LEVELS OF VITAMIN D AND CALCIUM AS RISK FACTORS FOR CHRONIC PERIODONTITIS

Sandy H. S. Hassan,* Nayroz Abdel-Fattah Tarrad* and Olfat Gameel Shaker**

ABSTRACT

Objectives: This study was conducted to estimate, compare and correlate the serum and salivary vitamin D3 (VitD3) in form of [25 (OH) D] and calcium (Ca) levels in healthy subjects and chronic periodontitis (C.P) patients.

Subjects and methods: The study included 50 subjects divided into 2 groups: 30 C.P patients and 20 periodontally healthy subjects. Clinical examination was performed for all subjects, salivary and blood samples were collected from all 50 subjects. Salivary and serum VitD3 levels were assessed by ELISA assay, while Ca was measured colourimetrically.

Results: Salivary and serum VitD3 levels were significantly higher in control subjects compared to C.P patients. Salivary Ca level was significantly higher in C.P group than control group but serum Ca level showed insignificance higher level in control group than C.P patients. No significance correlation was observed except for saliva and serum VitD3 levels there was a statistical significance correlation in the control group.

Conclusion: Low salivary and serum VitD3 and high salivary Ca levels are associated with periodontitis, they might be considered as risk factors for periodontal diseases but this association need to be confirmed in more future studies.

KEYWORDS: Chronic periodontitis, vitamin D3, calcium, saliva, serum, Egyptian population

INTRODUCTION

Chronic periodontitis (C.P) is a multifactorial chronic inflammatory disease that may affect more than half of the population, caused by group of specific microorganisms that elicit activation of immune-inflammatory mechanisms which cause several adverse effects on bone and connective tissue remodeling resulting in progressive destruction of periodontal ligament and alveolar bone loss (1,2).

Vitamin D3 (VitD3) deficiency is a public health concern as 1 billion people worldwide have VitD3 deficiency [3]. Current knowledge about the non hormonal, intracrine, and paracine actions of the active form of VitD3 [25 (OH) D] has shown that it has

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antibacterial effect via increase transcription of antibacterial peptides by cells of monocyte macrophage lineage which can defend against oral pathogens. Also it has anti-inflammatory action as it can inhibit antigen-induced T-cell proliferation and cytokine production. In addition to its well-known role in regulation of calcium (Ca) homeostasis in blood by promoting Ca absorption in the intestine, reabsorption of Ca from kidneys and bone formation in order to maintain balanced plasma Ca concentration and bone mineralization, moreover its stimulation for osteoblasts to enable normal bone growth.\(^{[4-6]}\)

Ca and VitD3 are two essential elements in bone mineralization as mentioned in literature there is a tie relationship between both. Chronically low intake of Ca and VitD3 lead to negative Ca balance and bone loss as it causes a secondary increase in Ca removal from bone, consequently decreased general bone mineralization including the alveolar bone. This manifestation may contribute to weakening of the tooth attachment apparatus which is a well-recognized symptom of C.P\(^{[7]}\).

Despite the causative role of specific bacteria in the onset of C.P, its progression and severity are affected by other recently investigated modifiers as nutrition. Nutrition plays an important role in reducing the risk of developing periodontal diseases basically including vitamins and minerals as their imbalance aggravate destruction of the periodontium. Therefore, the present study was conducted to estimate, compare and correlate the serum and salivary VitD3 and Ca levels in healthy subjects and C.P patients \(^{[8]}\).

**SUBJECTS & METHODS**

**Study population**

Fifty subjects were included in this study both males and females, age range (23 - 48) years. The subjects were divided into two groups,

**Group I (Control Group):** 20 periodontally healthy volunteers who served as control subjects.

**Group II (Chronic Periodontitis Group):** 30 patients suffering from C.P. All subjects had signed an informed written consent form and the study was approved by the research ethics committee, after explaining the study procedures.

**Inclusion Criteria**

All participants were free from any systemic diseases according to the modified Cornell Medical Index \(^{[9]}\). C.P patients were selected from the Outpatient Clinic, Department of Oral Medicine, Periodontology and Diagnosis, Faculty of Dentistry, Fayoum University. Patients were diagnosed with severe C.P when having a pocket depth (PD) of $\geq$5 mm and a clinical attachment level (CAL) $\geq$5 mm, according to the *Periodontology TAAo, (2000)* \(^{[10]}\). The control group ($n=20$) was selected from healthy subjects who attended the restorative dental clinic and had clinically healthy gingiva with PD $\leq$ 3 mm, zero plaque index (PI), gingival index (GI) and CAL.

**Exclusion Criteria**

(1) pregnant women, (2) subjects taking any type of medication and/or antibiotic therapy during the 3 months before the study, (3) subjects who received periodontal treatment within the past 6 months, and (4) smokers.

**Clinical examination**

Clinical examination for all patients was performed including the following periodontal parameters PI, GI, PD and CAL. These measurements were recorded by a single calibrated expert examiner at six sites for all teeth (mesiobuccal, mesiolingual, midbuccal, distobuccal, distolingual and midlingual). PI was assessed by measuring the presence or absence of supragingival biofilm with a sweeping movement of the probe around the surfaces of all teeth \(^{[11]}\). Marginal gingival bleeding was recorded
with GI \[12\]. PD was measured from the free-gingival margin to the base of the periodontal pocket and CAL was measured from the cementoenamel junction to the base of the periodontal pocket. Measurements were rounded to the highest whole millimetre using the Michigan 0 probe with Williams’ markings.

Saliva Samples Collection.

Collection of unstimulated whole saliva was done using standard techniques \[13\]. Samples were obtained by requesting subjects to swallow first, tilt their head forward and expectorate all saliva into a tube for 5 min without swallowing. After collection, saliva samples were centrifuged at 2000xg and the supernatants were separated and stored at -80°C until subsequent analysis.

Blood samples collection.

Two milliliters of blood was collected from the antecubital vein under aseptic conditions and then transferred into a sterile test tube with anticoagulant to prevent coagulation of blood. Serum samples were separated after centrifugation and stored in aliquots at -70°C until further assays.

Method of Vitamin D3 detection

VitD3 was measured for all patients and controls using Human 25-Dihydroxy vitamin D (25-OH-D) enzyme-linked immuno sorbent assay (ELISA) kit provided by Sunredbio, Shanghai, China. This ELISA kit is based on the principle of double antibody sandwich technique to detect human 25-OH-D. The kit uses a double-antibody sandwich ELISA to assay the level of Human 25-OH-D in samples. 25-OH-D was added to monoclonal antibody enzyme well which is pre-coated with Human 25-OH -D. Incubation was done; then, 25-OH-D antibodies labeled with biotin was added, and combined with Streptavidin-HRP to form immune complex; then incubation and washing were carried out again to remove the uncombined enzyme. Then chromogen Solution A, B, were added, the color of the liquid was changed into blue. The concentration of the Human Substance 25-OH-D of sample was positively correlated.

Calcium measurement

The calcium was measured colourmetrically \[14\].

Statistical Analysis

Mean and SD values for clinical parameters, levels of Ca and VitD3 are calculated for all the study groups. Student’s t-test was used to compare between C.P and control groups for normal parametric distribution. Pearson’s correlation coefficient was used to determine correlations of VitD3 and Ca levels and also between saliva and serum for each group. P-values that were less than or equal to 0.01 level were considered statistically significant. Statistical analysis was performed using analysis of variance for multiple comparisons of means of the different groups using Tukey Contrasts at 0.01 significance level. All statistical work was done using R statistical software version 3.3.3, \[15\] using ‘Rcmdr’ package \[16,17\].

RESULTS

In the control and C.P groups the mean age range was found to be 34.4 and 37.8 years, respectively. (Table 1)

Results of Ca levels: The normal salivary Ca level in humans is 7.5–8 mg/dl. The study results demonstrated that the salivary Ca level was higher in group II than group I with a mean value of 9.4 and 7.2 mg/dl respectively. There was statistically significant difference in the salivary Ca level between C.P and control group. But for serum Ca level, the normal level in humans is 9–11 mg/dl, mean value for group I was 10.3 mg/dl while for group II was 10.1 mg/dl, showed no statistically significant difference between both groups. (Table 1, figure 1)
**Results of VitD3 levels:** Statistically significant difference was found when comparing mean values of salivary VitD3 levels in the studied groups as **group I** mean value was 462.7 pg/ml and for **group II** was 240.7 pg/ml. The serum mean value for control group was 27.25 pg/ml but it decreased with statistically significant difference in C.P group to be 15.2 pg/ml (Table 1, figure 2)

When the correlation of VitD3 level between saliva and serum was done for each of the study groups, there was a statistical significance correlation in the control group but no correlation existed in the C.P group, while there was no significance correlation between saliva and serum in both groups regarding Ca levels (Table 2)

As for the correlation between VitD3 and Ca levels in saliva in each of the included groups, no significance correlation was observed in both groups and the same regarding their correlation in serum levels (Table 2)

**TABLE (1): Characteristics of subjects presented as mean values, (±SD) and P & F values**

<table>
<thead>
<tr>
<th></th>
<th>Periodontitis group (n=30)</th>
<th>Control group (n=20)</th>
<th>Results of Student’s t-test</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>37.8 (±4.6)</td>
<td>34.4 (±5.7)</td>
<td></td>
</tr>
<tr>
<td>Ca level in serum</td>
<td>10.1mg/dl (±0.29)</td>
<td>10.3 mg/dl (±0.6)</td>
<td>F value= 2.515 P&lt; 0.12</td>
</tr>
<tr>
<td>Ca level in saliva</td>
<td>9.4mg/dl (±0.44)</td>
<td>7.2 mg/dl (±0.29)</td>
<td>F value= 335.3 P&lt;0.0001*</td>
</tr>
<tr>
<td>VitD3 level in serum</td>
<td>15.16 pg/ml (± 1.84)</td>
<td>27.25 pg/ml (±3.8)</td>
<td>F value=211.4 P&lt;0.0001*</td>
</tr>
<tr>
<td>VitD3 level in saliva</td>
<td>240.7 pg/ml ((±20.1)</td>
<td>462.7pg/ml (±27.3)</td>
<td>F value= 985.8 P&lt;0.0001*</td>
</tr>
</tbody>
</table>

*considered significant

**Analysis of variance and mean comparisons at 0.01 level**

**TABLE (2) Results of Pearson’s correlation coefficient for the correlation between VitD3 and Ca levels in saliva and serum in the study groups**

<table>
<thead>
<tr>
<th></th>
<th>Group</th>
<th>Correlation coefficients</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>VitD3 levels in saliva and serum</td>
<td>Control</td>
<td>0.63937</td>
<td>0.0077</td>
</tr>
<tr>
<td></td>
<td>VitD3 level in saliva and serum</td>
<td>0.12311</td>
<td>0.5169</td>
</tr>
<tr>
<td>Ca levels in saliva and serum</td>
<td>Control</td>
<td>0.34731</td>
<td>0.1875</td>
</tr>
<tr>
<td></td>
<td>Ca level in saliva and serum</td>
<td>0.13892</td>
<td>0.4641</td>
</tr>
<tr>
<td>VitD3 &amp; Ca levels in saliva</td>
<td>Control</td>
<td>-0.45727</td>
<td>0.0749</td>
</tr>
<tr>
<td></td>
<td>VitD3 &amp; Ca level in saliva</td>
<td>-0.23312</td>
<td>0.2151</td>
</tr>
<tr>
<td>VitD3 &amp; Ca levels in serum</td>
<td>Control</td>
<td>0.25356</td>
<td>0.3433</td>
</tr>
<tr>
<td></td>
<td>VitD3 &amp; Ca level in serum</td>
<td>0.23914</td>
<td>0.2031</td>
</tr>
</tbody>
</table>
Low intakes of specific nutrients have been found to be related to periodontal diseases. Thus, nutritional imbalance can impair tissue regeneration, healing and can increase susceptibility to oral infections, so it might affect the development, growth and maintenance of the periodontal tissues. In addition it could be a contributing factor in resistance and virulence of the infective process which determines the extent of the periodontal disease [18,19].

Ca is an essential ion within the human body, maintenance of its concentration is biologically important for the function of tissues. Literature suggests that, salivary Ca level plays an important role in the initiation of periodontal diseases. Hence VitD3 is a major modulator in Ca homeostasis and immunity plus its antibacterial function, there is a biologic rationale to assume that VitD3 and Ca levels might affect periodontal diseases. Many studies have investigated the effect of a single nutrient on periodontal disease. Although, minerals and vitamins could have interactions between them that affect their homeostasis and functionality. Therefore, the goal of this exploratory study was to investigate whether serum and salivary levels of Ca and VitD3 associated with C.P and to investigate the possibility of interactions between them [20-22].

In the herein investigation, results revealed a statistically significant increase in salivary Ca level of C.P patients compared to the healthy control group. These results are consistent with the findings of Sewon et al., 1990, they hypothesized that periodontitis affected patients had a higher remineralization potential. It seems that salivary Ca due to its high affinity for being readily taken up by plaque is a risk factor for onset of periodontitis[23]. In 2010 Khalili & Biloklytska demonstrated that there is an increase in salivary Ca concentration in periodontal disease. Moreover, increased Ca concentration was related to the clinical periodontal status of patients and associated with the severity of the disease [24]. In the same line Acharya et al., 2011 and Fyaz et al., 2013, studies results showed that, the subjects in the periodontitis group had significantly higher levels of salivary Ca than healthy groups [25,26]. Recently Patel et al., 2016 and in same year Vasavi et al., investigations concluded that, a positive correlation existed between high salivary Ca content and periodontitis [27, 28]. On the contrary, in a similar study higher salivary Ca level was related to good dental health and there was no relation to periodontal bone destruction [29].

A study performed by VargheSe et al., 2015 aiming to evaluate the existence of any disturbances in Ca metabolism and absorption induced by smoking, by quantitatively assessing the variations in the salivary Ca levels between smokers and non-smokers with periodontitis. The findings of
their study support the view that, increased salivary Ca reflecting altered Ca turnover, fluctuations in dietary Ca and enhanced mineral dissolving action of osteoclast induced bone resorption among the smokers. It might act as a risk factor for the development and progression of periodontal diseases by hampering the normal structure and function of alveolar bone. They recommended more studies to prove the clinical significance of their findings [30].

Dental plaque is the primary etiological factor responsible for periodontal disease, a polymicrobial biofilms established on tooth surfaces and retained if not removed frequently. Hence saliva is the primary source for mineralization of supragingival plaque and salivary minerals, Ca in particular is taken up by plaque. The rate of this process depends on salivary minerals concentration, if increased it harden plaque more rapidly thus indirectly influencing the level of oral hygiene so contributing in initiation and progression of C.P [28,31]. Therefore, the presented results of higher level of salivary Ca in C.P patients than in healthy controls could be attributed to these postulations.

Previous studies found a statistically significant association between low serum Ca and periodontal disease, suggesting that low serum Ca may be considered a risk factor for periodontal disease progression [32-34]. But the mean total serum Ca levels observed in our study were within the normal range. Although serum Ca was higher in control group than C.P group however, the difference was statistically insignificant. This difference may be owed to different in number of subjects and differences in the sensitivity of various ELISA kits employed in both studies.

In contrast to the presented data, Vavasi et al., 2016, demonstrated a statistically significant increase in serum Ca level between C.P and healthy group. They stated that, low dietary intake of Ca results in bone loss to meet the needs of the body which stimulates secretion of parathyroid hormone causing bone resorption which might be a cause for increased serum Ca level. In periodontitis affected individuals, alveolar bone destruction might be a cause of increased serum Ca level [28]. But recent study in 2017 conducted to correlate between serum Ca level and periodontitis, no significant correlation was observed in their results [38].

Concerning VitD3 serum level, our results demonstrated an inverse association between serum level of VitD3 and C.P as it significantly decreased in group II than group I. These results are in accordance with former studies where, Dietrich et al., 2005, had analyzed 6,700 individuals, found that individuals in the highest quintile of serum VitD3 presented significantly less bleeding as well as lower mean PD, CAL and number of missing teeth. They proposed that VitD3 has anti-inflammatory properties [36]. These results were supported by Flavia et al., 2012 as VitD3 showed positive correlation with adiponectin and negative correlation with IL-6 and leptin in addition to inverse relation with level of pathogenic bacteria [37]. Moreover, Hiremath et al., 2013 investigation showed that the optimal ranges of serum VitD3 reduce susceptibility to gingivitis as VitD3 has a dose dependent anti-inflammatory effect on gingivitis [38]. Furthermore, another conformation to the anti-inflammatory property of VitD3 demonstrated by Andrukhov et al., 2014 as there was significant reduction in secretion of proinflammatory mediators as interleukin (IL) 6, IL-8, monocyte chemotactic protein 1 by 25(OH)D in periodontal ligament cells after lipopolysaccharides stimulation [39].

A study performed by, Zhan et al., 2014 showed that increase of serum VitD3 was associated with decreased risk of tooth loss suggesting that VitD3 is a protective factor for tooth loss [40]. Recently, investigations found significantly lower levels of serum VitD3 in C.P patients compared to controls suggesting that insufficient VitD3 level might be involved in periodontal disease progression [41,42]. The previous results support our hypothesis that VitD3 hypovitaminos is a risk factor for periodontal diseases.
Oral epithelial cells are capable of converting inactive VitD3 to the active form of 25 (OH) D, which has been shown to induce expression of the antimicrobial peptide LL-37 and other defensins that combat bacterial infection so reduces tissue production of destructive matrix metalloproteinases associated with periodontal disease. Moreover, a study using saliva from healthy subject found that it could inhibit proteolysis of LL-37 by *Porphyromonas gingivalis*, a bacterium associated with C.P. This mechanism clarify how VitD3 enhances innate immune defenses against periodontal pathogenic bacteria (antibacterial property)[43-45]. Another mode of action was found in diabetic model rats with periodontitis as it decreased alveolar bone loss by lowering the expression of nuclear factor kappa beta and the phosphorylation of Janus family kinase[46]. Thus these protective properties confirm our hypothesis again of its potential role in pathogenesis of C.P considering its deficiency as risk factor.

In addition to VitD3 anti-inflammatory, antibacterial, protective property, bone and Ca homeostasis, it is owed immunomodulatory action by its capability to affect the adaptive immune response through selectively stimulating specific T-helper subsets (Th2) that might contribute in resolution of inflammation. Monocyte/macrophages can be induced to secrete molecules that have a strong antibiotic effect. Therefore these mentioned properties can explain to some extent why VitD3 level could has a role in progression of periodontal diseases[47-49].

In contrast to the presented result, Liu et al., 2009 found that plasma VitD3 level was higher among aggressive periodontitis individuals compared with control individuals [50]. This difference with the current results may be due to different studied groups as our enrolled patients were C.P affected individuals. Also Antonoglou et al., 2015 stated that there was practically no association between VitD3 level and periodontal health status [51].

Not only the serum level of VitD3 has been shown to be lower in C.P group when compared to healthy volunteers. To the best of our knowledge this investigation estimated for the first time the level of VitD3 in saliva. Hence saliva is a complex biological fluid which is an ultra filtrate of plasma, it might be used as sample of body fluid that could be collected more easily, less costly, with minimally trained personals. In addition to its well-known role in induction of periodontal diseases. One might envision that salivary VitD3 level could be used to assess periodontal health[52].

Regarding interaction between Ca and VitD3 in C.P patients, limited to our results no interaction was demonstrated as our result didn’t show significance correlation between their levels neither in serum nor saliva.

Conclusively, our findings corroborate the hypothesis that high salivary Ca level might aggravate C.P initiation and progression however we didn’t find significant relation between serum Ca level and C.P. In addition lower serum and salivary VitD3 levels in C.P group were observed suggesting its involvement in progression of periodontal diseases. Consequently they could be considered as risk factors, their estimation aid in diagnosis of C.P and improvement of the treatment modality applied. Although more investigations are required to confirm our first trial to asses salivary VitD3 level in C.P patients.

REFERENCES

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