# **Original Article**

# Clinical and histopathological spectrum of lichen planus

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## **Abstract**

**Objective** To determine histopathological correlation with clinical presentation of lichen planus (LP).

*Methods* From the patients attending the outpatient department of Dermatology of Dayanand Medical College & Hospital, Ludhiana, those who were clinically diagnosed as LP constituted the study material.

Results Oral LP was the most common clinical variant with 24 (40%) cases, out of which 18 (30%) were of reticular oral LP, 6 (10%) were of erosive oral lichen planus. 19 (31.7%) cases were of classical LP, 6 (10%) were of hypertrophic LP, 5 (8.3%) were of LP pigmentosus, 3 (5%) were of eruptive LP, 1 (1.7%) was of genital LP, 1 (1.7%) was of lichen planopilaris, 1 (1.7%) was of actinic LP with DLE overlap. Both oral LP and classical LP were the most common histopathological variants with both having 21 (35%) cases. Second most common histopathological variant was that of hypertrophic LP with 6 (10%) cases. Majority of patients were in the age group of 41-60 years. Our study had a female preponderance. Most of the cases sought medical help within 6-10 months of development of symptoms. Lower limbs were the most common site involved in cutaneous LP. Burning sensation was the main presenting complaint in the patients of oral LP. In cutaneous LP moderate to severe itching was the most common symptom. The commonest recognized histopathological features were saw-toothed rete ridges/ irregular acanthosis, vacuolar degeneration of basal layer and band like infiltrate in our study.

**Conclusion** Oral LP is the most common clinical variant followed by classical LP with histopathological findings adequately consistent with the clinical diagnosis.

#### Key words

Clinical features, histopathology, lichen planus.

### Introduction

Lichen planus (LP) is an idiopathic inflammatory skin disease affecting the skin and mucous membranes, often with a chronic course with relapses and periods of remission. Its prevalence is approximately 0.5% of the

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# population.1

The incidence varies between 0.22% and 1% of the adult population worldwide.<sup>2</sup> In contrast, oral LP seems to be more frequent with a reported incidence between 1% and 4% of the population.<sup>1</sup> LP is rare in children and commonly affects adults during their fourth to sixth decade. According to one study, LP represents 0.38% of all dermatology outpatients in India.<sup>3</sup>

Several hypotheses have been made regarding its etiology including genetic, infective, psychogenic and autoimmune factors.<sup>4</sup> Several studies have suggested a role for hepatitis C virus (HCV) in LP.<sup>5</sup> Both antigen-specific and nonspecific mechanisms are involved in initiation of the immune reaction.<sup>6</sup>

The classic clinical presentation of LP includes primary lesions consisting of firm, shiny, polygonal, 1-3 mm diameter papules with a red to violet colour. "Pruritic, Purple, Polygonal, Planar, Papules and Plaques" are the traditional 6 "P's" of LP.7 Annular lesions are especially common on the penis and rarely may be the predominant type of lesion present, later leading to atrophy.8 When the palms and soles are affected, the lesions tend to be firm and rough with a yellowish hue.9 Linear LP occurring in Blaschko's lines can also be observed. 10 Mucous membrane lesions are very common, occurring in 30-70% of cases. The buccal mucosa and tongue are most often involved. White streaks, often forming a lacework, on the buccal mucosa are highly characteristic. Oral LP carries a significant risk of malignant transformation. studies have reported malignant transformation rate between 0.4 and 1.5%.11 Nail involvement occurs in up to 10% of cases.12

On standard histopathology, LP is characterized by the presence of a band-like lymphohistiocytic infiltrate at the dermal-epidermal junction with hydropic degeneration of the epidermis. Resultant dyskeratosis is represented by the presence of necrotic keratinocytes (Civatte bodies or cytoid bodies), which are extruded into the papillary dermis. Subepidermal clefts (Max-Joseph spaces) may form as a consequence of interface inflammation. Irregular acanthosis may assume a saw-toothed appearance. Hyperkeratosis is also seen.<sup>13</sup>

### **Methods**

Sixty clinically diagnosed patients of LP attending the outpatient department of Dermatology of Dayanand Medical College & Hospital, Ludhiana, constituted the study material.

An informed consent was taken from the patients regarding the biopsy and participation in the study. The patients were subjected to detailed clinical examination and skin/mucosal biopsy was done which was subjected to detailed histopathological examination.

#### Results

Out of 60 patients of LP enrolled in the study, 24 (40%) were of oral LP, 19 (31.7%) were of classical LP (**Table1**).

Oral LP and classical LP constituted the most common histopathological variants (35%). Biopsy was inconclusive in 10 (16%) cases (**Table 2**).

Majority of patients i.e. 24 (40%) were in age group of 41-60 years. LP in our study showed a female preponderance with 38 (63.3%) patients enrolled in the study being females. Majority of the cases (36.7%) sought medical help within 6-10 months of development of symptoms. Only 5% cases waited for more than 15 months before seeking medical help. Maximum number of cases (45%) showed only cutaneous involvement while 43.3% cases showed only mucosal involvement (**Table 3**).

Oral cavity was the most common site involved with 28 (46.7%) cases showing oral involvement. Lower limbs were the second most common site involved in 27 (45%) cases. Burning sensation was reported in 48.3% cases. It was closely followed by pruritus in 45% cases.

**Table 1** Various clinical variants of lichen planus (n=60).

Clinical diagnosis	N (%)
Classical lichen planus	19 (31.7)
Reticular oral lichen planus	18 (30.0)
Erosive oral lichen planus	6 (10.0)
Hypertrophic lichen planus	6 (10.0)
Lichen planus pigmentosus	5 (8.3)
Eruptive lichen planus	3 (5.0)
Genital lichen planus	1 (1.7)
Lichen planopilaris	1 (1.7)
Actinic lichen planus with discoid lupus erythematosus overlap	1 (1.7)

**Table 2** Histopathological variants of lichen planus (n=60).

Histopathological diagnosis	$N\left(\%\right)$
Classical lichen planus	21 (35)
Reticular oral lichen planus	19 (31.7)
Hypertrophic lichen planus	6 (10.0)
Erosive oral lichen planus	2 (3.3)
Lichen Planopilaris	1 (1.7)
Lichen planus pigmentosus	1 (1.7)
Inconsistent	10 (16.7)

**Table 3** Sites of involvement (n=60).

Sites involved	N (%)
Cutaneous	27 (45.0)
Mucosal	26 (43.3)
Cutaneous + nail	4 (6.7)
Cutaneous + oral	2 (3.3)
Cutaneous + nail+ oral	1 (1.7)

**Table 4** Spectrum of signs and symptoms of lichen planus (n=60).

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Signs and symptoms	N (%)
Burning sensation	29 (48.3)
Pruritus	27 (45.0)
Pain	14 (23.3)
Ulcers/erosions	7 (11.7)
Postinflammatory hyperpigmentation	5 (8.3)
Cicatricial alopecia	1 (1.7)
Longitudinal ridging	6 (10.0)
Nail plate thinning	2 (3.3)
Onycholysis	1 (1.7)
Koebner's phenomenon	10 (16.7)
Wickham striae	21 (35)

23.3% of patients complained of pain. Wickham's striae were observed in 35% of cases while Koebner's phenomenon (**Figure 1**) was seen in 16.7% of cases.

**Table 5** Various histopathological features (n=60).

Clinical diagnosis	N (%)
Hyperkeratosis/ orthokeratosis	38 (63.3)
Wedge shaped hypergranulosis	37 (61.7)
Irregular acanthosis/sawtooth rete ridges	51 (85.0)
Vacuolar degeneration of basal layer	49 (81.7)
Band-like infiltrate	43 (71.7)
Melanophages in upper dermis	26 (43.3)
Civatte bodies	27 (45.0)
Max-Joseph spaces	10 (16.7)
Pigment incontinence	25 (41.7)
Follicular plugging	5 (8.3)
Perivascular infiltrate	17 (28.3)
Papillomatosis	1 (1.7)
Chronic inflammatory infiltrate	6 (10.0)
Atrophy	5 (8.3)
Erosion	2 (3.3)

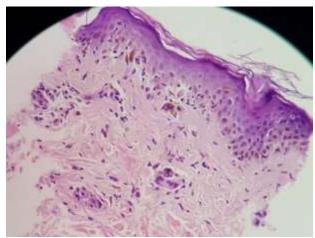
In nail changes, longitudinal ridging was the most common finding observed in 10% cases (**Table 4**).

Of all the cases, 16.7% patients complained of intense pruritus, 15% complained of moderate pruritus and only 13.3% experienced mild pruritus. 5 (8.3%) cases presented with only hyperpigmented violaceous papules and plaques without any associated symptoms. The commonest histopathological features recognized in our study were sawtooth rete ridges/ irregular acanthosis in 85%, vacuolar degeneration of basal layer in 81.7%, band-like infiltrate in 71.7%, hyperkeratosis/ orthokeratosis in 63.3% and wedge-shaped hypergranulosis in 61.7% of cases (Figure 2).

Classical LP was the most common histopathological diagnosis, followed by oral reticular LP. LP pigmentosus showed sawtoothed rete ridges/ irregular acanthosis, vacuolar degeneration of the basal layer, bandlike infiltrate, melanophages in upper dermis, Civatte bodies, pigment incontinence, chronic inflammatory infiltrate and perivascular infiltrate. Single case of lichen planopilaris enrolled in our study showed vacuolar



Figure 1 Wickham's striae and Koebner's phenomenon.



**Figure 2** Hyperkeratosis, irregular acanthosis and vacuolar degeneration of basal layer along with pigment incontinence.

**Table 6** Clinico-histopathological correlation of lichen planus (n=60).

	Histopathological diagnosis			Sensitivity to	
Subtype	Clinical diagnosis	Consistent with subtype	Inconsistent	Consistent with lichen planus but not subtype	dig lichen planus
Classical lichen planus	19	17	2-inconsistent	17-lichen planus	89.50%
Reticular oral lichen planus	18	17	1-inconsistent	17- oral lichen planus	94.40%
Erosive oral lichen planus	6	2	2-inconsistent	2-oral lichen planus	33.30%
Eruptive lichen planus	3	0	0	3-lichen planus	100.00%
Genital lichen planus	1	0	0	1-lichen planus	100.00%
Hypertrophic lichen planus	6	6	0	0	100.00%
Lichen planopilaris	1	1	0	0	100.00%
Actinic lichen planus with DLE overlap	1	0	1-inconsistent	0	0.00%
Lichen planus pigmentosus	5	0	4-inconsistent	1-lichen planus	20.00%

degeneration of basal layer, melanophages in upper dermis and perivascular infiltrate on histopathological examination. Ten cases did not show any features of lichen planus and were labelled as inconsistent (**Table 5**).

In the present study, 50 out of 60 clinically diagnosed cases were confirmed on histopathology and only 10 cases were found inconsistent hence giving a sensitivity of 83.3%. Histopathological examination showed a sensitivity of 71.6% in identifying the various subtypes of lichen planus (**Table 6**).

Regarding various associated comorbidities, hypertension was present in 18.3% cases, diabetes mellitus was seen in 6.7% cases, hepatitis C infection was present in 6.7% cases.

# **Discussion**

Lichen planus (LP) is an idiopathic inflammatory skin disease affecting the skin and mucous membranes, often with a chronic course with relapses and periods of remission. In our study, majority of patients i.e. 24 (40%) were in age group of 41-60 years, close second i.e. 20 (33.4%) patients were in the age group of 21-40

years. These findings are similar with other studies of the western population<sup>14,15</sup>; however, in the studies of Indian population by Bhattacharya et al.,3 Singh and Kanwar,16 a younger age group has been reported. However, in one Indian study by Arora et al. 17 an older age group of 31-60 years has been reported. Furthermore, in the western literature LP is considered to be rare in children.<sup>18</sup> Even in those studies, which are published from the European countries, a proportion of patients were Indians.<sup>19</sup> In our study, we found that female gender is more commonly affected with lichen planus than males. Even though in the literature there has been no consensus regarding any sex preference of LP, but most of the studies have shown that females are more commonly affected than males<sup>20,21,22</sup>; however, study by Kachhawa et  $al.^{23}$  have shown male preponderance.

In our study, the most common variant was oral LP seen in 40% of all enrolled cases. Reticular oral LP was the most common variant seen in 75% of total cases of oral LP followed by erosive oral LP seen in 25%. These findings are consistent with the study done by Munde et al.24 Classical LP (31.7%) was the second most common subtype overall and the most common subtype (52.7%) in cutaneous LP followed by hypertrophic LP (16.6%). These findings are consistent with another study by Abdallat et al.25 Oral cavity was the most common site involved in 46.7% of cases. In cutaneous LP lower limbs was the most commonly involved site seen in 75% of cases, which is in concordance with other studies done by Kachhawa et al.23 and Kanwar et al.26 Lichen planus pigmentosus showed a predilection for back involving 60% of cases. In another study by Parihar et al.27 it was observed that face and neck were the most frequent initial sites of involvement followed by the trunk. Similar findings have been reported in previous studies by Bhutani et al.28 and Kanwar et al.29 Only one case of lichen planopilaris was enrolled in our study and it had lesions on the scalp. In the study by Parihar *et al.*<sup>27</sup> also, the most common site involved was scalp, seen in 82% patients.

In the present study, burning sensation was the most common symptom in oral LP, although pruritus was the most common symptom in cutaneous LP as observed in the studies done by Parihar et al.27 and Arora et al.17 In the present study, Wickham's striae were observed in 35% of cases while Koebner's phenomenon was seen in 16.7% of cases. However, Koebner's phenomenon was observed in only 6% patients in the study done by Kanwar and De.19 In our study, postinflammatory hyperpigmentation was seen in 8.3% of cases, while it was seen in 15.9% of patients in a study done by Abdallat et al. 25 In our study, nail involvement in the form of longitudinal ridging was seen in 10% of cases similar to a study by Garg et al.30 On the contrary, Kanwar and De26 have observed nail involvement in 19% of their patients

In our study, the commonest histopathological features recognized were saw-toothed rete ridges/ irregular acanthosis seen in 85%, vacuolar degeneration of basal layer in 81.7%, band-like infiltrate in 71.7%, hyperkeratosis/ orthokeratosis in 63.3% and wedge-shaped hypergranulosis in 61.7% of cases followed by Civatte bodies in 45%, melanophages in upper dermis in 43.3%, pigment incontinence in 41.7%, perivascular infiltrate in 28.3% and Max-Joseph spaces in 16.7% of cases.

In a study done by Arora et al.<sup>17</sup> and Garg *et al.*,<sup>30</sup> epidermal changes included hypergranulosis (82%), hyperkeratosis (92%) and basal cell vacuolization (100%). The dermis showed band-like lymphocytic inflammatory cell infiltrate, predominantly perivascular in location.<sup>17,30</sup>

In another study done by Parihar *et al.*<sup>27</sup> all the cases showed orthokeratosis. Irregular acanthosis was seen in 94% cases. Pointed rete ridges and dome-shaped papillae were identified in 76% cases, wedge-shaped hypergranulosis in 96.5%. The infiltrate in the upper dermis was band-like in 94% of cases, Civatte bodies or necrotic keratinocytes were present in 82% of cases in the lower epidermis and especially in the papillary dermis. Pigment incontinence was seen in 99% of cases. Max-Joseph spaces were apparent in 29.5% of cases.

In our study, single case of lichen planopilaris enrolled showed vacuolar degeneration of basal layer, melanophages in upper dermis and perivascular infiltrate on histopathological examination. In the study done by Garg *et al.*<sup>30</sup> lichen planopilaris showed more marked keratotic plugging than classical LP and inflammatory infiltrate was predominantly perifollicular in nature.

LP pigmentosus showed saw-toothed rete ridges/irregular acanthosis, vacuolar degeneration of basal layer, band-like infiltrate, melanophages in upper dermis, Civatte bodies, erosion, pigment incontinence, chronic inflammatory infiltrate and perivascular infiltrate. In study by Parihar *et al.*<sup>30</sup> all the cases showed epidermal thinning and pigment incontinence, and 85% cases showed basal layer vacuolation.

In our study hypertension was present in 18.3% of cases and diabetes mellitus in 6.7% of cases. While in a study done by Kachhawa *et al.*,<sup>23</sup> hypertension was seen in 2.3% of cases and association with diabetes was also noted. Association with hypertension and diabetes was also noted by Bajaj *et al.*<sup>31</sup> Hepatitis C infection was present in 6.7% of cases enrolled in our study, while study done by Das *et al.*<sup>32</sup> showed associated HCV infection in 3.07% of cases

while no association was found in the study done by Prabhu *et al.*<sup>33</sup>

#### Conclusion

Lichen planus is a common entity encountered in our day-to-day practice. In the majority of cases diagnosis can be made clinically. However, in difficult cases diagnosis can be clinched by histopathological examination as it shows a high sensitivity. Histopathological examination also shows a high sensitivity to effectively identify the various subtypes of lichen planus.

It is important to identify the various subtypes of lichen planus as the prognosis, duration of treatment and advice to the patient varies for different subtypes.

#### References

- 1. Darier J. *Précis de Dermatologie*. Paris: Masson, 1909. p. 118.
- 2. Boyd AS, Neldner KH. Lichen planus. *J Am Acad Dermatol*. 1991;**25**:593-619.
- 3. Bhattacharya M, Kaur I, Kumar B. Lichen planus: a clinical and epidemiological study. *J Dermatol.* 2000;**27**:576-82.
- 4. Sugerman PB, Satterwhite K, Bigby M. Autocytotoxic T-cell clones in lichen planus. *Br J Dermatol.* 2000;**142**:449-56.
- 5. Deguchi M, Aiba S, Ohtani H, Nagura H, Tagami H. Comparison of the distribution and numbers of antigen-presenting cells among T lymphocyte-mediated dermatoses: CD1a+, factor XIIIa+, and CD68+ cells in eczematous dermatitis, psoriasis, lichen planus and graft-versus-host disease. *Arch Dermatol Res.* 2002;**294**:297-302.
- 6. 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.
- Lazar AJF, Murphy GF. The Skin. In: Kumar V, Abbas A, Fausto N, Aster JC. Robbins & Cotran Pathologic Basis of Disease, 8<sup>th</sup> edn. Philadelphia: Saunders; 2009. P. 1165-204.

- 8. Weedon D. The lichenoid tissue reaction. *Int J Dermatol.* 1982;**21**:203-6.
- 9. Gorouhi F, Davari P, Fazel N. Cutaneous and mucosal lichen planus: a comprehensive review of clinical subtypes, risk factors, diagnosis, and prognosis. *Sci World J.* 2014;**2014**:742826.
- 10. Schmidt H. Frequency, duration and localization of lichen planus. A study based on 181 patients. *Acta Derm Venereol*. 1961:**41**:164-7.
- 11. Shen ZY, Liu W, Zhu LK, Feng JQ, Tang GY, Zhou ZT. A retrospective clinicopathological study on oral lichen planus and malignant transformation: analysis of 518 cases. *Med Oral Pathol Oral Cir Bucal*. 2012;**17**:e943-7.
- 12. Daramola OO, Ogunbiyi AO, George AO. Evaluation of clinical types of cutaneous lichen planus in anti-hepatitis C virus seronegative and seropositive Nigerian patients. *Int J Dermatol.* 2003;**42**:933-5.
- 13. Lehman JS, Tolleffson MM, Gibson LE. Lichen Planus. *Int J Dermatol*. 2009;**48**:682-94.
- 14. Andreason J. Oral lichen planus. A clinical evaluation of 115 cases. *Oral Surg*. 1968:**25**:31-42.
- 15. Scully C, EI-Kom M. Lichen planus-review and update of pathogenesis. *J Oral Pathol*. 1985;**14**:431-58.
- 16. Singh OP, Kanwar AI. Lichen planus in India an appraisal of 441 cases. *Int J Dermatol*. 1976;**15**:752-6.
- 17. Arora KS, Chhabra S, Saikia UN, Dogra S, Minz RW. Lichen planus: A clinical and immuno-histological analysis. *Indian J Dermatol.* 2014;**59**:257-61
- 18. Mellgren L, Hersle K. Lichen planus—a clinical study with statistical methods. *Indian J Dermatol.* 1965;**11**:1.
- 19. Kanwar AJ, De D. Lichen planus in childhood: report of 100 cases. *Clin Exp Dermatol.* 2010;**35**:257-62.
- 20. White CJ. Lichen Planus: a critical analysis of 64 cases. *J Cutan Dis.* 1919;**37**:671-9.
- 21. Altman J, Perry HO. The variations and course of lichen planus. *Arch Dermatol*. 1961;**84**:179-91.

- 22. Little EG. Lichen Planus. *J Cutan Dis*. 1919:**37**:639
- 23. Kachhawa D, Kachhawa V, Kalla G, Gupta L. A clinico-aetiological profile of 375 cases of lichen planus. *Indian J Dermatol Venereol Leprol*. 1995;**61**:276-9.
- 24. Munde AD, Karle RR, Wankhade PK, Shaik SS, Kulkumi M. Demographic and clinical profile of oral lichen planus: A retrospective study. *Contemp Clin Dent*. 2013;4:181-5.
- 25. Abdallat SA, Maaita TJ. Epidemiological and clinical features of lichen planus in Jordanian patients. *Pak J Med Sci.* 2007;**23**:92-9.
- 26. Kanwar AJ, De D. Lichen planus in children. *Indian J Dermatol Venereol Leprol*. 2010;**76**:366-72.
- 27. Parihar A, Sharma S, Bhattacharya SN, Singh UR. A clinicopathological study of cutaneous lichen planus. *J Dermatol Surg*. 2015;**19**(1):21-6.
- 28. Bhutani LK, Bedi TR. Lichen planus pigmentosus. *Dermatologica*. 1974;**149**:43-5.
- 29. Kanwar AJ, Dogra S, Handa S, Parsad D, Radotra BD. A study of 124 Indian patients with lichen planus pigmentosus. *Clin Exp Dermatol*. 2003;**28**:481-5
- 30. Garg VK, Nangia A, Logani K, Sharma RC. Lichen Planus-a Clinico-histopathological. *Indian J Dermatol Venereol Leprol.* 2000;**66**:193-5.
- 31. Bajaj DR, Khoso NA, Devrajani BR, Matlani BL, Lohana P. Oral lichen planus: A clinical study. *J Coll Physicians Surg Pak.* 2010;**20**:154-7.
- 32. Das A, Das J, Majumdar G, Bhattacharya N, Neogi DK, Saha B. No association between seropositivity for hepatitis C virus and lichen planus: A case control study. *Indian J Dermatol Venereol Leprol*. 2006;**72**:198-200.
- 33. Prabhu S, Pavithran K, Sobhanadevi G. Lichen planus and hepatitis c virus (HCV) Is there an association? A serological study of 65 cases. *Indian J Dermatol Venereol Leprol.* 2002;**68**:273-4.