

Correlation Of Clinical Severity Of Oral Lichen Planus With Treatment Prognosis - A Retrospective Institution Based Study

Research Article

Krishnapriya Umashankar¹, Hannah. R²¹Department of Oral Pathology, Saveetha Dental College and Hospital, Saveetha Institute of Medical and Technical Sciences (SIMATS), Saveetha University, Chennai-600077, India.²Senior Lecturer, Department of Oral Pathology, Saveetha Dental College and Hospital, Saveetha Institute of Medical and Technical Sciences (SIMATS), Saveetha University, Chennai-600077, India.

Abstract

Lichen planus is an immunologically mediated mucocutaneous disease that is triggered by varied aetiological agents. Lichen planus shows many clinical features affecting skin, oral cavity, genital organ, nail and scalp. Lichen planus has well documented clinical findings and histological findings that aid in diagnosis. The objective of this retrospective study was to assess the clinical severity of oral Lichen Planus (OLP) with the treatment prognosis of the patients visiting Saveetha dental College and hospital. In the present study a total of 60 clinically diagnosed Lichen planus patients were included. Clinical and treatment details were recorded. All the collected data were analysed by appropriate statistics using SPSS software. The results revealed 60% of the cases to be females with 58.3% accounting for erosive type of lichen planus. 60% had involvement of bilateral buccal mucosa. Erosive variant showed 8 months duration of treatment using systemic steroids. Within the limitations of the study we can conclude that females are more commonly affected than males with erosive lichen planus being the most common variant which has shown maximum treatment duration using systemic steroids. Follow up is advocated for these patients.

Keywords: Lichen Planus; Females; Buccal Mucosa; Steroids.

Introduction

Oral Lichen Planus (OLP) is defined as a chronic mucocutaneous inflammatory disease of immune origin [7, 28]. There are various factors which play a major role in the progression of the disease [28, 34]. These factors include stress, anxiety, hormonal imbalance, menopause, drugs [7, 28]. It is the most common type of mucocutaneous lesion affecting 2 to 5% of the general population [7, 28]. Females are more commonly affected [28]. Its onset is in the 4th to 5th decade of life [28]. Intraorally it involves the buccal mucosa, tongue although more commonly and also other sites such as floor of the mouth are rarely affected [28]. It presents clinically as a wide range from asymptomatic white keratotic lesions to painful erosions and alterations [5, 6]. It is clinically seen in various forms such as reticular, papular, plaque like, erosive, atrophy, bullous [6, 17]. The most common types are reticular and erosive form [6]. The epidemiological distribution of the type of

OLP varies in each geographical region depending on their lifestyle, habits and other associated immune related factors [11, 23].

OLP is a T-cell mediated autoimmune disease in which the cytotoxic CD8+ cells triggers apoptosis of the basal cells of the oral epithelium [2, 19, 30, 31]. Further the T cells migrate into the epithelium either due to random encounter of antigen during routine surveillance in the basal keratinocyte [2, 19, 30, 31]. These migrated T cells directly bind to the MHC1 on keratinocyte or via the activated CD4 positive lymphocytes [2, 19, 30, 31]. This releases various factors such as IL2, IFN gamma, TNF alpha which in turn destroys the basal keratinocytes [2, 19, 30, 31].

Histopathological OLP is characterised by hydropic degeneration of basal epithelial cells with intra epithelial and dense subepithelial lymphocytes infiltrate [4, 13, 24, 39]. The WHO classified OLP as a potentially malignant disorder with a malignant transforma-

*Corresponding Author:

Dr. Hannah. R,

Senior Lecturer, Department of Oral Pathology, Saveetha Dental College and Hospital, Saveetha Institute of Medical and Technical Sciences (SIMATS), Saveetha University, Chennai-600077, India.

Tel: +91-9962071806

Email Id: hannahr.sdc@saveetha.com

Received: January 12, 2021**Accepted:** January 22, 2021**Published:** January 30, 2021

Citation: Krishnapriya Umashankar, Hannah. R. Correlation Of Clinical Severity Of Oral Lichen Planus With Treatment Prognosis - A Retrospective Institution Based Study. *Int J Dentistry Oral Sci.* 2021;08(01):1518-1522. doi: <http://dx.doi.org/10.19070/2377-8075-21000302>

Copyright: Hannah. R[©]2021. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution and reproduction in any medium, provided the original author and source are credited.

tion rate of approximately 1.37% [4, 35, 36].

Many studies have been done on the pathogenesis of OLP, risk factors, treatment, disease characteristics and its rate of malignant transformation. This is the first kind of study done at an institutional level to assess the clinical severity of OLP mainly based on the symptoms and the type of OLP and also to correlate this with the treatment prognosis. Thereby it helps us to identify the association of clinical severity with the duration of treatment of OLP with steroid therapy. This study could serve as a basis to understand the pattern of disease with its treatment. Hence it could help the clinician in emphasising the need for treatment to the patient and also to predict the duration of treatment of OLP.

Materials And Methods

A cross-sectional, observational retrospective study was conducted. This study was approved by the scientific review board of Saveetha dental College and Hospital, Chennai. The sample consisted of patients with a diagnosis of OLP which had been followed up between July 2019 to February 2020 consisting of 60 patients. The clinical data of the patients visiting the institution were retrieved from the DIAS online patient portal. The following data were obtained: age, gender, symptoms, clinical presentation, Habits, treatment done and duration of treatment done with associated skin lesions were also evaluated.

The descriptive variables are quantified using bar graphs and frequency tables. Chi-square test was done for further clinical presentation, treatment done and duration of the treatment. $P < 0.05$ was considered to be statistically significant.

Results And Discussion

Among 60 patients diagnosed during the period of June 2019 to February 2020 60% of the affected individuals with females and 40% were males. 72% of the affected were not associated with any habit and 28% had the habit of smoking and chewing. The common site was buccal mucosa (8.3%) followed by tongue (8.3%) and Gingiva (3.3%). Most of the patients, 58.3% had erosive type of lichen planus followed by reticular type which accounts for 35%. Pigmented lichen planus accounted for 6.7 % of the population. All these details are described in Table 1, Figure 1 and 2 respectively. Burning sensation was the most common symptom seen in patients which accounts for nearly 60%. When the clinical variants of oral lichen planus was correlated with gender, it was not found to be statistically significant $P = 0.769$ (Figure 3). However, Occurrence of erosive type of oral lichen planus was more among females than males. When duration of treatment was correlated with the type of OLP, it was not found to be statistically significant $P = 0.134$ (Figure 4). However erosive lichen planus exhibits maximum treatment duration when compared to reticular type. There was no evidence of malignant transformation in the OLP cases reported during the period of study.

The clinical characteristics of patients included in this study was similar to that of the previous studies, although few differences were noted. Retrospective studies have limitations and cannot be compared satisfactorily to prospective studies. However they are useful in evaluating patient populations.

According to the clinical and histopathological criteria of the WHO the results of the study revealed that OLP is seen in middle-aged patients around 40 to 60 years with sex predilection for

Table 1. Demographic data of the population.

	Demographics	Percentage
Gender	Female	60%
	Male	40%
Age	20-40 years	34%
	40-60 years	66%
Type	Erosive	58.30%
	Reticular	35%
	Pigmented	6.70%

Figure 1. Bar graph depicting the frequency of site of occurrence of oral lichen planus. X axis showing the site of occurrence and Y axis indicating the frequency. 90% of the cases were seen in the buccal mucosa , 8.3% were seen in the tongue and 1.7% were seen in the gingiva.

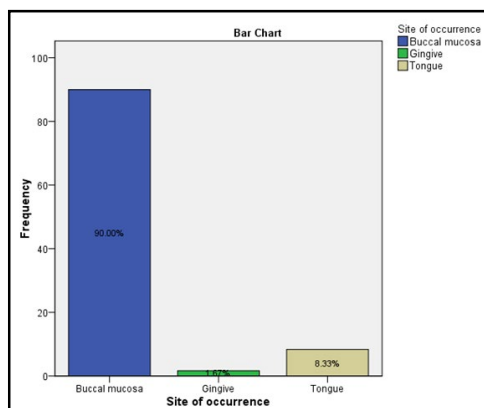


Figure 2. Frequency of occurrence of various clinical variants of oral lichen planus with X axis showing the clinical variants and Y axis showing the frequency. 58.3% of the cases were Erosive, 35% were Reticular and 6.7% were pigmented.

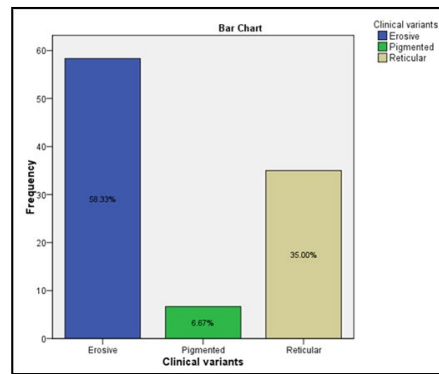


Figure 3. Correlation between gender and the different clinical variants of oral lichen planus with X axis depicting the clinical variants and Y axis depicting the frequency of occurrence in male and female. Erosive lichen planus was more commonly seen among the females than the males. However, Chi square analysis shows no statistical significance with P = 0.769 (P>0.05).

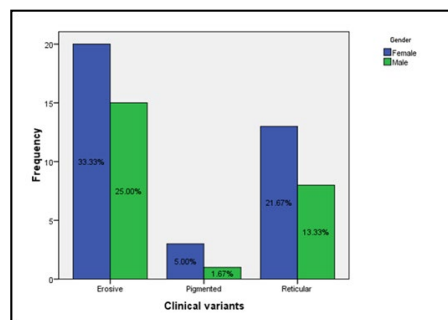
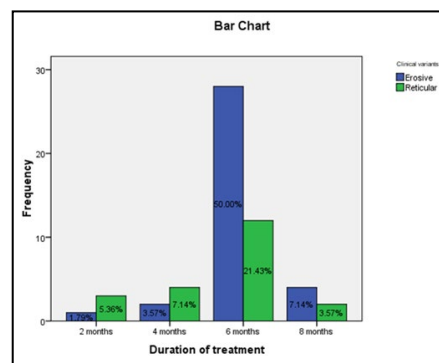


Figure 4. Correlation between the clinical variants of oral lichen planus and the duration of treatment. X axis depicts the duration of treatment and Y axis depicting the clinical Variants. Maximum duration of treatment is for erosive lichen planus when compared with reticular type. Chi square analysis shows no statistical significance with P=0.134 (P>0.05 statistically not significant).



females. The most commonly affected sites were buccal mucosa, gingiva and tongue. The male to female ratio is 2:3 which is in agreement with the other studies. Most of the studies in the other parts of the world had also a similar female predominance [12, 26-28, 37]. This could be attributed to hormonal imbalance, frequent use of medications such as paracetamol for pain, allergy to dentifrices [16]. OLP was more prevalent in the fourth decade of life in our study (mean age was 42.1 years) which is lower than the mean age group reported in central China (50.4 years), UK (52 years), Spain (56.4 years) and Italy (56.7 years) [12, 26-28, 37]. This was probably due to the ethnic population and geographic differences in our study when compared to previous studies. OLP in minor juveniles or children is uncommon and in our study childhood form of OLP was not observed [3, 16]. This could be attributed to the rarity of associated autoimmune conditions, exposure of drug and dental restorative materials, infective agents

and other environmental triggers that have been known to initiate lichen planus (Thapa and Malathi 2016).

As previously mentioned, the lesions of OLP were bilateral, symmetrical and buccal mucosa was the most commonly affected site [3, 14, 16, 37]. Buccal mucosa concomitant with gingiva was the most common multiple oral site [27, 40]. Isolated lesions on the floor of the mouth and palate were rare [28]. Erosive was the most common form and was present in 58.3% of the patients which was predominant in females. This could be attributed to hormonal imbalance due to menopause as most of the women were between 45-60 years of age and use of allergic dentifrice and application of clove oil for relief of burning sensation [37]. These findings were inconsistent with the previous studies in which the reticular type of OLP was most common among females [27, 28].

The association of pigmentation of the oral mucosa was a prominent feature of reticular form of OLP [3, 16]. It was noted to be 6.7%. This could be attributed to various factors such as race, skin type and habits such as chewing tobacco, smoking [16]. The pigmentation was diffuse or in patches which ranged from brown to black in colour and was especially seen in the buccal mucosa. This was similar to other Indian studies (S, Anandan and Prasanthi, 2013; Hartanto and Kallarakal, 2017; Institute and National Cancer Institute, 2020).

The majority of the patients (60%) complained of some degree of oral discomfort in the form of burning sensation as reported in other studies [3]. Nearly 75% of the erosive lichen planus was treated using systemic steroids like Prednisone for a maximum period of eight months. During the later follow up, it was noted that the patients responded well with systemic steroid therapy when compared to topical steroid therapy. This could be because of the recalcitrant nature of OLP to topical steroids therapy [1, 8, 20, 21, 25].

Even though there is no specific treatment for OLP, symptomatic treatment is indicated [8, 21]. Corticosteroids provide relief and are the first drug of choice [1, 38]. Reticular type has better response to steroids when compared to erosive form [3, 16]. This can be related to the chronicity and refractory course of erosive lichen planus (Romero et al. 2016). The Spontaneous remission is seen in 40% of oral lichen planus [16].

To overcome this remission, use of ultraviolet A (PUVA) and laser can be used as an alternative therapy [29]. Small and accessibility solutions can be treated by the use of adherent paste in the form of a custom tray which allows accurate control over the contact time it ensures that the entire regional surface is exposed to the drug [9-11]. Local drug therapy can provide a more targeted and efficient drug delivery option than systemic delivery for the disease of oral mucosa [11, 33]. However potential for novel drug delivery systems in dentistry has not yet been fully developed and further research is still needed to improve the treatment outcomes [9-11].

Conclusion

OLP accounts for nearly 28.4% of the OPMD reporting to Saveetha dental College and Hospital with erosive type of lichen planus being the largely reported type of OLP which has an increased rate for malignant transformation. Hence it is necessary to follow up the OLP patients regularly and to provide a precise treatment which prevents the remission of the disease in these patients.

References

- [1]. Alerraqi E. Steroid Therapy in Oral Lichen Planus. *J Steroids HormSci* [Internet]. 2016;7(2):10-3.
- [2]. Srinivas K, Aravinda K, Ratnakar P, Nigam N, Gupta S. Oral lichen planus-Review on etiopathogenesis. *National journal of maxillofacial surgery*. 2011 Jan;2(1):15.
- [3]. Bakhtiari S, Taheri JB, Toossi P, Azimi S, KawosiNezhad S. Prevalence of oral lichen planus in Iranian children and adolescents: a 12-year retrospective study. *Eur Arch Paediatr Dent*. 2017 Dec;18(6):419-422. Pubmed PMID: 29139037.
- [4]. Ben Slama L. Pathologies médicalespotentiellementmalignes de la muqueusebuccale [Potentially malignant disorders of the oral mucosa]. *Rev Prat*. 2019 Oct;69(8):856-860. Pubmed PMID: 32237647.
- [5]. Crincoli V, Di Bisceglie MB, Scivetti M, Lucchese A, Tecco S, Festa F. Oral lichen planus: update on etiopathogenesis, diagnosis and treatment. *ImmunopharmacolImmunotoxicol*. 2011 Mar;33(1):11-20. Pubmed PMID: 20604639.
- [6]. Dissemmond J. Oral lichen planus: an overview. *J Dermatolog Treat*. 2004 Jun;15(3):136-40. Pubmed PMID: 15204144.
- [7]. Farhi D, Dupin N. Pathophysiology, etiologic factors, and clinical management of oral lichen planus, part I: facts and controversies. *ClinDermatol*. 2010 Jan-Feb;28(1):100-8. Pubmed PMID: 20082959.
- [8]. Ferguson MM. Treatment of erosive lichen planus of the oral mucosa with depot steroids. *Lancet*. 1977 Oct 8;2(8041):771-2. Pubmed PMID: 71587.
- [9]. Ganesha R,Hadi P. 'Management of Oral Lichen Planus Due to Stress', Proceedings of the 7th International Meeting and the 4th Joint Scientific Meeting in Dentistry. 2017.
- [10]. Gheena S, Ezhilarasan D. Syringic acid triggers reactive oxygen species-mediated cytotoxicity in HepG2 cells. *Hum ExpToxicol*. 2019 Jun;38(6):694-702. Pubmed PMID: 30924378.
- [11]. Gonzalez-Moles MA, Ruiz-Avila I, Rodriguez-Archilla A, Morales-Garcia P, Mesa-Aguado F, Bascones-Martinez A, et al. Treatment of severe erosive gingival lesions by topical application of clobetasol propionate in custom trays. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod*. 2003 Jun;95(6):688-92. Pubmed PMID: 12789149.
- [12]. González-Moles MÁ, Warnakulasuriya S, González-Ruiz I, González-Ruiz L, Ayén Á, Lenouvel D, et al. Worldwide prevalence of oral lichen planus: A systematic review and meta-analysis. *Oral Dis*. 2020 Mar 7. Pubmed PMID: 32144836.
- [13]. Gupta V, Ramani P. Histologic and immunohistochemical evaluation of mirror image biopsies in oral squamous cell carcinoma. *J Oral BiolCraniofac Res*. 2016 Sep-Dec;6(3):194-197. Pubmed PMID: 27761383.
- [14]. Hannah R, Ramani P, Sherlin HJ, Ranjith G, Ramasubramanian A, Jayaraj G, Don KR, Archana S. Awareness about the use, ethics and scope of dental photography among undergraduate dental students dentist behind the lens. *Research Journal of Pharmacy and Technology*. 2018 Mar 1;11(3):1012-6.
- [15]. Hartanto FK, Kallarakal TG. Pigmented oral lichen planus: A case report. *Scientific Dental Journal*. 2017 Sep 28;1(1):11-6.
- [16]. Hasan S, Mansoori S, Ansari MI, Siddiqui S. Oral lichen planus in an 8-year-old child: A case report with a brief literature review. *Journal of oral and maxillofacial pathology: JOMFP*. 2020 Feb;24(Suppl 1):S128.
- [17]. Hema Shree K, Ramani P, Sherlin H, Sukumaran G, Jayaraj G, Don KR, et al. Saliva as a Diagnostic Tool in Oral Squamous Cell Carcinoma - a Systematic Review with Meta Analysis. *PatholOncol Res*. 2019 Apr;25(2):447-453. Pubmed PMID: 30712193.
- [18]. Institute, N. C. and National Cancer Institute (2020) 'Oral Lichen Planus', Definitions. doi: 10.32388/hup02c.
- [19]. Ismail SB, Kumar SK, Zain RB. Oral lichen planus and lichenoid reactions: etiopathogenesis, diagnosis, management and malignant transformation. *Journal of oral science*. 2007;49(2):89-106.
- [20]. Jangid K, Alexander AJ, Jayakumar ND, Varghese S, Ramani P. Ankyloglossia with cleft lip: A rare case report. *J Indian SocPeriodontol*. 2015 Nov-Dec;19(6):690-3. Pubmed PMID: 26941523.
- [21]. Jayaraj G, Sherlin HJ, Ramani P, Premkumar P, Anuja N. Cytomegalovirus and Mucoepidermoid carcinoma: A possible causal relationship? A pilot study. *J Oral MaxillofacPathol*. 2015 Sep-Dec;19(3):319-24. Pubmed PMID: 26980959.
- [22]. Jayaraj G, Ramani P, Sherlin HJ, Premkumar P, Anuja N. Inter-observer agreement in grading oral epithelial dysplasia-A systematic review. *Journal of Oral and Maxillofacial Surgery, Medicine, and Pathology*. 2015 Jan 1;27(1):112-6.
- [23]. Jayaraj G, Sherlin HJ, Ramani P, Premkumar P, Natesan A. Stromal myofibroblasts in oral squamous cell carcinoma and potentially malignant disorders. *Indian J Cancer*. 2015 Jan-Mar;52(1):87-92. Pubmed PMID: 26837985.
- [24]. Doshi B, Khopkar U. Histopathology of lichen planus and its variants. *Lichen Planus*. New Delhi: Jaypee Brothers Medical Publisher (P) LTD. 2013:123-47.
- [25]. Kurt MH, Kolsuz ME, Eren H. Corticosteroid injection in treatment of persistent oral lichen planus: Three cases. *DermatolTher*. 2019 Sep;32(5):e13015. Pubmed PMID: 31268212.
- [26]. Li C, Tang X, Zheng X, Ge S, Wen H, Lin X, Chen Z, Lu L. Global Prevalence and Incidence Estimates of Oral Lichen Planus: A Systematic Review and Meta-analysis. *JAMA Dermatol*. 2020 Feb 1;156(2):172-181. Pubmed PMID: 31895418.
- [27]. de Lima SL, de Arruda JA, Abreu LG, Mesquita RA, Ribeiro-Rotta RF, Mendonça EF, Arantes DA, Batista AC. Clinicopathologic data of individuals with oral lichen planus: A Brazilian case series. *J ClinExp Dent*. 2019 Dec 1;11(12):e1109-e1119. Pubmed PMID: 31824590.

- [28]. Omal P, Jacob V, Prathap A, Thomas NG. Prevalence of oral, skin, and oral and skin lesions of lichen planus in patients visiting a dental school in southern India. *Indian J Dermatol*. 2012 Mar;57(2):107-9. Pubmed PMID: 22615505.
- [29]. Pavlic V, Vujic-Aleksic V. Phototherapy approaches in treatment of oral lichen planus. *PhotodermatolPhotoimmunolPhotomed*. 2014 Feb;30(1):15-24. Pubmed PMID: 24118508.
- [30]. Payeras MR, Cherubini K, Figueiredo MA, Salum FG. Oral lichen planus: focus on etiopathogenesis. *Arch Oral Biol*. 2013 Sep;58(9):1057-69. Pubmed PMID: 23660124.
- [31]. Rebora A. Etiopathogenesis of lichen planus: The contribution of epidemiology. *Journal of Dermatological Science*. 1991 May 1;2(3):194.
- [32]. RAMAMURTHY J, GURUNATHAN D. Association Between Smoking and Oral Lichen Planus in Males-A Retrospective study. *Journal of Contemporary Issues in Business and Government*. 2021 Feb 7;27(2):623-32.
- [33]. Kumar A, Sherlin HJ, Ramani P, Natesan A, Premkumar P. Expression of CD 68, CD 45 and human leukocyte antigen-DR in central and peripheral giant cell granuloma, giant cell tumor of long bones, and tuberculous granuloma: An immunohistochemical study. *Indian J Dent Res*. 2015 May-Jun;26(3):295-303. Pubmed PMID: 26275199.
- [34]. Sivaramakrishnan SM, Ramani P. Study on the Prevalence of Eruption Status of Third Molars in South Indian Population. *Biology and Medicine*. 2015 Oct 1;7(4):1.
- [35]. Sridharan G, Ramani P, Patankar S, Vijayaraghavan R. Evaluation of salivary metabolomics in oral leukoplakia and oral squamous cell carcinoma. *J Oral Pathol Med*. 2019 Apr;48(4):299-306. Pubmed PMID: 30714209.
- [36]. Sridharan G, Ramani P, Patankar S. Serum metabolomics in oral leukoplakia and oral squamous cell carcinoma. *J Cancer Res Ther*. 2017 Jul-Sep;13(3):556-561. Pubmed PMID: 28862226.
- [37]. Srivastava R, Sharma L, Pradhan D, Jyoti B, Singh O. Prevalence of oral premalignant lesions and conditions among the population of Kanpur City, India: A cross-sectional study. *J Family Med Prim Care*. 2020 Feb 28;9(2):1080-1085. Pubmed PMID: 32318471.
- [38]. Swathy S, Gheena S, Sri VL. Prevalence of pulp stones in patients with history of cardiac diseases. *Research Journal of Pharmacy and Technology*. 2015 Dec 1;8(12):1625.
- [39]. Thangaraj SV, Shyamsundar V, Krishnamurthy A, Ramani P, Ganesan K, Muthuswami M, Ramshankar V. Molecular Portrait of Oral Tongue Squamous Cell Carcinoma Shown by Integrative Meta-Analysis of Expression Profiles with Validations. *PLoS One*. 2016 Jun 9;11(6):e0156582. Pubmed PMID: 27280700.
- [40]. Viveka TS, Shyamsundar V, Krishnamurthy A, Ramani P, Ramshankar V. p53 Expression Helps Identify High Risk Oral Tongue Pre-malignant Lesions and Correlates with Patterns of Invasive Tumour Front and Tumour Depth in Oral Tongue Squamous Cell Carcinoma Cases. *Asian Pac J Cancer Prev*. 2016;17(1):189-95. Pubmed PMID: 26838208.