Oxidative Stress and Periodontal Disease in Diabetic patients: A 3-month pilot study

MARCONCINI, S, et al.
Oxidative stress and periodontal disease in diabetic patients: a 3-month pilot study

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Abstract

Aim: Oxidative stress is involved in both Periodontal Disease (PD) and Diabetes Mellitus (DM). The present study aimed to evaluate the oxidative balance in diabetic patients diagnosed with PD before and after non-surgical periodontal treatment.

Materials and methods: Sixty patients were divided into three groups all receiving non-surgical periodontal treatment plus either chlorhexidine, ozone-therapy or antioxidant mouth-rinse. Probing depth (PPD), Percentage Plaque Index (%PI) and Percentage Bleeding on Probing (%BoP) were recorded. Free-radicals (dROMs), plasmatic antioxidants (PAT), salivary antioxidants (SAT) and glycated hemoglobin (HbA1c) were measured.

Results: Mean PPD at baseline was 3.14 mm. Mean PPD three months after was 2.05 mm. The overall %PI at baseline was 55% and %BoP 76%. Three months after treatment %PI was 34% and %BoP was 64%. The longitudinal analysis did not show differences between groups. Mean dROMs at baseline was 353 U. Carr (oxidative stress) and decreased after three months reaching 295 U. Carr (normal), SAT was 2083 U. Carr at baseline (inflammation) and decreased to 1337 U. Carr (ideal). The longitudinal analysis did not show differences between groups. Mean HbA1c at baseline was 6.92% and decreased significantly to 6.63% three months after treatment.

Conclusion: Based on the results of the present study, oxidative stress should be further investigated as a potential modulator of the clinical course of both DM and PD.

Introduction

Periodontal Disease is caused by a dysbiosis, an imbalance in the relative presence or influence of microbial species that take part of the oral microbiome [1,2]. The dysbiotic microbial community, if left untreated, might cause or, at least, exacerbate other systemic inflammatory disorders [3]. In the last decades, many researchers have focused on the association between periodontitis and various systemic inflammatory pathologies. An increasing number of studies have suggested that systemic inflammation might sometimes result from the entry of oral microbial agents and their virulence factors into the circulation [4].

Diabetes mellitus (DM) is increasingly common worldwide [5]. Diabetic subjects present an increased prevalence and severity of periodontitis [6,7]. The pathological microbial environment of periodontitis has been suggested as a metabolic influencer [8,9]. The nature of the relationship is still debated [10]. Several mechanisms have been proposed to explain it: increased oxidative stress, advanced glycation end-products, altered immune function, and changes in collagen. Nowadays, there is a broad consensus about the fact that periodontal treatment might improve glycaemic control: a significant reduction of HbA1c, 3 months after periodontal treatment, has been demonstrated [11-13]. Since periodontitis is greatly untreated, what measures would improve its management in persons at risk for complications of diabetes? Many inflammatory chemokines and endotoxins (c-reactive protein, interleukin 6, tumour necrosis factor α) have been investigated as markers of systemic inflammation in diabetic patients [14]. Lately, a large amount of literature has been directed to the study of reactive oxygen species and their role in the pathophysiology of different systemic inflammatory disorders [15]. Oxidative stress is caused by an excessive release of free radicals or by a deficiency of anti-oxidant agents. A great amount of pro-oxidant agents is produced during inflammatory conditions, such as chronic periodontal disease [16]. The overexposure to free radicals may worsen systemic inflammatory conditions leading to tissue damage [17]. Saliva is rich in molecules, such as uric acid, vitamin C, reduced glutathione (GSH), oxidized glutathione (GSSG), lacto-ferrine, and many others [18]. Since those molecules work in concert renewing each other, total antioxidant capacity may be useful to assess individuals’ defense capabilities. The aim of the present prospective clinical study was to investigate the putative correlation between oxidative stress, clinical periodontal indexes and glycaemic status in diabetic patients. The null hypothesis was that non-surgical periodontal treatment does not induce any remarkable difference in terms of markers of oxidative stress and glycaemic status.

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Key words: periodontal disease, oxidative stress, salivary antioxidant capacity

Received: May 01, 2017; Accepted: May 12, 2017; Published: May 15, 2017
Materials and methods

Sixty patients were recruited for this prospective randomized clinical study. All of the patients were diagnosed with Diabetes Mellitus (DM) type 1 or 2 at the Diabetology Department of the University of Pisa. Detailed information of their current medications were collected as well as their glycemic control status.

Each patient had to sign an informed written consent and to answer a specific anamnestic questionnaire.

The study has been conducted in accordance with the Helsinki declaration of 2002.

Selection criteria

Inclusion criteria for study participation were an age range of 18–70 years and the diagnosis of both Diabetes Mellitus (type 1 and type 2) and Periodontitis.

Exclusion criteria were

- Pregnancy, conditions requiring antibiotic prophylaxis, psychiatric disorders, severe physical handicaps, carcinoma, immunosuppressive therapy, hospital in-patient, vitamins supplements or antibiotics taken within 3 months before the study or periodontal therapy received within 6 months.

Study design

The study design was a prospective clinical study. The cohort size was determined by power calculation analysis according to previous published literature. Recruitment began in September 2015, and the follow-up is still on-going. The last recall visit is due to be within 12 months from baseline. All of the recruited patients underwent a periodontal examination (clinical assessment with a standard periodontal UCLA probe) and were eventually asked to participate in the experimental part of the study. Diagnosis of periodontal disease was assessed as the proximal attachment loss ≥3 mm in ≥2 non-adjacent, BOP on at least 25% of their total sites and documented radiographic bone loss [19].

At baseline (T₀) patients were randomly allocated to one of the three groups of treatment, so that a professional hygiene session was performed and an adjunctive domiciliary therapy was prescribed. At the same time, clinical and laboratory parameters were collected. The follow-up schedule comprised two recall visits: one month (T₁) and three months after periodontal treatment (T₃). Each visit consisted in a full periodontal examination and in the collection of saliva and blood samples in order to evaluate specific markers.

Periodontal assessment

One investigator performed full periodontal examinations. Periodontal probing depth (PPD) was measured using a UNC-PCP15 (Hu Friedy) probe at six sites (mesio-buccal, mid-buccal, disto-buccal, mesio-lingual and disto-lingual) per tooth (third molars excluded). Mean values of PPD per patient were calculated. Marginal bleeding on probing (BoP) and plaque index (PI) were assessed dichotomously (“1” present, “0” absent) at six sites per tooth. For each patient BoP and PI were then expressed as a percentage (BoP%, PI%) of positive sites.

Periodontal treatment and motivation

Randomization was obtained by a virtually-generated list. Diabetic patients were then assigned to one of the three groups of treatments (twenty patients per group):

- Control group: standard non-surgical periodontal treatment (ultrasonic debridement and manual scaling) plus a domiciliary prescription of a commercial chlorhexidine mouthwash for the following 7 days.
- Test group: standard non-surgical periodontal treatment (ultrasonic debridement and manual scaling) plus a domiciliary prescription of an anti-oxidant mouthwash for the following 7 days.
- Test group: standard non-surgical periodontal treatment (ultrasonic debridement and manual scaling) plus professional ozone administration and the domiciliary usage of an ozone delivery device (Aqualab s.r.l). The ultrasonic device used was the Mectron Multipizeo (Mectron s.p.a.). The ozone delivery device was a hydro-pulseur (Aqualab s.r.l) with a magnetic drive pump that produces a continuous release of ozonized-water. Professional usage was set at a higher range of ozone tension (12 V) with the smallest nozzle (0.6 mm). Domiciliary devices were calibrated and then handed to patients. Ozone tension was set at 12 V and the nozzle of choice was the largest available.

Motivation to proper oral hygiene had the duration of 30 to 60 minutes per patient; the operator informed each patient about the correct maneuvers to perform at home in order to achieve the best results in terms of personal oral hygiene.

Laboratory analysis

Clinical samples were collected early in the morning. Both saliva and blood samples were processed according to instructions furnished the company producing the collection kit and the dedicated spectrophotometer (H&D s.r.l).

The d-ROM (derived reactive oxygen molecules) test determines the concentration of hydro-peroxides in the blood. Its unit of measurement is the U. CARR (0.08 mg/dL of a solution of hydrogen peroxide). The PAT (plasmatic antioxidant test) determines the concentration of the water-soluble antioxidants in the blood that are able to reduce ferric ions to ferrous ions. Its unit of measurement is the U. CARR as well. The SAT (salivary antioxidant test) evaluates the salivary total antioxidant capacity. Saliva was immediately analyzed since it degenerates fast altering the absorbance properties of the sample.

Glycemic status and medical anamnesis

At baseline, information regarding physical activity, diet, major complications of diabetes, body mass index and smoking habits were recorded. In cooperation with the Department of Diabetology, the level of glycated hemoglobin was assessed before treatment. At three months, the new value/ post-treatment was registered.

Statistical analysis

Clinical and laboratory parameters were inserted into an Excel data-sheet. Outcome variables were defined as: mean probing depth (PPD), mean plaque index in percentage (PI%), mean bleeding on probing in percentage (BoP%), mean dROMs, mean PAT, mean SAT, mean HbA₁c. Descriptive and statistical analyses of data were performed by usage of two statistical tool packages (Stata® and R Studio®). Normal distribution was tested with the Shapiro-Wilkoxon method. All measurements in the text and Tables are described as mean and standard deviations (m ± std). Intergroup and intragroup longitudinal analysis was performed using the Brunner and Langer model for non-parametric longitudinal designs [20]. The inter-group analysis (between subject) was based on the stratification for the group...
of treatment (whole-plot factor). The intra-group analysis (within subjects) relied on the stratification for repeated measures on one sub-plot factor: time. The mixed method applied for longitudinal analysis consisted of Wald type statistic (WTS), ANOVA type statistic (ATS) and modified ANOVA-Type Statistic for the Whole-Plot Factors. The significance was set at 95% for all of the tests performed.

Results

A total of 75 Type I and 2 diabetes patients were screened; 11 patients were excluded based on their medical history and 4 for personal reasons, leaving a total of 60 patients eligible for study participation. Table 1 reports the demographic characteristics of the initial cohort. The cohort presented 22 females and 38 males; mean age was 60.9 years old with Type-I patients, being significantly younger than type II. Mean body mass index was 27.8, being higher among type 2 Diabetes patients. Eleven patients were smokers. Twenty-two out of 60 patients suffered from DM type 1 and the left 38 patients were affected by DM type 2.

Table 2 reports the periodontal findings for each group of treatment, for every moment of the follow-up. Mean probing depth (PPD) for the entire cohort at baseline was 3.14 mm. The overall mean plaque index at baseline was 55%. Bleeding on Probing (BOP) resulted positive in 76% of sites. Three months after treatment, the global mean of PPD was 2.05 mm. The mean percentage plaque presence decreased to 34% of positive sites. Bleeding on Probing showed a positive percentage at 3 months of 64% of sites analyzed. The oxidative balance descriptive statistics is reported in table 3. The mean amount of plasmatic free radicals (d-ROM) for the cohort at baseline was 353 U.Carr which is considered to be a normal range. The mean concentration of Plasmatic Antioxidant agents at baseline was 1838 U.Carr. which is considered to be a deficiency status. Coming to the analysis of Total Antioxidant Capacity of Saliva at baseline the mean value was 295 U.Carr which is considered ideal. The mean value of HbA1c at baseline was 6.92% which is considered a deficiency status. Three months after periodontal treatment (Table 4). Anyway, it was not possible to demonstrate a treatment effect in reducing glycated hemoglobin neither a different outcome between type I and type II diabetic patients.

Discussion

The present study was a randomized clinical investigation and it was based on emerging data that relate oxidative stress to both Diabetes and Periodontal disease [21]. The results of the present study.
agreed with the current literature about the capability of non-surgical periodontal treatment to reduce systemic inflammation [22]. It has been demonstrated that chronic inflammation following periodontal infection may enhance the release of inflammation mediators and free radicals, thus threatening the natural history of diabetes [23]. In gingival tissues of subjects who also have periodontitis, the presence of AGEs (advanced glycation end-products) and accompanying markers for increased oxidant stress have been demonstrated [24]. The present cohort of patients was characterized by its fair glycaemic control (mean value at baseline of 6.92%) due to the fact that all the patients joined a strict regimen within the unit of Diabetology of the University of Pisa. The mean probing depth at baseline was 3.14 mm and the mean bleeding on probing was 76%. The mean plaque index was 1.94. These values pictured a mild-to-moderate periodontitis at baseline for the entire cohort. The descriptive statistics for the mean initial oxidative balance showed a tendency towards pathological ranges. The mean count of plasmatic free radicals (d-ROM) was 353 U.Carr., a value that denoted a moderate level of oxidative stress. At the same time, the plasmatic antioxidant status (PAT) showed a deficiency if compared to the general population. The amount of plasmatic reactive oxygen species (dROMs) showed a significant reduction three months after periodontal treatment in each of the three groups of treatment. The intergroup analysis did not highlight any remarkable differences in terms of dROMs reduction between the three treatment modalities. PAT values remained in a deficiency status at a three-month evaluation. The mean total salivary antioxidant capacity of the present cohort at baseline was 2083 U.Carr which resulted higher than normal range thus denoting a potential state of inflammation in diabetic patients affected by moderate periodontitis. The over-production of free radicals caused by periodontal inflammation is in fact supposed to cause an up-regulation of the anti-oxidant defence [25,26]. The results of the present study agree with literature relating high levels of salivary antioxidants (SAT) to a possible periodontal inflammatory condition. In response to oxidative stress, antioxidant enzymes appeared up-regulated in inflamed periodontal tissues [27,28]. The longitudinal analysis of this trial showed a normalization of SAT values three months after non-surgical periodontal treatment. This inversion of tendency was coherent to the reduction of mean PPD, PI% and BOP% of the cohort during the follow-up. Since recently it has been hypothesized that oral dysbiosis might contribute to diabetes genesis among healthy individual [29], it would be crucial to explore new ways to monitor the oral microbiome behavior. The present clinical investigation has linked diabetes condition cross-sectionally to oxidative stress both systemic (plasmatic count) and local (salivary count). Periodontal treatment brought to periodontal indices normalization and to a reduction in terms of oxidative stress. Periodontal indexes showed a better outcome 1 month after treatment if compared to the three-month evaluation. This result highlighted how it is difficult to maintain an ideal balance within few months in problematic patients. In our opinion, it would be crucial to tailor the timing of hygiene recalls to each patient considering one’s specific “periodontal deadline”. Coming to glycaemic control, the mean reduction of glycated haemoglobin was 0.56. It is a key point to note that baseline values of HbA1c among this cohort were already almost acceptable for a group of diabetic patients. Therefore, a small reduction of HbA1c over three months, should not be ignored even if occurring within non-alarming values. At least, it would be worthwhile to investigate the same phenomenon over a larger sample including uncontrolled diabetic patients. The mean reduction of HbA1c was more evident among the ozone test group thus reinforcing the role of this particular type of adjunctive periodontal therapy in diabetic patients. The present study was designed as a pilot research in the field of perio-medicine. Given its limitations in terms of the number of patients and heterogeneity of the sample, it was impossible to establish any linear relationship between the variables of interest. The results obtained suggested the possibility to identify new markers describing the risk profile of diabetic patients to develop inflammatory-based-complications, periodontitis being included. Oxidative stress is an active factor both in the etiology and in the course of diabetes and periodontal disease. To monitor the oxidative balance of a complicated patient could mean preventing future major complications. It will be interesting in the future to further investigate the role of ozone therapy in the management of periodontal disease in diabetic patients.

### Disclosure statement

The paper has been submitted solely to Journal of Dental, Oral and Craniofacial Research and that it is not concurrently under consideration for publication in another journal.

All individuals listed as authors agree that they have met the criteria of authorship and agree to the conclusions of the study.

### Conflicts of interest

The authors report no conflicts of interest related to this study.

### Footnotes

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+ Mectron s.p.a. via Loreto 15° 16042 Carasco (GE) Italy
+ H&D s.r.l., Strada Langhirano n°264/1°, Località Fontanini 43124 Parma (PR), Italy
+ Stata, StataCorp LP 4905 Lakeway Drive College Station, Texas 77845-4512 USA
+ R Studio free software, 250 Northern Ave, Boston, MA 02210

### Authors contribution

Marconcini S: contributed to conception and design; contributed to acquisition, analysis, and interpretation; drafted manuscript; critically revised manuscript; gave final approval; agrees to be accountable for all aspects of work ensuring integrity and accuracy

Barone A: contributed to conception and design; contributed to analysis and interpretation; drafted manuscript; critically revised manuscript; gave final approval; agrees to be accountable for all aspects of work ensuring integrity and accuracy

Covani U: contributed to conception and design; contributed to analysis and interpretation; drafted manuscript; critically revised manuscript; gave final approval; agrees to be accountable for all aspects of work ensuring integrity and accuracy

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Table 4. Longitudinal analysis for non-parametric distributions.
Genovesi AM: contributed to conception and design; contributed to acquisition and interpretation; drafted manuscript; critically revised manuscript; gave final approval; agrees to be accountable for all aspects of work ensuring integrity and accuracy.

Soder B: contributed to conception and design; contributed to interpretation; drafted manuscript; critically revised manuscript; gave final approval; agrees to be accountable for all aspects of work ensuring integrity and accuracy.

Giammarinaro E: contributed to conception and design; contributed to acquisition, analysis, and interpretation; drafted manuscript; critically revised manuscript; gave final approval; agrees to be accountable for all aspects of work ensuring integrity and accuracy.

Giampietro O: contributed to conception and design; contributed to acquisition, analysis, and interpretation; drafted manuscript; critically revised manuscript; gave final approval; agrees to be accountable for all aspects of work ensuring integrity and accuracy.

Giampietro C: contributed to conception and design; contributed to acquisition and analysis; drafted manuscript; critically revised manuscript; gave final approval; agrees to be accountable for all aspects of work ensuring integrity and accuracy.

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