

Dermoscopic Keys in Extragenital Bullous Hemorrhagic Lichen Sclerosus

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Introduction

Lichen sclerosus (LS) is a chronic inflammatory dermatosis characterized by leukoderma and scarring, predominantly affecting the genital skin. It may sometimes involve extragenital areas. The suffix atrophicans is no longer used as a few cases are associated with hypertrophy rather than atrophy. Few atypical variants include bullous, hemorrhagic, pigmented, verrucous, and keratotic forms. Herein, we report a case of extragenital bullous hemorrhagic lichen sclerosus with its dermoscopic features.

Case Presentation

A 40-year-old male presented with a nine-month history of a slow-growing asymptomatic raised lesion on his back associated with occasional bleeding episodes after casual trauma. There was no history of similar lesions in the past or elsewhere on the body. Physical examination revealed a solitary, well-defined, 2.5 cm × 2.0 cm, non-tender, hemorrhagic bulla with crusting in the center and atrophy in the surrounding area (Figure 1 A). Dermoscopy revealed superficial yellowish

white and hemorrhagic crusts, follicular plugs, and multi-colored diffuse hemorrhagic area with varying shades ranging from black to red, with black color representing old hemorrhage and red color representing recent hemorrhage (Figure 1B). Surrounding skin revealed atrophy with follicular plugs (Figure 1C). Based on the clinical and dermoscopic examination, we considered hemorrhagic lichen sclerosus, irritated seborrheic keratosis, Bowen disease, and discoid lupus erythematosus as our differential diagnoses. Histopathology revealed follicular plugs, epidermal atrophy, subepidermal blister, and hyalinized compact collagen, which confirmed the case to be LS (Figure 1D).

Discussion

The extragenital form of LS is less common, and the bullous hemorrhagic form is very rare, with only a handful of cases in the literature. This form is generally associated with less pruritus and the absence of any malignancy, as seen in our patient as well. In our case, the lesion was present on the back, a site that has not been reported for this particular variant. The formation of bullous lesions has been described

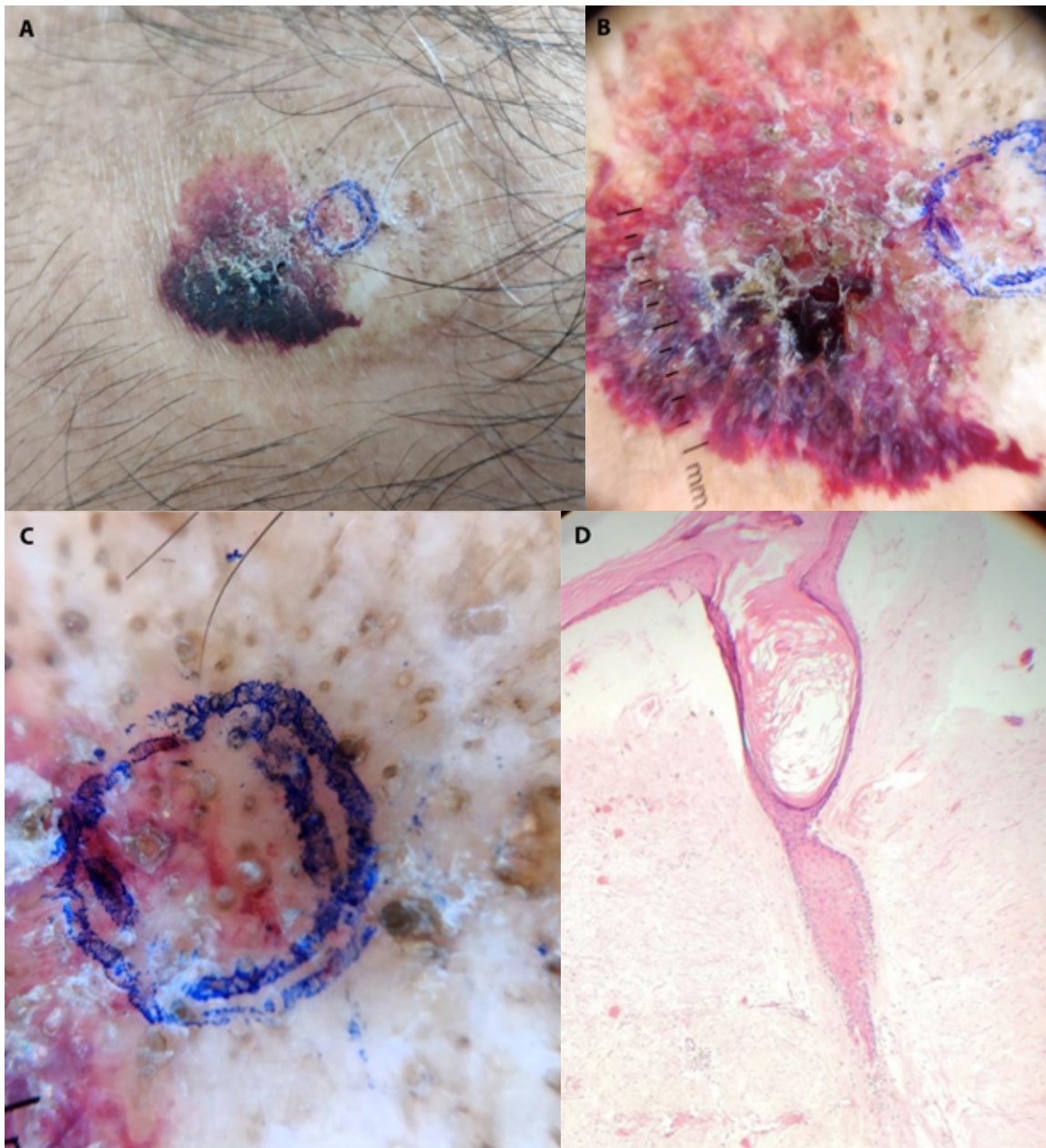


Figure 1. A. A solitary, well defined, 2.5 cm × 2.0 cm, non-tender, hemorrhagic bulla with crusting in the centre and atrophy in the surrounding area. B. Dermoscopic examination of lesion shows superficial yellowish white and hemorrhagic crusts, follicular plugs and multicolored diffuse hemorrhagic area with varying shades ranging from black to red (black color representing old hemorrhage and red color indicating recent hemorrhage). Blue color indicates marking for biopsy site. C. Dermoscopic examination of surrounding skin shows atrophy with follicular plugs. D. Skin biopsy showing presence of follicular plugs, epidermal atrophy, subepidermal blister and hyalinized compact collagen (H&E x 40).

in LS. A possible explanation for the formation of bulla and hemorrhage could be the pronounced edema within the skin that disrupts the capillaries collagen support, predisposing them to rupture with minimal trauma or damage [1].

Conclusions

Dermoscopy of extragenital LS has been described as white structureless areas, follicular plugs, white chrysalis-like structures, and variable vascular patterns being the essential

components [2]. Our case had superficial yellowish white and hemorrhagic crusts, a multicolored (black to red) hemorrhagic area, and a peripheral atrophic area with follicular plugs. There was no vascular pattern which commensurates with the chronicity of the lesion. The patient was managed with topical corticosteroids with a good response. This case report helps establish the fact that follicular plugs which have been reported in LS are seen in this rare variant also. In addition, the dermoscopic features of the hemorrhagic area of LS, which have not been previously described, have been

brought out. This report will enhance the existing repertoire of knowledge of dermoscopic features of LS which may aid diagnosis in future and avoid invasive procedures.

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