Effect of nonsurgical periodontal therapy and smoking status on hematological variables related to anemia of chronic disease in chronic periodontitis patient: a case-control study

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Background: Chronic infectious, inflammatory, or neoplastic disorders are associated with anemia of chronic disease. Chronic inflammatory diseases such as periodontitis may contribute to masked anemia, especially in smokers. This study was aimed at verifying and comparing the efficacy of nonsurgical periodontal therapy (NSPT) for improving anemia among chronic periodontitis patients with and without the habit of smoking.

Methods: Thirty systemically healthy individuals with chronic periodontitis were divided into two groups of 15 each, smokers (group A) and nonsmokers (group B). The groups were compared based on hematological parameters such as serum erythropoietin (SE) and serum ferritin (SF) levels at baseline and 3 months after NSPT for anemia evaluation.

Results: The baseline SE levels in groups A and B were 11.84 and 15.19 mIU/mL (p = 0.031), respectively; the corresponding levels at 3 months after NSPT were 13.00 and 17.74 mIU/mL (p = 0.022). The baseline SF levels in groups A and B were 95.49 and 44.86 ng/mL (p = 0.018), respectively; the corresponding levels at 3 months after NSPT were 77.06 and 39.05 ng/mL (p = 0.009). Group B showed a significant increase and decrease in the SE and SF levels, respectively, at 3 months after NSPT (p = 0.035 and p = 0.039, respectively), whereas group A showed insignificant changes (p = 0.253 and p = 0.618, respectively).

Conclusion: NSPT led to an improvement in anemia among chronic periodontitis patients. However, the improvement is less in smokers compared to that in nonsmokers. Furthermore, SF and SE levels might serve as effective biomarkers for assessing anemia in smokers and nonsmokers with chronic periodontitis.

Keywords: Anemia; Chronic periodontitis; Erythropoietin; Ferritin; Smoking
Introduction

Chronic infectious, inflammatory, or neoplastic disorders are associated with anemia of chronic disease (ACD) [1]. Pathophysiological factors such as limited iron availability to erythroid progenitor cells, a blunted response to erythropoietin, erythropagocytosis, diminished erythropoiesis, and microbial/tumor cell infiltration of bone marrow contribute to the development of ACDs. Patients with ACD demonstrate diminished levels of serum iron, normal to elevated serum ferritin (SF) levels, and normocytic to microcytic anemia [2]. Periodontitis is a chronic inflammatory disease characterized by increased production of inflammatory cytokines that can contribute to the prevalence of anemia by directly inhibiting erythropoiesis and inducing changes in iron absorption and release [3].

Patients with chronic periodontitis (CP) who are regular smokers show lower gingival redness and bleeding on probing due to potential vasoconstriction caused by the nicotine content in tobacco. This may lead to an inaccurate assessment of periodontal status and failure to diagnose the underlying pathogenic state [4]. In addition, smoking causes an increase in hemoglobin (Hb) concentration mediated by carbon monoxide, which bonds with Hb and forms inactive carboxyhemoglobin (COHb) with a reduced ability to deliver oxygen to the tissues. Therefore, as a compensatory mechanism, smokers maintain a higher Hb level than nonsmokers, which is referred to as secondary polycythemia [5,6]. Despite the higher Hb levels found in smokers, this hypoxic state triggers erythropoietin production, thereby increasing erythropoiesis. Erythropoietin is a large glycoprotein hormone produced by the peritubular cells lining the kidneys and hepatocytes; it is the principal regulator of the erythrocyte lineage [7]. This clearly implies that the underlying anemic state in smokers is masked by high Hb values, which may lead to an underestimate of the prevalence of anemia among smokers [8]. In such a scenario, an estimation of serum erythropoietin (SE) levels may aid in assessing the anemia status of smokers.

In chronic inflammatory conditions such as CP, proinflammatory cytokines such as interleukin (IL)-1α, IL-1β, IL-6, tumor necrosis factor-α, and transforming growth factor-β not only increase hemolysis and impair erythropoiesis via direct bone marrow suppressive effects but also release reactive oxygen species that inhibit erythropoietin gene expression [9]. Singh et al. [10] and Hutter et al. [11] found lower erythropoietin levels in patients with CP, thereby strengthening the hypothesis that CP may lead to ACD.

Acute-phase proteins are biomarkers that show changes in plasma concentration that increase (positive acute-phase proteins), such as ferritin, or decrease (negative acute-phase proteins) by at least 25% during inflammatory disorders, due to their altered production by hepatocytes [12]. Ferritin serves as the main iron storage protein in the body and contains 20% iron by weight. The serum iron concentration is directly proportional to its storage in the body, which increases under inflammatory conditions and iron overload. Chakraborty et al. [13] observed that SF levels were higher in patients with CP than in healthy controls, and a reduction was noted with remission of chronic inflammation following nonsurgical periodontal therapy (NSPT). Considering the prevalence of periodontal disease in the community at large, its deleterious effects on the systemic health of affected individuals, the increased use of tobacco in the form of cigarette smoking, and the resultant masking of anemia status, therapeutic measures such as scaling and root planning (both NSPTs) might significantly improve anemia status in such individuals.

The available literature suggests that NSPTs were not tested while evaluating erythropoietin levels in smokers and nonsmokers with CP. To date, only a few studies have evaluated NSPTs with a single hematological parameter to assess ACD in CP cases. Hence, multiple hematological parameters, such as SF and SE levels, collectively might serve as a better means to assess the anemia status of smokers with CP instead of any single parameter. The aim of the present study was to verify and compare the efficacy of NSPT in improving anemia status among patients with CP who do or do not smoke.

Methods

Ethical statements: This study was carried out following CONSORT (Consolidated Standards of Reporting Trials) guidelines and written informed consent was obtained from all participants who fulfilled the inclusion criteria and agreed to participate voluntarily. Details about the nature, risk, and benefits of the hematological investigations as well as the associated procedures were explained to all participants. The experimental protocol and consent form were approved by the Institutional Ethical Committee and Institutional Review Board (IRB) of Dr. R. Ahmed Dental College and Hospital (IRB No: DCH/07/18-19).

1. Study design

This clinico-biochemical study included 30 systemically healthy patients diagnosed with CP stage I/II (probing pocket depth of ≥ 4 mm but < 6 mm) requiring NSPT who were selected from the outpatient Department of Periodontics [14]. CP was confirmed clinically and radiographically according to the guidelines of the
2. Statistical analysis

The Shapiro-Wilk test was performed to assess the assumption of normality of the data. Data are presented as mean ± standard error of the mean (SEM). An unpaired Student t-test was performed to compare the parameters of the two groups that showed normal distributions. Normally distributed paired data of each group were compared using a paired Student t-test. Non-normally distributed unpaired data were evaluated using the Mann-Whitney U-test, and paired data were evaluated using the Wilcoxon matched-pairs signed-rank test. Correlations between two normally distributed parameters were evaluated using the Pearson correlation test. Sex distribution between the two groups was evaluated using Fisher exact test. The correlation between two non-normally distributed parameters was evaluated using Spearman nonparametric correlation. Direct and inverse correlations were indicated by positive and negative correlation coefficient (r) values, respectively. An absolute value of r of 1.0 to 0.5, 0.5 to 0.3, 0.3 to 0.1, and < 0.1 was considered strong, moderate, weak, and no correlation, respectively. Differences were considered statistically significant at p < 0.05. Statistical analysis was performed using Graph Pad Prism ver. 5, 2007 (Graph Pad Software Inc., San Diego, CA, USA).

Table 1. Demographic details of the different study groups

| Demographic variable | Group A | Group B | p-value |
|----------------------|---------|---------|---------|
| No. of patients      | 15      | 15      |         |
| Age (yr)             | 47.73 ± 2.33 | 41.93 ± 2.79 | 0.122<sup>a</sup> |
| Sex, male:female     | 9 (60.0):6 (40.0) | 7 (46.7):8 (53.3) | 0.457<sup>b</sup> |

Values are presented as number only, mean ± standard error of mean, or number (%). Group A, smokers; group B, nonsmokers.

<sup>a</sup>Analysis by independent samples t-test;<sup>b</sup>by chi-square test. p < 0.05, statistically significant.
(p = 0.253) after 3 months of periodontal therapy (Table 2). However, the SE level in group B increased significantly (p = 0.035) after 3 months compared to the baseline level (Table 2, Fig. 1).

Considering the SF level data, group A showed statistically significant higher values of SF than group B at baseline (group A, 95.49 ± 21.53 ng/mL and group B, 44.86 ± 4.69 ng/mL; p = 0.018). Whereas group B showed statistically lower (group A, 77.06 ± 14.06 ng/mL and group B, 39.05 ± 5.46 ng/mL; p = 0.009) SF level compared to group A after 3 months of NSPT (Table 3, Fig. 2). However, no significant alteration (p = 0.618) in SF level was observed in group A after 3 months of treatment compared to the baseline level. However, the SF level in group B decreased significantly (p = 0.039) after 3 months of NSPT compared with the corresponding baseline level (Table 3, Fig. 2).

**Discussion**

The underlying anemia status in smokers may be masked by relatively higher Hb values, which are usually tested to assess anemia. This may lead to an underestimation of the prevalence of anemia among smokers. The available literature suggests that only a few studies have analyzed the efficacy of periodontal interventions, such as NSPT, on the anemia status of such patients by assessing SE or SF.

In the present study, the baseline mean SE levels were significantly lower (p = 0.031) in smokers with CP than in nonsmokers with CP. In addition, SE levels were negatively correlated with smoking at baseline. These findings are consistent with those of Tanabe et al. [16] and Eisenga et al. [17]. During daytime smoking, higher SE levels lead to erythrocytosis, which in turn inhibits
further erythropoietin production through a negative feedback loop. Endogenous circulating erythropoietin with a half-life of 6 to 8 hours would result in low SE in the morning hours when blood samples are usually drawn [17]. This phenomenon is supported by the circadian rhythm of SE levels described by Miller et al. [18]. An alternative explanation may be derived from the study by Weinberg et al. [19] who observed the JAK2 V617F mutation in cigarette smokers and suggested that the erythrocytosis observed in smokers occurs via an erythropoietin-cell-intrinsic erythropoietin-independent mechanism. They also stated that this may be an unidentified direct effect of smoking on erythropoiesis. Chronic smoking initially induces an increase in erythrocyte volume, plasma volume, and erythropoietin concentration, the latter of which is reduced when the erythrocyte volume increases. Hence, erythropoietin production represents a balance between stimulation by hypoxia and negative feedback by increasing erythrocyte volume [14]. The release of proinflammatory cytokines from peripheral neutrophils and various parameters of inflammation in plasma seem to be affected more by cigarette smoking than periodontal disease, which might contribute to the downregulation of erythropoietin production. Elevation of these inflammatory mediators leads to inhibition of the hormone erythropoietin and erythropoiesis, leading to the development of anemia [20].

In the present study, 3 months following the NSPT intervention, SE levels increased from baseline values in group A, although they were not statistically significant \( (p = 0.253) \). In contrast, the group B patients showed a statistically significant \( (p = 0.035) \) improvement in SE levels following periodontal intervention. This is in agreement with the results of Miller et al. [21], who failed to detect differences in SE levels when the COHb concentration changed following smoking cessation. As a possible explanation, they mentioned that small changes in COHb were not sufficient to trigger an erythropoietin response in persons with normal lung function. In the present study, the mean SF levels at baseline were significantly higher \( (p = 0.018) \) in group A than in group B. Ghio et al. [22] supported this finding of increased SF levels among smokers compared with those among nonsmokers. They correlated this finding with the systemic accumulation of iron after cigarette smoke exposure and concluded that cigarette smoke alters iron homeostasis both in the lung and systemically. However, that study did not include patients with CP. Contradictory findings were obtained in a study by Erdemir et al. [23], who noted similar SF levels among smokers and nonsmokers. These findings were in agreement with those of patients with ACD, who had normal to elevated SF levels. However, the possible cause of the similar SF values in both groups was not explained in that report. A cross-sectional study conducted by Prakash et al. [24] assessed the anemia status of nonsmoking patients with CP by evaluating various hematological parameters. No significant changes in SF levels were observed between the study groups. In the present study, the mean SF levels in group B were significantly lower \( (p = 0.0397) \) at 3 months after NSPT than at baseline. This is in agreement with the study of Chakraborty et al. [13], who detected relatively higher SF levels in smokers with CP than in nonsmokers with CP, and these levels were restored to normal following NSPT intervention. The mean SF levels in the present study at 3 months after NSPT were reduced in group A compared to baseline; however, the difference was not statistically significant \( (p = 0.618) \). The available literature does not include any comparable studies.

One limitation of the present study is its relatively small sample size. Further studies involving larger sample sizes and other parameters may be conducted in the future if required.

Overall, NSPT leads to a relative increase in SE levels and a relative decrease in SF levels, thereby indicating an improvement in the anemia status of both smokers and nonsmokers. However, the magnitude of the changes was less in smokers. Hematological parameters such as SF and SE might serve as effective biomarkers for assessing anemia status in nonsmokers with CP. For smokers with CP, further studies with larger sample sizes may clearly demonstrate the effect of NSPT on SE and SF levels among these individuals.

**Notes**

**Conflicts of interest**

No potential conflicts of interest relevant to this article was reported.

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**Author contributions**

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