Lack of association between diabetes mellitus and oral lichen planus in Zahedan (South-East of Iran)

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Abstract

Introduction: Oral lichen planus (OLP) is a chronic immunological disorder with unknown etiology. Some studies have reported an association between oral lichen planus and diabetes mellitus. The aim of this study was to compare the frequency of diabetes mellitus in patients with oral lichen planus and healthy persons.

Materials & Methods: This case–control study was performed on 50 patients with OLP and 50 healthy individuals. Diagnosis of OLP was confirmed by typical clinical and histopathological findings. The control group were selected randomly from healthy individuals after matching for age and sex. Blood samples were taken to achieve 5 mL for measuring fasting serum blood glucose and HbA₁C. Data were analyzed using Student’s t-test.

Results: In this case–control study, 50 patients with OLP (39 females, 11 males) with a mean age of 44.5±13.24 years and 50 healthy individuals (33 females, 17 males) with a mean age of 41.3±11.44 years were evaluated. The mean fasting blood glucose and HbA₁C levels in patient with OLP were 95.1±8.1 and 5.1±1.3 mg/dL, and 89.1±7.7 and 4.6±1.1 mg/dL in healthy individuals (P=1).

Conclusion: The frequency of diabetes mellitus was not significantly different between the case and the control groups. The results showed that diabetes mellitus does not have a direct role in the OLP etiology.

Keywords: Diabetes mellitus, Lichen planus, Oral, Etiology

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هدف: لیسانس دیابت ملیت‌س با لیکه پلان داود در زاّدان (جیّب شرق ایران)

طاهره نصرت زهی، فاطمه اربابی کلاتی، حسین عارف بور

چکیده
مقدمه: لیکه پلان دهانی یک پیام‌ری ایمونولوژیک زمینه با انتانولوژی ناشناخته است. در مطالعات مختلف همراهی لیکه پلان میلیت‌س با دیابتی اشاره کرده‌اند. هدف از این مطالعه بررسی فراوانی دیابت ملیت‌س در بیماران مبتلا به لیکه پلان دهانی در مقایسه با افراد سالم است.

مواد و روش ها: ۱۰۰ مبتلا به لیکه پلان دهان و ۱۰۰ فرد سالم انجام شد. بیمارانی که از نظر شکل بالینی شایعات دهانی و میزان فاقدی شناسی تشخیص لیکه پلان مطابقت داشتند، به عنوان بیمار وارد مطالعه شدند. آزمون گروهی از افراد فاقد بیماری پاتی ۱۴۰ موردی از جمعیت عمومی انتخاب شدند. این تعداد از افراد گرفته شد و اطلاعات ثابت گردید. جهت بررسی فاکتوری که با میزان سی ۵ سال از افراد هر دو گروه مورد آنالیز قرار گرفتند.

یافته ها: با طراحی یک آزمایش مورد شاهدی شرکت کرد. در افراد مبتلا به لیکه پلان دهانی (۴۹ (۵۷٪) از ناری و ۳۳ (۴۳٪) از مردان) رابطه ای بین لیکه پلان دهانی و میزان سی ۵ سال در افراد مبتلا به لیکه پلان دهانی (۴۴±۷/۱۸٪) با میزان سی ۵ سال در افراد سالم (۴۵±۷/۱۸٪) آزمون هیبسون (P=1، P<0.05) داشت. نتیجه گیری: فراوانی دیابت ملیت‌س تفاوت معناداری بین دو گروه مبتلا به لیکه پلان دهانی و افراد سالم را نشان داد. یافته‌ها نشان داد دیابت ملیت‌س نقش مستقیم در انتانولوژی لیکه پلان دهانی بازی نمی‌کند.

واژگان کلیدی: دیابت ملیت‌س، لیکه پلان دهانی، ایمونولوژی

Introduction
Oral Lichen Planus (OLP) is a chronic immunological disorder. Seventy-five percent of the patients with cutaneous lichen planus also experience oral lesions.[1] The etiology of OLP is unknown. In recent years, it has been found that the immune system plays a primary role in the disease.[2] On the other hand, it has been established that OLP is an immune-related disorder, and stress and anxiety are two factors causing it.[3] Diabetes Mellitus is a chronic disease involving different systems of the body with skin involvement.[4] According to reports, diabetes mellitus is associated with oral lichen planus in 14–85% of the cases.[5,6] The autoimmune background of lichen planus could support diabetes mellitus, too, due to the same pathogenesis of both diseases.[7]

Ahmed et al studied the incidence of oral lichen planus in patients with non-insulin-dependent diabetes mellitus. The study population consisted of 86 patients (49 (57%) females and 37 (43%) males). The patient’s age ranged from 40 to 70 with an average age of 51.3. They were divided into three groups: a) 40–50 years old, b) 51–60 years old, and c) over 60 years old; 6 patients (6.9%) showed histopathological signs of oral lichen planus compared with the control group with one patient (1.2%). They concluded that OLP has a significant relationship with non-insulin-dependent diabetes compared with the normal population.[8]

Bagewadi et al studied the relationship between oral lichen planus and diabetes mellitus. In the study, 150 patients were divided into three groups. The first group consisted of 50 oral lichen planus patients; the second one included 50 diabetic patients and the third group was hypertensive patients. Four and eight of 50 oral lichen planus patients, were diabetic and hypertensive patients, respectively. Only one of them had all the three diseases.[9] They argued that diabetes mellitus and
hypertension play no direct role in the etiology of oral lichen planus. However, it seems that more studies should be carried out in different geographical regions in order to confirm the co-dependency of the two above-mentioned disorders.

Materials & Methods
One hundred cases were selected from the Department of Oral Medicine in Zahedan University of Medical Sciences. The subjects were divided into two groups.

Group 1: The study group was 50 patients with oral lichen planus, which was clinically and histopathologically confirmed. OLP may exhibit both red and white components and provide, together with differing textures, the basis for the clinical classification of this disorder. The white and red components of the lesion can be a part of the following textures: reticular, plaque-like papules, bullous, erythematous, and ulcerative. The histopathological features of OLP are: (1) areas of hyperparakeratosis or hyperorthokeratosis, often with thickening of the granular cell layer and a saw-toothed appearance to the rete pegs; (2) “liquefaction degeneration” or necrosis of the basal cell layer; and (3) an eosinophilic band may be seen just beneath the basement membrane and represent fibrin covering the lamina propria. A dense subepithelial band-shaped infiltrate of lymphocytes and macrophages is also characteristic of the disease.

Group 2: The control group consisted of 50 healthy individuals from the general population (same age and sex distribution). Prior to participate in the research, patients signed an informed consent. The study was approved by the Ethics Committee of Research Deputy of Zahedan Medical Sciences University with the code of 91-1252. Patients who had more than one dermatologic disease, pregnant women, subjects with a history of neoplasia or malignancy, patients undergoing radiotherapy or chemotherapy, patients with autoimmune diseases such as lupus erythematosus, rheumatoid arthritis, Sjögren’s syndrome or metabolic diseases such as diabetes, patients taking any systemic medications (including benzodiazepines, antidepressants, anabolic steroids, OCP, corticosteroids suppressing the immune system) within the last 30 days and smokers were excluded from the study. Then, 5 mL of blood serum was collected from antecubital vein and sent to the laboratory to check FBS levels. Photometric method was used for quantitative identification of glucose in plasma. FBS≥126 mg/dL and HbA1C≥6.5 mg/dL were considered as a measure of definite diabetes (table 1).

| Table 1. Criteria for the diagnosis of diabet mellitus |

<table>
<thead>
<tr>
<th>Objective Criteria</th>
<th>Definition</th>
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<tr>
<td>1. Symptoms of diabetes plus casual plasma glucose level of 200 mg/dL or greater</td>
<td>Casual is defined as any time of day irrespective of the time elapsed since the last meal. Classic symptoms of diabetes include polyuria, polydipsia, and unexplained weight loss.</td>
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<tr>
<td>2. Fasting plasma glucose of 126 mg/dL or greater</td>
<td>Fasting is defined as no caloric intake for 8 hours or longer.</td>
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<tr>
<td>3. 2-hour plasma glucose level of 200 mg/dL or greater during an oral glucose tolerance test</td>
<td>The test should be performed using a glucose load containing the equivalent of 75 g of anhydrous glucose dissolved in water; this test is not recommended as a routine clinical examination.</td>
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<tr>
<td>4. HbA1C</td>
<td>HbA1C is used for general assessment of the long-term level of hyperglycemia in patient with diabetes.</td>
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Results
In this study, 50 patients with OLP aged 21–71, with the mean age of 44.5±13.24 years, and 50 healthy subjects aged 18–35, with the mean age of 44.3±11.44 years were studied. Of 50 patients with OLP, 29 (63%) were female and 11 (37%) were male. The most common site of involvement was the buccal mucosa (43
cases, 76%), followed by the tongue (10%), the lips (10%) and the palate (4%). The most common forms of OLP were reticular and papular (70%).

The mean fasting blood glucose and HbA1C levels in patient with OLP were 95.1±8.1 and 5.1±1.3 mg/dL, and 89.1±7.7 and 4.6±1.1 mg/dL in healthy individuals. Based on Student’s t-test, no significant differences were observed in the serum FBS levels between the two studied groups (P=1). Based on Student’s t-test, no significant differences were observed in the serum HbA1C levels between the two studied groups (P=1).

Discussion
Wilson (1869) first described lichen planus as a disease that involves skin, nails, and oral mucosa. OLP is a chronic inflammatory mucocutaneous disease mediated by T cells and has an unknown etiology. OLP exhibits periods of remission and relapse. It is a cell-dependent condition in which T lymphocytes are accumulated under the epithelium of the oral mucosa and increase the differentiation of stratified squamous epithelium, leading to hyperkeratosis and redness, with or without a wound. This disease involves 2–5% of the general population with predominance in women. Its onset is in the 4th and 5th decades of life.

In the current study, a total of 30 patients with OLP (19 women and 11 men), with a mean age of 44.5±13.24 years, and 30 healthy subjects (13 women and 17 men), with a mean age of 44.3±11.44 years, were evaluated. Lichen planus may involve different areas in the oral mucosa. The buccal mucosa is the most common site for bilateral lesions. The floor of the mouth is the rarest incidence site.

Our study showed that the most common sites of involvement, in descending order, were as follows: buccal mucosal (76%), tongue (10%), lips (10%) and palate (6%). The most common form in this study was the reticular form with 21 cases (70%).

Recently, the etiology of oral lichen planus has been considered multifactorial, where mechanical, electrochemical, trauma, infection, allergy, endocrine disorders, salivary gland disorders, heredity, immunological reactions, stress, overworking, mucous exciting factors, and habits make people susceptible to lichen planus.

For the first time, the prevalence of DM among OLP patients was reported 40% by Grinspan, et al. In last decade, In 2007 the prevalence of DM was found as 26/7% in Turkish people with lichen planus. In 2011 Ara, et al observed DM in 10% of their patients with oral lichen planus. In 2013, Narayan among 2000 diabetic patients 15 cases (0/75%) of oral lichen planus were seen. In Iran in 2012 the prevalence of DM was announced as 20% among 80 people with the age range of 44-60 years old.

Atefi and Chalkoo and Ara revealed that the prevalence of DM among patients with OLP is more than normal population. However, similar to the studies of Ansari, Borhan Mojabi, Bagewadi and Saini, our study illustrated no significant differences in FBS test results between the two groups. The present study indicated that there was no positive dependency between diabetes mellitus and oral lichen planus as 100% of oral lichen planus patients had normal FBS values and no FBS disorder was observed in the control group.

The results of the present study were not consistent with those of some other studies in terms of significant co-dependency between diabetes and oral lichen planus. The difference might be attributed to reasons such as test methods, differences in laboratory methods, unknown reasons contributing to the disease pathogenesis and non-definite relationship between the two diseases.

Conclusion
It can be concluded that oral lichen planus may not be directly associated with diabetes mellitus but could be contributing to oral lichen planus like lesions in oral cavity as a result of various medications.

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Conflict of interest: We declare no conflict of interests.
References


