Glimpses of gold: Unveiling the enigma of multiple lichen aureus

Dear Editor,

A 53-year-old woman presented with a six-month history of multiple, mildly itchy, reddish-brown lesions on both legs. The lesions started appearing after four months of surgery for varicose veins and gradually increased in number over the next 4 months. On examination, there were multiple, well-defined, variably sized and shaped, reddish-brown papules and plaques on both legs ranging between 1 × 1 cm to the largest being 5 × 3 cm. The two most prominent plaques on her left leg showed a classic reddish-brown appearance surrounded by a yellowish-golden rim with the surface showing mild scaling forming a collarette at some places and punctate red to brown vascular structures [Figures 1a and 1b]. The surrounding skin showed signs of venous flares, such as dilated tortuous capillaries and veins on a generalised yellowish-to-brownish background. Dermoscopic examination revealed the presence of multiple discrete red dots and globules in a coppery brown background, serpentine vessels, reticular brown lines and subtle brown dots [Figure 2].

A skin biopsy performed on one of the lesions for histopathological examination revealed epidermal hyperkeratosis, focal patchy parakeratosis with focal basal cell vacuolisation and lymphocytic exocytosis [Figure 3]. The upper dermis showed moderate lymphocytic inflammatory infiltrate, predominantly in the perivascular region. The dermal vessels had endothelial swelling, and extravasated red blood cells (RBCs) were noted in the upper dermis. No evidence of vasculitis was identified.

Based on the clinical, dermatoscopic and histopathological findings, a diagnosis of multiple lichen aureus was made. Mid-potency topical steroid, once daily application, in combination with oral vitamin C supplementation and avoidance of prolonged standing, were suggested for treatment. The acute appearance of characteristic, rust-yellow or gold-coloured papules or circumscribed patches and plaques characterises lichen aureus. The name “lichen aureus” draws attention to the eruption’s traditional gold-brown hue. Notably, lesions that are copper-orange or violaceous may also be seen in

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lichen aureus. Unilateral lower extremity involvement is more common than bilateral involvement. The lesions can appear in the upper extremities and trunk and can occur in segmental, dermatomal, and blaschkoid patterns. The lesions can follow the course of superficial or deep veins. Aberrant distributions and patterns have been observed more frequently in children.

Multiple lichen aureus has been rarely reported. Torraca et al. described a case of multiple segmental lesions of lichen aureus that evolved over six years. One of the possible factors for the multiplicity of lesions in our patient can be the presence of venous insufficiency. In a previous study by Kim et al., venous insufficiency was detected in 35 of the 56 (62.5%) patients with pigmented purpuric dermatosis. Significant correlations were found between pigmented purpuric dermatosis and venous insufficiency. Longer duration of pigmented purpuric dermatosis, darker, and widespread lesions were seen in patients with established venous insufficiency.

Managing a case of widespread pigmented purpuric dermatosis in the presence of established venous insufficiency, when surgical management for venous insufficiency has already been tried without much improvement in lichen aureus, remains a challenge. Patients need counselling and an understanding of the interplay of various factors involved in the genesis of these skin lesions. Topical steroids, in combination with oral vitamin C supplementation, can be tried. Behavioural modifications, like intermittent sitting during hours of prolonged standing, elevation of limbs while sitting and interrupting the itch-scratch cycle, must be emphasised to the patients.

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