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**Dermatology** 

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# **Unilateral Blaschkoid Lichen Planus: A Case Series of Rare Entity**

Dr. Sukhmani Kaur Brar<sup>1</sup>, Dr. Jaskirat Kaur<sup>2\*</sup>

#### <sup>1</sup>Fortis Medcentre Chandigarh, India

<sup>2</sup>Post Graduate Resident, Department of Dermatology, Venereology and Leprology, Guru Gobind Singh Medical College and Hospital, Faridkot-151203, Punjab, India

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### \*Corresponding author: Dr. Jaskirat Kaur

Abstract	Case Report

Lichen planus is an idiopathic inflammatory dermatological condition affecting skin and mucosa. Blaschkoid lichen planus is a relatively linear pattern of the disease. We herein report case series of 3 patients with unilateral blaschkoid lichen planus along with their characteristic clinical and histopathological profile. Keywords: blaschkoid, lichen planus

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## **INTRODUCTION**

Lichen planus is an idiopathic inflammatory dermatological condition affecting skin and mucosa. Certain variants of lichen planus may present a difficulty in diagnosis especially when the lesions are arranged in linear pattern. We herein report case series of 3 patients with unilateral blaschkoid lichen planus.

## **CASE HISTORY**

## Case-1

A 25 year old male patient presented with chief complaint of hyperpigmented raised lesion over right axilla since 5 months. The lesions were pruritic but there was no pain or discharge. The lesions were gradually expanding. There was no history of trauma, fever, or any stress factor. No history of any drug intake could be elicited. On examination the lesions were, voilaceous to hyperpigmented macules involving T1 and T2 dermatome (Figure-1). Nails, mucosa and scalp examination was completely normal. Systemic examination was essentially normal. Dermoscopic examination of the lesion with Dermlite Dl4 showed wickhiam straie with multiple pigmeted globules in surrounding (Figure-2).

With the provisional diagnosis of progressive cribriform and zosteriform hyperpigmentation and blashkoid lichen planus, skin biopsy of the lesion was done. Histopathological examination of the lesion revealed basket weave hyperkeratosis and a cluster of colloid bodies. Prominent melanin incontinence along with features of interface dermatitis were seen (Figure3). These findings were consistent with findings of lichen planus. The patient was given topical potent corticosteroids and antihistaminics for 2 weeks, pigmentation slightly improved.



Fig-1: Voilaceous hyperpigmented macules can be seen in the axilla

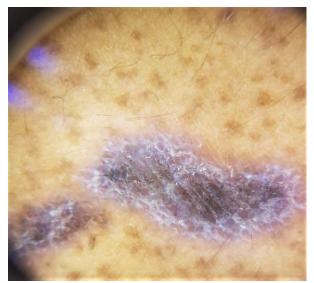


Fig-2: Dermoscopic examination shows wickhiam straie and multiple pigmented globules

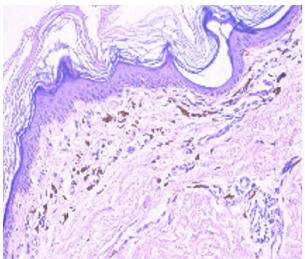


Fig-3: Histopathological examination shows prominent melanin incontinence and features of interface dermatitis

#### Case-2

A 34 year old female presented with pruritic rash over left upper limb, left lower limb and left side of trunk. There was no significant past history. On examination, the lesions were voilaceous flat topped 2-3 mm papules, along the blaschko's lines (Figure 4). The lesions did not crossed midline. Nail examination revealed onychoschizia, prominent longitudinal striations, thinning and melanonyhia (Figure-5). A preliminary diagnosis of lichen planus was made. Histopathological examination revealed parakeratosis, abuting the basal layer showing vacuolization, squamatization and occasional apoptotic keratinocytes. Lymphoplasmocytic infilterate was also seen in lamina propria (Figure-6). The features were consistent with the findings of lichen planus.

All the labortary investigations including G6PD were within normal limit. The patient was prescribed oral dapsone 100mg h.s. along with topical corticosteroid and oral antistaminics. The lesions

improved over 2 months, but the hyperpigmentation still remained



Fig-4: Blaschkoid lichen planus lesions on lef upper trunk



Fig-5: Nail examination revealed onychoschizia, prominent longitudinal striations, thinning and melanonyhia

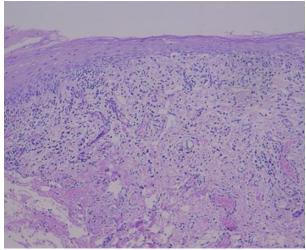


Fig-6: Histopathological examination revealed parakeratosis, abuting the basal layer showing vacuolization, squamatization and occasional apoptotic keratinocytes

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#### Case-3

A 28 year old male presented with multiple spiral purple coloured lesions over left upper limb, lower limb and left side of trunk (Figure 7a and b). The lesions were associated with itching. No history of trauma, any topical or oral medication was present. Nail examination was normal. Oral mucosa showed reticulate pattern of lichen planus. Serological investigations were within normal limit. Histopathological examination showed basket weave stratum corneum covering epidermis. Sparse perivascular lymphocytic infiltrate with numerous melanophages was seen (Figure-8). On the basis of clinical and histopathological examination, the diagnosis of blaschkoid lichen planus was made. The patient was given treatment in form of potent topical corticosteroids and antihistaminics. The response to treatment was favorable, with decrease in itching and improvement in the clinical picture.



Fig-7: The figure shows spiral hyperpigmented lesion on (a) left side of trunk and left upper limb and (b) left lower limb

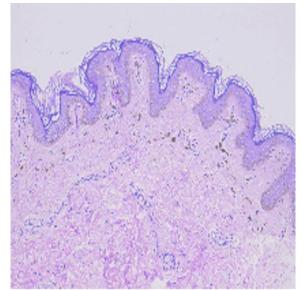


Fig-8: Histopathological examination shows sparse perivascular lymphocytic infilterate with numerous melanophages was seen

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## DISCUSSION

The classical presentation of lichen planus include purple, polygonal, pruritic papules and plaques [1]. Other morphological variants include hypertrophic, atrophic, actinic, guttate, erosive etc. Blaschkoid lines demonstrate a type of mosaicism, which determine the localization of many dermatological diseases. They are V –shaped over upper spine, S- shaped over abdomen and U-shaped over chest and upper arm. Blaschkoid lichen planus is one of the dermatosis in this pattern. It is a rare clinical entity with prevalence of 0.5 to 1 % in various studies [2]. The differential diagnosis for the condition include lichen striatus, progressive cribriform and zosteriform hypermelanosis, inflammatory linear verrucous epidermal nevus, linear psoriasis, linear and whorled nevoid hypermelanosis etc.

Apart from blashkoid lichen planus, linear lesions have also been reported in other variants of lichen planus like atrophic lichen planus [3], lichen planus follicularis [4] and lichen planus pigmentosus [5].

Lichen planus in dermatomal distribution which has been rarely reported in literature. Thus,

blaschoid lichen planus should be considered in the differentials of linear lesions in dermatology. Correct diagnosis and treatment of this condition are necessary to prevent any undue investigation and for cosmetic appearance.

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