion. All periodontal interventions were completed by a single periodontist (Jingsong Liu) within two weeks.
Biological measurements

Blood samples were collected after overnight fasting, and the sera were obtained by centrifugation at 3,000 revolutions/minute for 10 minutes. The separated serum samples were collected in Eppendorf tubes and stored at -80°C until further testing. The levels of TNF-α, IL-6, APN, and FGF21 (test kits from Shanghai Tong Wei Industrial Co., Ltd., China) were tested using enzyme-linked immunosorbent assays (ELISAs) according to standard protocols at baseline and after 3 months. The HbA1c levels were tested with a D-10™ Hemoglobin Analyzer (Bio-Rad Laboratories, Inc., USA).

Statistical analysis

The statistical analyses were performed with SPSS ver. 17.0. The baseline balance between the groups was determined using independent-samples t-tests or the Pearson χ² test. One-sample Kolmogorov-Smirnov tests were used to test the distributions of the data for the periodontal parameters, HbA1c, and adipokines. Because the data for the PD, AL, HbA1c, TNF-α, APN and FGF21 were normally distributed, the differences are presented with bar graphs. The inner-group differences (i.e., baseline vs. after 3 months) were analyzed using paired-sample t-tests, and the intergroup differences were determined using independent-sample t-tests. Because the IL-6 data were not normally distributed, the difference is illustrated with a box plot, and the inner-group differences were analyzed with a non-parametric method (Wilcoxon signed-rank test).

Results

The demographic variables are presented in Table 1. The study groups were similar in age, gender, degree of education, BMI, the duration of diabetes, smoking, alcohol drinking, and physical exercise (p > 0.05). The levels of HbA1c, PD, and AL were similar in the study groups at baseline (p > 0.05).

As presented in Table 2, compared with baseline, the levels of PD and AL were significantly improved in the intervention group after 3 months (all p < 0.001). The mean changes in the PD and AL were statistically significant in the intervention group after 3 months compared with those in the control group (all p < 0.001).

At baseline, the HbA1c level exhibited no significant difference between the intervention group and the control group (p > 0.05). After 3 months, a greater reduction in the HbA1c level was observed in the intervention group, i.e., from 7.63 ± 0.89 to 6.99 ± 0.75 (p < 0.001), whereas a reduction of 7.67 ± 1.32 to 7.41 ± 1.31 was observed in the control group (p < 0.05). When the changes over time were compared between the two groups, the intervention group exhibited a significantly greater change in the HbA1c level than the control group (p < 0.05; Table 2).

As illustrated in Figure 2, the concentrations of serum TNF-α and IL-6 significantly decreased after periodontal therapy in the intervention group (p < 0.01, p<0.05), whereas these concentrations in control group at baseline and at 3 months were not significantly different (both p > 0.05; Figure 2A and Figure 2B). The APN and FGF21 levels increased significantly in the intervention group after 3 months of the periodontal treatment (both p < 0.01) but only changed slightly in the control group (both p > 0.05; Figure 2C and Figure 2D).

Discussion

Although the association between T2DM and CP has been extensively discussed, the mechanism by which CP affects T2DM remains unclear. In the present study, we evaluated the effects of PT on the levels of adipokines and HbA1c in patients with T2DM and CP with the purpose of examining the possible mechanisms by which CP influences T2DM. Four adipokines were selected for study, including two insulin-antagonizing adipokines (TNF-α and IL-6), and two insulin-sensitizing adipokines (APN and FGF21).

The etiology of periodontitis is closely related to dental plaque and calculus deposits in the periodontal tissues. Both healthy individuals and diabetic patients exhibit similar outcomes after periodontal therapy in terms of PD and AL reductions.18,19,20 In the present study, we also found that the levels of PD and AL were improved significantly after PT.
Table 1. Demographic characteristics of the study groups at baseline.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Intervention group (n = 19)</th>
<th>Control group (n = 20)</th>
<th>t/χ²</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td>61.58 ± 4.69</td>
<td>61.90 ± 6.75</td>
<td>0.172</td>
<td>0.865</td>
</tr>
<tr>
<td>BMI</td>
<td>24.32 ± 2.70</td>
<td>23.72 ± 3.46</td>
<td>0.606</td>
<td>0.548</td>
</tr>
<tr>
<td>Duration of DM (y)</td>
<td>8.47 ± 3.08</td>
<td>7.70 ± 4.69</td>
<td>1.243</td>
<td>0.214</td>
</tr>
<tr>
<td>HbA1c (%)</td>
<td>7.63 ± 0.89</td>
<td>7.67 ± 1.32</td>
<td>0.119</td>
<td>0.906</td>
</tr>
<tr>
<td>PD (mm)</td>
<td>3.66 ± 0.60</td>
<td>3.85 ± 0.58</td>
<td>0.988</td>
<td>0.329</td>
</tr>
<tr>
<td>AL (mm)</td>
<td>4.12 ± 0.89</td>
<td>4.28 ± 0.77</td>
<td>0.568</td>
<td>0.574</td>
</tr>
<tr>
<td>Gender (n [%])</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>12 (63.16)</td>
<td>14 (70.00)</td>
<td>0.205</td>
<td>0.741</td>
</tr>
<tr>
<td>Female</td>
<td>7 (36.84)</td>
<td>6 (30.00)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Education</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤ Primary school</td>
<td>6 (26.32)</td>
<td>3 (15.00)</td>
<td>1.703</td>
<td>0.427</td>
</tr>
<tr>
<td>Middle school</td>
<td>9 (47.33)</td>
<td>13 (65.00)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>≥ College/university</td>
<td>4 (21.05)</td>
<td>4 (20.00)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Smoking* (n [%])</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>6 (31.58)</td>
<td>3 (15.00)</td>
<td>1.509</td>
<td>0.219</td>
</tr>
<tr>
<td>No</td>
<td>13 (68.42)</td>
<td>17 (75.00)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Drinking** (n [%])</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>12 (61.15)</td>
<td>17 (75.00)</td>
<td>2.838</td>
<td>0.242</td>
</tr>
<tr>
<td>Seldom</td>
<td>3 (15.79)</td>
<td>2 (10.00)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Often</td>
<td>4 (21.06)</td>
<td>1 (5.00)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Physical exercise*** (n [%])</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Seldom</td>
<td>15 (88.95)</td>
<td>14 (70.00)</td>
<td>0.409</td>
<td>0.522</td>
</tr>
<tr>
<td>Often</td>
<td>4 (21.05)</td>
<td>6 (30.00)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Hypertension**** (n [%])</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>4 (21.05)</td>
<td>8 (40.00)</td>
<td>1.642</td>
<td>0.200</td>
</tr>
<tr>
<td>No</td>
<td>15 (78.95)</td>
<td>12 (60.00)</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

*: more than 1 cigarettes/day and continuing for more than 1 year; **: "No": never drinking; "Seldom": drinking ≤1 time/week; "Often": drinking>1 time/week; ***: "Often"= regular exercise ≥2 times/week, each time more than 30 minutes; ****: diagnosis of hypertension at a grade three hospital.

Table 2. Comparisons of the PD, AL and HbA1c levels in the two groups after three months.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Control group (n = 20)</th>
<th>Intervention group (n = 19)</th>
<th>Baseline</th>
<th>After 3 mos</th>
<th>Δ</th>
<th>Baseline</th>
<th>After 3 mos</th>
<th>Δ</th>
</tr>
</thead>
<tbody>
<tr>
<td>PD</td>
<td>3.85 ± 0.58</td>
<td>3.92 ± 0.56</td>
<td>0.07 ± 0.28</td>
<td>3.66 ± 0.60</td>
<td>-0.57 ± 0.30**</td>
<td>3.09 ± 0.63b</td>
<td>-0.57 ± 0.30**</td>
<td></td>
</tr>
<tr>
<td>AL</td>
<td>4.28 ± 0.77</td>
<td>4.36 ± 0.81</td>
<td>0.08 ± 0.32</td>
<td>4.12 ± 0.90</td>
<td>-0.50 ± 0.37**</td>
<td>3.62 ± 0.84b</td>
<td>-0.50 ± 0.37**</td>
<td></td>
</tr>
<tr>
<td>HbA1c</td>
<td>7.67 ± 1.33</td>
<td>7.41 ± 1.32</td>
<td>-0.27 ± 0.43</td>
<td>7.63 ± 0.89</td>
<td>-0.64 ± 0.66*</td>
<td>6.99 ± 0.77b</td>
<td>-0.64 ± 0.66*</td>
<td></td>
</tr>
</tbody>
</table>

The results are given as the mean ± the SD. Intragroup differences were analyzed using paired-sample t-tests. *: p<0.05. **: p<0.001. Intergroup differences were determined using the independent-samples t-test. *: p < 0.05. **: p < 0.001.
Therefore, PT, specifically the removal of dental accretions, can relieve the level of periodontal inflammation and improve periodontal conditions. The HbA1c level reflects the glycemic level over the previous 1 to 3 months. Whether periodontal therapy reduces the HbA1c level in periodontitis patients remains controversial. Very recently, Teshome T performed a meta-analysis, and the results revealed a statistically significant HbA1c reduction of 0.48 (95%CI: 0.18, 0.78) in the treatment group compared with the control group, and similar results have been obtained in other studies conducted by Li and Engebretson. In contrast, a multicenter, randomized clinical trial reported that, at 6 months, the mean HbA1c level in the periodontal therapy group increased 0.17% compared with 0.11% in the control group with no significant difference between the groups. Several outcomes of meta-analyses also do not support the notion that PT lowers the level of HbA1c. In the present study, the intervention group exhibited a significantly greater change in the HbA1c level, which indicated that PT may improve glycemic control.

It is believed that pro-inflammatory mediators, such as TNF-α and IL-6, are expressed by the inflamed periodontal sites due to microbial stimuli or a host response. These mediators enter the systemic circulation, interfere with the function of insulin receptors and thereby derange the process of insulin signaling. TNF-α was the first inflammatory marker that was thought to play a role in the development of obesity-induced insulin resistance in the 1990s. TNF-α causes impaired insulin-derived peripheral uptake of glucose by increasing the serine phosphorylation of insulin receptor substrate 1 (IRS-1) and consequently inhibits the translocation of glucose transporter type 4 (GLUT4) to the plasma.

![Graphs showing changes in TNF-α, IL-6, APN, and FGF21 concentrations](image-url)

*Figure 2. Serum TNF-α, IL-6, APN, and FGF21 concentrations in the two groups at baseline and after 3 months.*
membrane, which results in peripheral IR. Some studies have suggested that treatment with anti-TNF-α antibody can improve insulin sensitivity in IR patients. IL-6 is another inflammatory adipokine that is involved in the pathogenesis of IR. Agarwal A compared 40 patients with impaired fasting glucose (IFG) with 40 healthy people, and IL-6 was found to be significantly elevated in the IFG group and to significantly correlate with IR. A prospective cohort study conducted by Bertoni found that serum IL-6 levels are significantly associated with T2DM. IL-6 induces IR through the prevention of non-oxidative glucose metabolism and the suppression of lipoprotein lipase, which increases the triglyceride level.

In the present study, significant reductions in serum TNF-α and IL-6 after 3 months of therapy were observed. These findings agree with those of other studies. Recently, the outcome of a meta-analysis of 6 randomized-controlled trials and 3 controlled clinical trials revealed a statistically significant mean difference (MD) in TNF-α (-1.33 pg/ml, 95%CI: -2.10; -0.56) favoring the periodontal intervention versus the control. In a prospective blind intervention trial, the level of IL-6 exhibited a significant decrease. These results indicate that PT may relieve periodontal inflammatory conditions and further decrease pro-inflammatory, insulin-antagonizing adipokines.

APN is a protein that is secreted from adipose tissue and is an insulin-sensitizing, anti-inflammatory adipokine that has been demonstrated to improve glucose tolerance and insulin resistance in humans. Studies have demonstrated that adiponectin improves insulin sensitivity through the inhibition of TNF-α-induced adhesion molecule expression and the inhibition of NF-κB activation. Many studies have revealed that effective periodontal intervention can increase the serum APN levels in T2DM patients with periodontitis. FGF-21 was discovered to be a potent regulator of glucose uptake in mouse 3T3-L1 cells and primary human adipocytes. FGF-21 can improve insulin sensitivity and lipid metabolism in obese and diabetic animal models and has been proposed as a potential therapeutic agent for the treatment of T2DM, obesity and related complications. FGF-21 significantly improves insulin signaling by enhancing the phosphatidylinositol 3-kinase (PI3K)/AKT pathway, up-regulating glucose uptake, promoting the release of insulin-sensitizing adipokines, such as adiponectin, and reducing the release of insulin-antagonizing leptin. This study represents the first introduction of FGF21 into a periodontal intervention study that aimed to investigate the effect of PT on FGF21. We found that the serum APN and FGF21 levels were significantly increased after therapy, which indicated that the effective control of inflammation by periodontal treatment may contribute to increasing systemic insulin-sensitizing adipokines, such as APN and FGF21, thus improving the overall health status.

There are several limitations to this study. First, the subjects were elderly; thus, it remains uncertain whether the results of this study can be generalized to other diabetic populations. Second, this study only observed the effect after three months, and the long-term effects after intervention should be considered. Therefore, a larger and long-term epidemiological study should be conducted in the future.

Conclusion

In summary, PT relieved the periodontal inflammatory status, which in turn caused reductions in insulin-antagonizing adipokines and increases in insulin-sensitizing adipokines that were reflected by an improvement in glycemic control in T2DM patients with CP. Large-scale and long-term human epidemiological studies and animal model studies exploring the role of adipokines in the mechanisms by which CP influences T2DM are necessary.

Acknowledgments

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References


