

THE CLINICAL PROFILE OF ORAL LICHEN PLANUS

Syed Murad Ali Shah¹, Muhammad Ilyas¹, Jawad Ahmad Kundi¹

1. Sardar Begum Dental College

ABSTRACT

OBJECTIVES

The objective of the study was to determine the clinical profile of oral lichen planus.

METHODOLOGY

This retrospective cohort study was carried from January 2011 to December 2015 at the Department of Oral and Maxillofacial Surgery, Sardar Begum Dental College, Peshawar. A total of 36 with 14 male and 22 female patients having oral lichen planus from were selected. On defined and population-based sample the age selected was ≥ 20 years and divided into four categories i.e., 20-29 years, 30-39 years, 40-49 years and 50-59 years. The diagnostic criteria proposed by van der Meij et al²³ in 2003 based on the WHO definition of oral lichen planus were used to identify the cases of oral lichen planus. That entire patient's with incomplete records and aged > 60 years were excluded. The data was analyzed through SPSS 22 at the significance level of $p < 0.05$ and Chi-square statistics was applied for site and gender association.

RESULTS

The mean age presentation was 39.2 (SD \pm 15.49) years. The female to male ratio was 1:57:1. The dominant aged group was 30-39 years with $n=16$ (44.44%). The buccal mucosa was the most common site involved $n=28$ (77.8%). Reticular type of oral lichen planus was the most common form and was present in $n=22$ (61.1%) patients however, bilaterally involved mucosa was commonly seen. Chi-square statistics showed a significant association between bilateral involvement of oral mucosa in oral lichen planus with both male and female ($\chi^2= 5.833$, $p= 0.016$).

CONCLUSION

The most common site involved in oral lichen planus was buccal mucosa, most common form was atrophic with female predominance and bilaterally involved oral mucosa was significantly associated with gender.

KEY WORDS

Oral Lichen Planus, Clinical Profile, Oral Mucosa.

INTRODUCTION

Oral lichen planus is chronic mucocutaneous disorder that affects oral mucosa, skin and

Correspondence:

Dr. Muhammad Ilyas
Sardar Begum Dental College
Contact: 0302-8809637
Email: ilyas_khan526@yahoo.com

<https://doi.org/10.37762/jgmids.3-01.45>

other mucous membrane and may be due to the immune response of CD8+ lymphocytes to antigen on lesional keratocytes^{1,2}. English physician Erasmus Wilson in 1866 described this condition to the world while Louis-Frédéric Wickham allocate Wickham Striae in

1895 to the interlacing white keratotic lines in the lesion^{3,4}. Oral lichen planus are characteristically raised multiform white lesions, accompanied by areas of erosions and pigmentation¹. Oral lichen planus has six clinical variants which may occur individually or in combination: papular, reticular, plaque-like, atrophic, erosive and bullous⁵. The reticular form has better prognosis as 40% of cases has spontaneous remission⁶, the erosive type being long standing with frequent exacerbations, severe pain and complications. Oral lesions are accompanied by skin lesions in approximately 50% of patients and may occur before, at the same time or after the skin lesions⁷. The lesions more commonly involved are bilateral buccal mucosa, mucobuccal fold, gingiva and less commonly tongue, palate and lips⁸. Reticular type is most

commonly involved in male as compared to female and usually asymptomatic however, erosive and atrophic are painful and causing burning sensation ⁹.

Approximately 20 % and 15 % of oral lichen planus is related to genital and cutaneous lichen planus respectively ^{10,11}. World Health Organization (WHO) classify oral lichen planus as potentially malignant disorder due to progression of oral lichen planus to the development of oral squamous cell carcinoma with the frequency of malignant transformation of 0.4-5.3% ^{12,13}. The research available from the developed countries well- described the demographic and clinical profile of oral lichen planus ¹⁴⁻¹⁸ however such sequence of research is rare from developing countries ¹⁹⁻²¹. The aim of the study was to determine the clinical profile of oral lichen planus.

METHODOLOGY

This retrospective cohort study was carried from January 2011 to December 2015 at the Department of Oral and Maxillofacial Surgery, Sardar Begum Dental College, Peshawar. Patients were identified by their medical records. A total of 36 patients with 14 male and 22 female patients having oral lichen planus from January 2011 to December 2015 were selected. On defined and population-based sample the age selected was ≥ 20 years and divided into four categories i.e., 20-29 years, 30-39 years, 40-49 years and 50-59 years. The diagnostic criteria proposed by van der Meij et al ²² in 2003 based on the WHO definition of oral lichen planus were used to identify the cases of oral lichen planus which include clinical as well as histopathological features but here only clinical features were included to diagnose oral lichen planus which was made through the clinical evaluation of patient's oral cavity by a specialist team at the Department of Oral & Maxillofacial Surgery. That entire patient's with incomplete records and aged > 60 years were excluded. The data was analyzed through SPSS 22 at the significance level of $p < 0.05$ and Chi-square statistics was applied for site and gender association.

RESULTS

The mean age presentation was 39.2 (SD \pm 15.49) years. The female to male ratio was 1.57:1. The dominant aged group was 30-39 years with $n=16$ (44.44%) followed by 40-49 years aged group, $n=12$ (33.33%). The least effective aged group is 20-29 years, $n=03$ (8.33%) as shown in the figure 1. The buccal mucosa was the most common site involved $n=28$ (77.8%). Gingiva and tongue were affected in $n=03$ (8.3%) and $n=05$ (13.9%) patients respectively. Reticular type of oral lichen planus was the most common form and was present in $n=22$ (61.1%) patients. Erosive form was observed in $n=12$ (33.3%) patients while atrophic oral lichen planus was seen in $n=02$ (5.6%) patients however, bilaterally involved mucosa was commonly seen as shown in the table.1. Chi-square statistics showed a significant association between bilateral involvement of oral mucosa in oral lichen planus with both male and female ($\chi^2= 5.833$, $p= 0.016$) as shown in table.2

Figure # 1: Age group distribution of Oral lichen planus

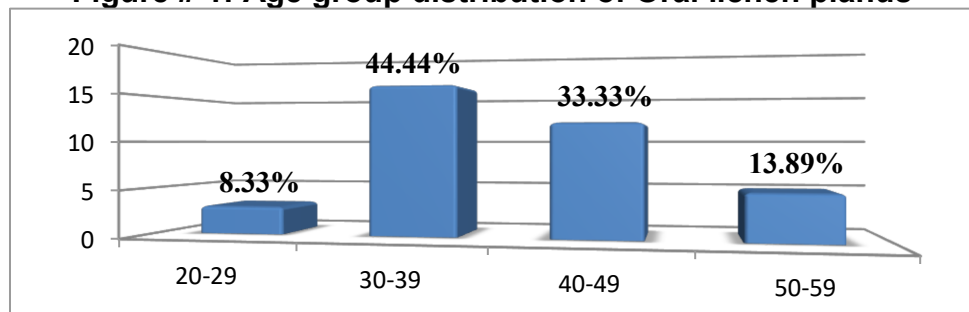


Table 1: Oral Lichen Planus involving sites and types

Bilateral n(%)	Unilateral n(%)	Reticular n(%)	Erosive n(%)	Atrophic n(%)	Buccal mucosa n(%)	Tongue n(%)	Gingiva n(%)
23(63.9%)	13(36.1%)	22(61.1%)	12(33.3%)	02(5.6%)	28(77.8%)	05(13.9%)	03(8.3%)

Table 2: Chi-Square statistics showed a significant association with site Involvement ($p < 0.05$)

	Unilateral	Bilateral	Mean	Standard Deviation	Chi-square Statistics	P value ($p < 0.05$)
Male	05	09	7	± 2.828	5.833	0.016
Female	08	14	11	± 4.242		

DISCUSSION

This retrospective study was done to determine the clinical characteristics of oral lichen planus patients in relatively small cohort from Peshawar, Pakistan. In this study, we observed that female outnumbered male with ratio of (F:M=1.57:1), which is in agreement with the report of Eisen D¹⁴ however Munde et al²³ in their retrospective study observed that male predominate female (M:F=1.61:1) which contradict our study. The bilaterally involved oral mucosa is more effected which support the study done by Ingafou M et al²⁴. According to two categories of clinical form classified by Gandolfo et al¹⁷ and Carbone et al¹⁸ the prevalence of reticular oral lichen planus in their series were 59.7% and 58.9% respectively likewise, the prevalence in our study which is 61.1% and is not in agreement with Munde et al²⁴. Our study also revealed that the most common site involved is buccal mucosa which support the study of Gandolfo et al¹⁷ and Carbone et al¹⁸. Oral lichen planus is more prevalent in the 4th decade of life in our study (mean age=39.4 years), which is lower than the mean age reported in central China (50.4 years)²¹, UK (52.0 years)²⁴, Spain (56.4 years)²⁵ and Italy (56.7 years)¹⁷ and support the study done by Munde et al²³.

A retrospective study has many restrictions and cannot be balanced decently with prospective study but they are applicable in assessing patient populations. The clinical characteristics of oral lichen planus show consistency in most of the results with the previous studies while few are not in agreement with our study. The lack of uniformity may be due to the different geographic areas. Oral lichen planus is a chronic disease and it is mandatory for Oral health care professionals to thoroughly examine the patient and evaluate the status of the patient accordingly with long term follow up which is obligatory for oral lichen planus patients.

CONCLUSION

The most common site involved in oral lichen planus was buccal mucosa, most common form was atrophic with female predominance and bilaterally involved oral mucosa was significantly associated with gender.

REFERENCES

1. Sugerman PB, Savage NW, Walsh LJ, Zhao ZZ, Zhou XJ, Khan A, et al. The pathogenesis of oral lichen planus. *Crit Rev Oral Biol Med* 2002;13:350-65.
2. Lodi G, Scully C, Carrozzo M, Griffiths M, Sugerman PB, Thongprasom K. Current controversies in oral lichen planus: Report of an international consensus meeting. Part 1. Viral infections and etiopathogenesis. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 2005;100:40-51.

3. Scully C, el-Kom M. Lichen planus: Review and update on pathogenesis. *J Oral Pathol* 1985;14:431-58.
4. Steffen C, Dupree ML. Louis-Frédéric Wickham and the Wickham's striae of lichen planus. *Skinmed* 2004;3:287-9.
5. Scully C, Carrozzo M. Oral mucosal disease: Lichen planus. *Br J Oral Maxillofac Surg.* 2008; 46(1): 15-21.
6. Andreason JO. Oral lichen planus: A clinical evaluation of 115 cases. *Oral Surg Oral Med Oral Pathol.* 1968; 25: 31-42.
7. Soames JV, Southam JC. Dermatological causes of white patches: Lichen planus. J.V Soames and JC Southam's *Textbook of Oral Pathology.* Oxford University Press Pakistan. 2005; 4th ed.:128-130.
8. Bermejo-Fenoll A, Sánchez-Siles M, López-Jornet P, Camacho-Alonso F, Salazar-Sánchez N. A retrospective clinicopathological study of 550 patients with oral lichen planus in south-eastern Spain. *J Oral Pathol Med.* 2010;39:491-6.
9. Chainani-Wu N, Silverman S, Jr, Lozada-Nur F, Mayer P, Watson JJ. Oral lichen planus: Patient profile, disease progression and treatment responses. *J Am Dent Assoc.* 2001;132:901-9.
10. Farhi D, Dupin N. Pathophysiology, etiologic factors, and clinical management of oral lichen planus, part I: Facts and controversies. *Clin Dermatol.* 2010;28:100-8.
11. Edwards PC, Kelsch R. Oral lichen planus: Clinical presentation and management. *J Can Dent Assoc.* 2002;68:494-9.
12. Eisen D. The evaluation of cutaneous, genital, scalp, nail, esophageal, and ocular involvement in patients with oral lichen planus. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod.* 1999;88:431-6.
13. Shi P, Liu W, Zhou ZT, He QB, Jiang WW. Podoplanin and ABCG2: Malignant transformation risk markers for oral lichen planus. *Cancer Epidemiol Biomarkers Prev.* 2010;19:844-9.
14. Eisen D. The clinical features, malignant potential, and systemic associations of oral lichen planus: A study of 723 patients. *J Am Acad Dermatol.* 2002;46:207-14.
15. Andreason JO. Oral lichen planus. 1. A clinical evaluation of 115 cases. *Oral Surg Oral Med Oral Pathol.* 1968;25:31-42.
16. Silverman S, Jr, Gorsky M, Lozada-Nur F. A prospective follow-up study of 570 patients with oral lichen planus: Persistence, remission, and malignant association. *Oral Surg Oral Med Oral Pathol.* 1985;60:30-4.
17. Gandolfo S, Richiardi L, Carrozzo M, Brocchetto R, Carbone M, Pagano M, et al. Risk of oral squamous cell carcinoma in 402 patients with oral lichen planus: A follow-up study in an Italian population. *Oral Oncol.* 2004;40:77-83.
18. Carbone M, Arduino PG, Carrozzo M, Gandolfo S, Argiolas MR, Bertolusso G, et al. Course of oral lichen planus: A retrospective study of 808 northern Italian patients. *Oral Dis.* 2009;15:235-43.
19. Murti PR, Daftary DK, Bhonsle RB, Gupta PC, Mehta FS, Pindborg JJ. Malignant potential of oral lichen planus: Observations in 722 patients from India. *J Oral Pathol.* 1986;15:71-7.

20. Pakfetrat A, Javadzadeh-Bolouri A, Basir-Shabestari S, Falaki F. Oral Lichen Planus: A retrospective study of 420 Iranian patients. *Med Oral Patol Oral Cir Bucal*. 2009;14:E315–8.
21. Xue JL, Fan MW, Wang SZ, Chen XM, Li Y, Wang L. A clinical study of 674 patients with oral lichen planus in China. *J Oral Pathol Med*. 2005;34:467–72.
22. van der Meij EH, Schepman KP, van der Waal I. The possible premalignant character of oral lichen planus and oral lichenoid lesions: A prospective study. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod*. 2003; 96: 164-171..
23. Munde AD, Karle RR, Wankhede PK, Shaik SS, Kulkurni M. Demography and clinical profile of Oral Lichen Planus: A retrospective study. *Contemp Clin Dent*. 2013 Apr;4(2):181-5.
24. Ingafou M, Leao JC, Porter SR, Scully C. Oral Lichen planus: A retrospective study of 690 British patients. *Oral Dis*. 2006; 12: 463-468.
25. Bermejo-Fenoll A, Sa`nchez-Siles M, Lo`pez-Jornet P, Camacho-Alonso F, Salazar-Sa`nchez N. A retrospective clinicopathological study of 550 patients with oral lichen planus in south-eastern Spain. *J Oral Pathol Med*. 2010; 39: 491-496.



LICENSE: JGMDS publishes its articles under a Creative Commons Attribution Non-Commercial Share-Alike license (CC-BY-NC-SA 4.0).
COPYRIGHTS: Authors retain the rights without any restrictions to freely download, print, share and disseminate the article for any lawful purpose. It includes scholarly networks such as Research Gate, Google Scholar, LinkedIn, Academia.edu, Twitter, and other academic or professional networking sites.