#### **ORIGINAL ARTICLE**

CONTENTS 🔼

# Evaluation of clinical and demographical finding in patients with oral lichen planus: A retrospective cross sectional study

#### Shahad Ihsan Kadhum, Ban Fadhil Al-Drobie

DEPARTMENT OF ORAL DIAGNOSIS, COLLEGE OF DENTISTRY, UNIVERSITY OF BAGHDAD, BAGHDAD, IRAQ

#### ABSTRACT

Aim: To learn more about Oral Lichen Planus Iraqi patients, including their background information, symptoms, and prognosis.

**Materials and Methods:** From the Oral and Maxillofacial Pathology Department, College of Dentistry, Baghdad University, we retrospectively reviewed the medical records of 68 patients with a histologically confirmed clinical diagnosis of oral lichen planus and subsequently contacted the patients by phone to evaluate their prognosis.

**Results:** Females were more likely than males to experience severe pain; the reticular form of Oral Lichen Planus was the most prevalent at 38.2%, but the erosive type was more prevalent among females. Only 53 of 68 patients responded to phone calls. More than 37% of those respondents reported involvement at a second location intra-orally following the first oral manifestation, and 20% had extraoral Lichen Planus, and approximately 22.6% of them observed changes in the morphology and behavior of the lesion after a brief period of time, while 26.4% experienced complete remission.

**Conclusions:** Females were more likely to have oral lichen planus. Females and elderly persons were more likely to have severe pain than other. The lesion must be monitored for symptomatic flare-ups over time.

KEY WORDS: epidemiology, lichen planus, oral lichen planus, retrospective study

Wiad Lek. 2024;77(5):1025-1032. doi: 10.36740/WLek202405123 Dol 2

### **INTRODUCTION**

Oral lichen planus (OLP) is a common inflammatory mucocutaneous condition that affects the oral mucosa [1, 2]. An important involvement for the immune system is suspected, however the exact reason is yet unclear, while there are several therapy options, the disease has a protracted clinical course [3, 4]. The incidence of OLP in the general population ranges from 1% to 2% [5]. Typically, it affects middle-aged and elderly people with a female/male ratio of 2:1 [6] and the age of onset is generally between the fourth and sixth decades of life [7]. Intraoral, buccal mucosa, tongue, and gingiva are most commonly involved while other areas like mucosa of the palate and floor of the mouth are rarely affected [8]. Approximately 20% and 15% of OLP are linked to genital and cutaneous lichen planus, respectively [9]. Although OLP may cause a wide variety of oral mucosal symptoms, the most common ones are bilateral and/or numerous symmetric lesions, often accompanied by other clinical patterns [10]. It has been found that aggravation and remission occur in cycles [11]. It is common for lichen planus to appear as OLP, either alone or in association with other lesions [12], according to Shah JS et al. [13] traditionally classified

into six forms: reticular, plaque-like, popular, atrophic, erosive, and bullous. The most prevalent form of lichen planus is reticular, whereas the second most common type is erosive, which causes painful symptoms and has been linked to a probable malignant transformation of lichen planus [14, 15].

One of the most serious outcomes of OLP is oral squamous cell carcinoma (OSCC), with rates of malignant transformation ranging from 0.5% to 7%. [16, 17], As a result, the World Health Organization (WHO) designated OLP as a potentially malignant disease [18]. Therefore, it is crucial that patients with OLP be evaluated by a team of specialists, since there may be extraoral site involvement and increased risk of oral cancer. Dense infiltration of lymphocytes into the subepithelial space, lymphocyte penetration of the epithelium, and hydropic degeneration of the basal keratinocytes are histopathological hallmarks of OLP [19]. The therapy issue is complicated. The severity of the disease, the predominant clinical form of lesions, and the patient's symptoms should all be taken into account while planning a course of treatment. Asymptomatic reticular lesions often don't need treatment and may be monitored for progression [20]. The primary goals of therapy are the resolution of symptoms and the healing of atrophic and ulcerative lesions.

### AIM

The purpose of this investigation was to examine the demographic and clinical data of 68 OLP patients in Iraq.

# MATERIALS AND METHODS

The current research, according to design was an observational retrospective, and was initiated in December 2022 and continued through May 2023 at the University of Baghdad's College of Dentistry's Department of Oral and Maxillofacial Pathology. Before beginning the research, permission was granted by the Ethics Committee at the University of Baghdad's College of Dentistry (project No. 695722, December 2022). For this study, the authors accessed the medical records of all patients diagnosed with oral Lichen Planus (LP) from the oral pathological laboratory between January 2017 and May 2023. These records included information such as the patients' ages, genders, diagnosis years, types of LP, oral manifestations, and biopsy, the authors were given permission to contact the LP patients via the phone numbers provided in the reports. The primary objective of this communication was to assess the disease prognosis and record any change in the morphology or behavior of the lesion. Additional questions were asked about extra oral LP lesions, including the time of lesion onset, the type of medication being taken, and the patient's family and medical history. The interview was conducted entirely in the native language. Each participant's replies and permission were gathered after an explanation of the study's goals was provided.

STATISTICAL ANALYSIS

SPSS (version 11.5) was used to analyze the data; descriptive statistics included the usage of frequency, percentage, mean, and standard deviation. Pain, oral clinical presentation, location, and type of OLP served as independent factors studied using the chi-square test for correlation with dependent variables (sex and age), a significant level was set at p<0.05.

# RESULTS

In total, the medical records of 68 patients with a clinically and histopathologically confirmed diagnosis of OLP were studied (Table 1). The sample population was composed of 43 (63.2%) females and 25 (36.8%) males (ratio F:M = 2:1). The mean age at diagnosis was 49.25  $\pm$  14.88 (46.36  $\pm$  16.28 years for males) and (50.34  $\pm$  14.36 years for females), and the peak of age-frequency distribution was the fifth decade (32.35%) of life (Fig.1).

Over 70% of the oral OLP had lesions in multiple oral cavity regions. The buccal mucosa was most commonly affected, followed by the tongue and gingiva, and the floor of the mouth was least affected (Fig.2), the most prevalent form of OLP was the reticular type (Fig.3).

Patients with reticular OLP accounted for 38.23% of all cases, erosive type for 29.41%, pigmented type for 4.4%, while patients with bullous OLP and atrophic OLP each accounted for 2.9% of all cases.

Buccal mucosa was the most prevalent place (69.1%), followed by tongue (17.6%) and gingiva (10.3%), and OLP lesions were lowest on the mouth floor (2.9%). The erosive, pigmented, bullous, and atrophic types were also documented. Predominantly males exhibited the reticular form (n=15) and while the erosive type was predominantly observed in females (n=20). Pain perception was shown to vary significantly across the sexes statistically. Patients with oral LP who were older than 40 years old reported significantly higher pain levels than their younger counterparts. There were no statistically significant differences between sexes or age groups in terms of clinical presentation, implicated locations, or OLP categories (Table 2).

Dysplastic alterations were seen on histology in 13 (19.1%) of the samples. There were five reticular samples and eight erosive instances. No correlations were found between dysplastic alterations and demographic factors such as age, gender, location, or OLP type. Only 53 patients could be reached by phone, according to the author in charge of communication (Table III) including 31 women and 22 males, with an average age of 49.64  $\pm$  15.66 years and the average of the age of onset was 47.59 ± 14.38, most of them were non-smoker 33 (62.26%) and non-drinker 52 (98.11%), and about 11 (20.8%) of them have family history of OLP and while 21 (39.6%) have medical history of systemic disease. Lesion involved another site intraorally in 20 (37.73%) of patients after period of time, and another involved extraoral LP in 25 (47.2%) after diagnosed the lesion orally. After re-call, 12 (22.64%) of them have a change in morphology and behavior of lesion, with 14 (26.41%) of the having total remission of lesion. No malignant transformation was documented. Corticosteroids and analgesic were the drugs prescribed for 22 (38.59%) of them, while 31 (58.49%) of patients don't used any medication (Table 3).

# DISCUSSION

Oral lichen planus is a chronic inflammatory mucucatenous disease that has the potential to become malignant

|                       | pianas (0=1) patients |       |  |  |
|-----------------------|-----------------------|-------|--|--|
| Davamatar             | Patients              |       |  |  |
| Parameter             | Abs.                  | %     |  |  |
| Sex                   |                       |       |  |  |
| Male                  | 25                    | 36.8% |  |  |
| Female                | 43                    | 63.2% |  |  |
| Male: female          | 1:2                   |       |  |  |
| Age (years) Mean ± SD | 49.25±14.88-          |       |  |  |
| Min–max               | 13–7                  | б     |  |  |
| Site of oral biopsy   |                       |       |  |  |
| Cheek                 | 47                    | 69.1% |  |  |
| Tongue                | 12                    | 17.6% |  |  |
| Gingiva               | 7                     | 10.2% |  |  |
| Type of OLP           |                       |       |  |  |
| White type            | 37                    | 54.4% |  |  |
| Red type              | 31                    | 45.5% |  |  |
| Pain                  |                       |       |  |  |
| Symptomatic           | 40                    | 58.8% |  |  |
| Asymptomatic          | 28                    | 41.1% |  |  |

#### Table 1. The demographic and clinical characteristics of oral lichen planus (OLP) patients

Table 2. Correlation between age and gender and other socioeconomic variables

| Parameter          | Male     | Female      | Age (years) <40<br><sup>n,%</sup> | <b>≥40</b><br>n,% |
|--------------------|----------|-------------|-----------------------------------|-------------------|
|                    |          | Pain        |                                   |                   |
| Symptomatic        | 6, 8.8   | 34, 50.0    | 6, 15.0                           | 34, 85.0          |
| Asymptomatic       | 19, 27.9 | 9, 13.2     | 12, 42.8                          | 16, 57.1          |
| p-value*           | 0.001    |             | 0.01                              |                   |
|                    |          | Site        |                                   |                   |
| Buccal mucosa      | 15, 31.9 | 32, 68.08   | 14, 51.9                          | 13, 48.1          |
| Tongue             | 5, 41.66 | 7, 58.33    | 10, 62.5                          | 6, 37.5           |
| Gingiva            | 4, 57.1  | 3, 42.85    | 2, 50.0                           | 2, 50.0           |
| Floor of the mouth | 1, 50.0  | 1, 50.0     | 12, 57.1                          | 9, 42.9           |
| p-value*           | 0.8      |             | 0.3                               |                   |
|                    |          | Type of OLP |                                   |                   |
| White type         | 16, 43.2 | 21, 56.75   | 12, 52.2                          | 11, 47.8          |
| Red type           | 9, 29.0  | 22 ,70.96   | 3, 50.0                           | 3, 50.0           |
| p-value*           | 0.4      |             | 0.5                               |                   |

\*Bold font indicates significance at p < 0.05 by Chi-square test.

[21]. The primary conclusions of the present observational research were that the main age was 43 years and that women were more impacted than men. In the majority of instances, the oral cavity was where the initial OLP lesions appeared. Aging and the gender (female) were linked to the pain becoming worse. In all, only 53 people answered the phone, and after receiving corticosteroids and analgesics, 14 of them claimed complete remission. According to our research, OLP is more common among women than in men, that is in consistent with the results of prior international epidemiological investigations [22-24]. It is probable that estrogen has a substantial role in the increased occurrence of OLP in women, since estrogen has been proven to promote immunological reactivity [25]. However, some researchers have presented the opposite, showing a greater rate of OLP in men than females [26]. In this sample of patients, OLP often first appeared in the fifth decades of age. Comparable findings have been reported from other surveys in other nations [27-29], however, other











Fig. 3. Distribution of pa-

tients based on OLP type at the time of diagnosis.

| <b>.</b> .                                     | Patients                            |              |  |  |  |  |
|--|-------------------------------------|--------------|--|--|--|--|
| Parameter —                                    | No.                                 | %            |  |  |  |  |
|  | Sex                                 |              |  |  |  |  |
| Male   | 22                                  | 41.50%       |  |  |  |  |
| Female   | 31                                  | 58.49%       |  |  |  |  |
| Age (years) Mean±SD                            | 4                                   | 9.64 ± 15.66 |  |  |  |  |
| Medical history of systemic disease            |                                     |              |  |  |  |  |
| Yes  | 21                                  | 39.6%        |  |  |  |  |
| No   | 32                                  | 60,3%        |  |  |  |  |
|  | Family history of OLP               |              |  |  |  |  |
| Yes  | 8                                   | 15%          |  |  |  |  |
| No   | 45                                  | 84%          |  |  |  |  |
|  | Another site involved later intraor | ally         |  |  |  |  |
| Yes  | 20                                  | 37.7%        |  |  |  |  |
| No   | 33                                  | 62.2%        |  |  |  |  |
|  | Another site involved later extraor | ally         |  |  |  |  |
| Yes  | 11                                  | 20.7%        |  |  |  |  |
| Skin   | 7                                   | 63.6%        |  |  |  |  |
| Genital area                                   | 3                                   | 27.2%        |  |  |  |  |
| Nail   | 1                                   | 9.0%         |  |  |  |  |
| No   | 42                                  | 79.2%        |  |  |  |  |
| Healing  |                                     |              |  |  |  |  |
| Total remission of lesion                      | 14                                  | 26.41%       |  |  |  |  |
| No change in morphology and behavior of lesion | 25                                  | 47.16%       |  |  |  |  |
| Change in morphology and behavior of lesion    | 12                                  | 22.64%       |  |  |  |  |
|  | Medication                          |              |  |  |  |  |
| Don't administrate any medication              | 31                                  | 58.49%       |  |  |  |  |
| Administrate medication                        | 22                                  | 38.59%       |  |  |  |  |
| Topical medication                             | 14                                  | 63.63%       |  |  |  |  |
| Systemic medication                            | 8                                   | 36.36%       |  |  |  |  |

research has shown that the average age of OLP is in the fourth decade of life [30]. These variations in findings may have several causes, the most prominent of which are sample size, genetic predisposition, and other confounding variables. Oral manifestations of LP lesions include reticular lesions in 38.23% and erosions in 29.41% of cases. These trends mirrored those seen by Gotmare et al. [31], who reported that reticular and erosive OLP were the most prevalent types. Lesions caused by oral lichen planus (OLP) may appear anywhere in the mouth, however the buccal mucosa is the most usually affected region [32, 33] which also was the pattern observed in the present study. Patients' primary complaint in this study was severe pain. Consistent with earlier research, this study confirmed that pain is a common symptom of OLP [34]. A cluster of cytotoxic (CD8) T cells very near to

the surface of the epithelium, leading to an exaggerated reaction to environmental triggers [35]. In addition, the pain experienced by women was much more intense than that experienced by men. It has been argued in the past that men have a greater pain tolerance or threshold than females [36]. There is currently no agreed-upon explanation for the observed disparity in pain perception between the sexes. However, it is possible that greater sensitivity to pain as we age contributes to the gradual decline in estrogen levels in women [37, 38]. Of the 68 OLP patients that were called, only 53 (77.94%) answered. There might be a variety of factors preventing contact with non-responding patients, including a change in contact information, relocation to another country, or even death. According to the responding patients, eight patients have a positive family history

of OLP. This result was consistent with findings from previous studies indicating that lichen planus patients have a positive family history. A higher frequency of human leukocyte antigen B7 (HLA-B7) has been identified in affected families [39]. Since OLP patients may be carriers of a disease with systemic consequences, a multidisciplinary team may be necessary for their management [40]. Twenty-one individuals with systemic diseases (such as hypertension, diabetes, or thyroid disease) were included in our research. OLP is a chronic inflammatory disorder that may be a precursor to cancer [16]. Inflammation has been shown to be a significant risk factor for the development of cancer in a number of studies. Oral and pharyngeal mucosal diseases that may progress to cancer are strongly linked to cigarette smoking [41], we tracked 20 smokers throughout the course of our study and found that some of them observed a progression from one lesion type to another. It's important to highlight that a patient with oral lichen planus may also have lichen planus lesions in other places of his body, according to a previous study [32]. Approximately 20% of patients contacted had further oral lesions from OLP after the original lesion appeared and 11% of patients had extraoral lesions 1-2 years after the initial oral presentation. In addition, we identified fourteen patients whose OLP spontaneously remitted, contrary to a previously reported finding that spontaneous remission of OLP is exceedingly uncommon [42, 43]. The development of oral cancer is the most serious consequence of OLP, although this was not seen in our study. The reported rate of malignant transformation

of OLP is 0 to 10% [44], giving our patient group one of the lowest incidences of malignant transformation. As the etiology of OLP remains obscure, no etiological treatment is currently available [45]. The goal of therapy is to reduce the disease's functional effect and relieve symptoms. The majority of patients with asymptomatic reticular lesions do not require any form of treatment. In contrast, the majority of erosive lesions are extremely excruciating, necessitating treatment in these patients. Corticosteroids are typically the treatment of choice for OLP. The key limitations of this research are the absence of a clinical evaluation and the need of a bigger sample size. The prognosis was also determined through telephone conversation rather than via a physical examination. Furthermore, higher-level clinical studies establish causality, whereas observational research data just show correlation. Caution is warranted until further research confirms the conclusions of the present study, which provided results of illnesses linked to a recurrence rate and malignant transformation.

# CONCLUSIONS

Results indicated that OLP was more prevalent among female Iraqis. Severe pain in OLP patients is strongly associated with females and older age groups; clinically, the disease undergoes remission and exacerbation, and all patients must be carefully monitored. All OLP patients should be followed up on a periodic basis. The correct diagnosis of any pathology is crucial for producing effective treatment and minimizing iatrogenic harm.

### REFERENCES

- 1. Al-Azzawi LM, Al-Ani LS. Immunohistochemical expression of p53 and PCNA proteins in oral lichen planus and oral dysplasia. Journal of Baghdad College of Dentistry. 2014;26(1):98-102.
- 2. Mohammed AJ, Diajil AR. Salivary vitamin E and uric acid in patients with OLP and healthy individuals. Journal of Baghdad College of Dentistry. 2019;31(3):39-43. doi:10.26477/jbcd.v31i3.2699. •••
- 3. Sriram S, Hasan S, Alqarni A et al. Efficacy of Platelet-Rich Plasma Therapy in Oral Lichen Planus: A Systematic Review. Medicina (Kaunas). 2023;59(4):746. doi:10.3390/medicina59040746.
- 4. Andabak-Rogulj A, Vindiš E, Aleksijević LH et al. Different Treatment Modalities of Oral Lichen Planus-A Narrative Review. Dent J (Basel). 2023;11(1):26. doi:10.3390/dj11010026.
- 5. Agha-Hosseini F, Moosavi MS, Ghaffarpour M. Investigating the factors proposed in oral lichen planus malignant transformation: A literature review. Health Sci Rep. 2023;6(5):e1267. doi:10.1002/hsr2.1267. 00120
- 6. Gamal-Abdelnaser A. Oral Lichen Planus in Childhood with Unique Histological Finding: A Case Report. Duzce Medical Journal. Duzce Med J. 2023;25(1):85-88. doi:10.18678/dtfd.1182644.
- 7. Shen ZY, Liu W, Zhu LK et al. A retrospective clinicopathological study on oral lichen planus and malignant transformation: analysis of 518 cases. Med Oral Patol Oral Cir Bucal. 2012;17(6):e943-e947. doi:10.4317/medoral.17778.
- 8. locca O, Copelli C, Rubattino S et al. Oral cavity carcinoma in patients with and without a history of lichen planus: A comparative analysis. Head Neck. 2023;45(6):1367-1375. doi:10.1002/hed.27350.
- 9. Sulewska ME, Tomaszuk J, Sajewicz E et al. Treatment of Reticular Oral Lichen Planus with Photodynamic Therapy: A Case Series. J Clin Med. 2023;12(3):875. doi:10.3390/jcm12030875.

- 10. Iqbal MA, Yesmin S, Maaisha F et al. Oral lichen planus and its recent management: A review. Update Dental College Journal. 2020;10(2):29-34. doi:10.3329/updcj.v10i2.50179.
- 11. El-Howati A, Thornhill MH, Colley HE et al. Immune mechanisms in oral lichen planus. Oral Dis. 2023;29(4):1400-1415. doi:10.1111/ odi.14142.
- 12. Radochová V, Dřízhal I, Slezák R. A retrospective study of 171 patients with oral lichen planus in the East Bohemia Czech Republic single center experience. J Clin Exp Dent. 2014;6(5):e556-e561. doi:10.4317/jced.51784.
- 13. Shah JS, Prajapati MN. Various Faces of Lichen Planus: A Clinical Study. National Journal of Integrated Research in Medicine. 2017;8(1):82-87.
- 14. Patigaroo SA, Ali I, Maqbool T et al. Reticular Oral Lichen Planus: A Clinical Experience of ENT Surgeons. Indian J Otolaryngol Head Neck Surg. 2023;75(2):390-396. doi:10.1007/s12070-022-03267-y.
- 15. Veneri F, Bardellini E, Amadori F et al. Efficacy of ozonized water for the treatment of erosive oral lichen planus: a randomized controlled study. Med Oral Patol Oral Cir Bucal. 2020;25(5):e675-e682. doi:10.4317/medoral.23693.
- 16. Diajil A, Robinson CM, Sloan P et al. Clinical outcome following oral potentially malignant disorder treatment: a 100 patient cohort study. Int J Dent. 2013;2013:809248. doi:10.1155/2013/809248. DOI 2010
- 17. Luqman N, Asgher R. Frequency of oral involvement in cutaneous lichen planus patients. Journal of Pakistan Association of Dermatologists. 2021;31(3):454-458.
- 18. Agha-Hosseini F, Barati H, Moosavi MS. Aquaporin3 (AQP3) expression in oral epithelium in oral lichen planus. Exp Mol Pathol. 2020;115:104441. doi:10.1016/j.yexmp.2020.104441.
- 19. Georgieva I. Oral lichen planus-clinical characteristics and diagnosis. A review. Scripta Scientifica Medicinae Dentalis. 2022;7(2):39-44. doi:10.14748/ssmd.v7i2.8088.
- 20. Santonocito S, Polizzi A, De Pasquale R et al. Analysis of the Efficacy of Two Treatment Protocols for Patients with Symptomatic Oral Lichen Planus: A Randomized Clinical Trial. Int J Environ Res Public Health. 2020;18(1):56. doi:10.3390/ijerph18010056.
- 21. Tsushima F, Sakurai J, Uesugi A et al. Malignant transformation of oral lichen planus: a retrospective study of 565 Japanese patients. BMC Oral Health. 2021;21(1):298. doi:10.1186/s12903-021-01652-7.
- 22. Liao H, Luo Y, Long L et al. Anxiety and oral lichen planus. Oral Dis. 2021;27(3):506-514. doi:10.1111/odi.13569.
- 23. Giuliani M, Troiano G, Cordaro M et al. Rate of malignant transformation of oral lichen planus: A systematic review. Oral Dis. 2019;25(3):693-709. doi:10.1111/odi.12885.
- 24. Radochová V, Koberová Ivančaková R, Heneberk O et al. The Characteristics of Patients with Oral Lichen Planus and Malignant Transformation-A Retrospective Study of 271 Patients. Int J Environ Res Public Health. 2021;18(12):6525. doi:10.3390/ijerph18126525. DOI 2010
- 25. Gupta A, Mohan RP, Gupta S et al. Roles of serum uric acid, prolactin levels, and psychosocial factors in oral lichen planus. J Oral Sci. 2017;59(1):139-146. doi:10.2334/josnusd.16-0219.
- 26. Munde AD, Karle RR, Wankhede PK et al. Demographic and clinical profile of oral lichen planus: A retrospective study. Contemp Clin Dent. 2013;4(2):181-185. doi:10.4103/0976-237X.114873.
- 27. Lauritano D, Arrica M, Lucchese A et al. Oral lichen planus clinical characteristics in Italian patients: a retrospective analysis. Head Face Med. 2016;12:18. doi:10.1186/s13005-016-0115-z. DOI 2012
- 28. Joseph BK, Ali MA, Dashti H et al. Analysis of oral and maxillofacial pathology lesions over an 18-year period diagnosed at Kuwait University. J Investig Clin Dent. 2019;10(4):e12432. doi:10.1111/jicd.12432. DOI 2019
- 29. Queiroz EP, Collicchio LA, Utumi ER et al. Simultaneous presentation of plaque-like and reticular lichen planus: Case Report. SVOA Dentistry. 2022;3(1):1-4.
- 30. Pavlic V, Vujic-Aleksic V. Phototherapy approaches in treatment of oral lichen planus. Photodermatol Photoimmunol Photomed. 2014;30(1):15-24. doi:10.1111/phpp.12074. DOI 2014;30(1):15-24. doi:10.1111/phpp.12074.
- 31. Gotmare SS, Gupta AA, Waghmare M et al. A Comparison of Proliferative Capacity of Reticular and Erosive Variants of Oral Lichen Planus by Argyrophilic Nucleolar Organizer Regions Method. J Microsc Ultrastruct. 2022;11(1):12-16. doi:10.4103/jmau.jmau\_104\_20.
- 32. Gupta S, Jawanda MK. Oral Lichen Planus: An Update on Etiology, Pathogenesis, Clinical Presentation, Diagnosis and Management. Indian J Dermatol. 2015;60(3):222-229. doi:10.4103/0019-5154.156315.
- 33. Elsabagh HH, Gaweesh YY, Ghonima JK et al. A novel comprehensive scoring system for oral lichen planus: A validity, diagnostic accuracy, and clinical sensitivity study. Oral Surg Oral Med Oral Pathol Oral Radiol. 2021;131(3):304-311. doi:10.1016/j.oooo.2020.12.016.
- 34. Serafini G, De Biase A, Lamazza L et al. Efficacy of Topical Treatments for the Management of Symptomatic Oral Lichen Planus: A Systematic Review. Int J Environ Res Public Health. 2023;20(2):1202. doi:10.3390/ijerph20021202. DOI 20
- 35. Alshami ML, Al-Rikaby HH, Majeed AA. Oral lichen planus: A review study. Journal of Pakistan Association of Dermatologists. 2022;32(3):574-584.
- 36. Rahim-Williams FB, Riley JL 3rd, Herrera D et al. Ethnic identity predicts experimental pain sensitivity in African Americans and Hispanics. Pain. 2007;129(1-2):177-184. doi:10.1016/j.pain.2006.12.016.

- 37. Alshami ML, Aswad F, Abdullah B. A clinical and demographic analysis of oral pemphigus vulgaris: A retrospective cross-sectional study from 2001 to 2021. Health Sci Rep. 2022;5(5):e832. doi:10.1002/hsr2.832.
- Roeder HJ, Leira EC. Effects of the Menstrual Cycle on Neurological Disorders. Curr Neurol Neurosci Rep. 2021;21(7):34. doi:10.1007/ s11910-021-01115-0.
- 39. Dissemond J. Oral lichen planus: an overview. J Dermatolog Treat. 2004;15(3):136-140. doi:10.1080/09546630410030720.
- 40. Cassol-Spanemberg J, Rodríguez-de Rivera-Campillo ME, Otero-Rey EM et al. Oral lichen planus and its relationship with systemic diseases. A review of evidence. J Clin Exp Dent. 2018;10(9):e938-e944. doi:10.4317/jced.55145.
- 41. Sami A, Elimairi I, Stanton C et al. The Role of the Microbiome in Oral Squamous Cell Carcinoma with Insight into the Microbiome-Treatment Axis. Int J Mol Sci. 2020;21(21):8061. doi:10.3390/ijms21218061.
- 42. Carbone M, Arduino PG, Carrozzo M et al. Course of oral lichen planus: a retrospective study of 808 northern Italian patients. Oral Dis. 2009;15(3):235-243. doi:10.1111/j.1601-0825.2009.01516.x.
- 43. Budimir V, Richter I, Andabak-Rogulj A et al. Oral lichen planus retrospective study of 563 Croatian patients. Med Oral Patol Oral Cir Bucal. 2014;19(3):e255-e260. doi:10.4317/medoral.18940.
- 44. Landini G, Mylonas P, Shah IZ et al. The reported rates of transformation of oral lichen planus. Journal of Oral and Maxillofacial Surgery, Medicine, and Pathology. 2014;26(2):213-220. doi:10.1016/j.ajoms.2013.04.015.
- 45. Elenbaas A, Enciso R, Al-Eryani K. Oral lichen planus: a review of clinical features, etiologies, and treatments. Dentistry Review. 2022;2(1):100007. doi:10.1016/j.dentre.2021.100007. 00120

The research received ethical clearance from the Institutional Review Committee (IRC) of the University of Baghdad's faculty of dentistry (project NO.6957722, Dec. 2022).

### **CONFLICT OF INTEREST**

The Authors declare no conflict of interest

### **CORRESPONDING AUTHOR**

### Shahad Ihsan Kadhum

Department of Oral Diagnosis, College of Dentistry, University of Baghdad, Baghdad, Iraq e-mail: ahad.ihsan2206@codental.uobaghdad.edu.iq

### **ORCID AND CONTRIBUTIONSHIP**

Shahad Ihsan Kadhum: 0009-0005-4845-7653 B C D E Ban Fadhil Al-Drobie: 0000-0002-9796-9322 A F

A – Work concept and design, B – Data collection and analysis, C – Responsibility for statistical analysis, D – Writing the article, E – Critical review, F – Final approval of the article

**RECEIVED:** 27.01.2024 **ACCEPTED:** 25.04.2024

