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Adult blaschkolinear acquired inflammatory skin eruption (BLAISE) with simultaneous features of lichen striatus and blaschkitis

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Abstract
Blaschkitis and lichen striatus are generally distinguished in the literature by the age of onset, lesion distribution, and histopathology. However, there is currently no clear consensus among authors about whether to consider blaschkitis and lichen striatus different clinical entities or a spectrum of the same disease. We present a case of adult BLAISE with features of both lichen striatus and blaschkitis, which seems to support the theory that these clinical entities may in fact represent a spectrum of the same pathological process.

Keywords: blaschkitis; lichen striatus; BLAISE; blaschkolinear skin eruption

Introduction
Blaschkolinear acquired inflammatory skin eruption (BLAISE) comprises a diversity of dermatologic entities characterized by the presence of inflammatory changes distributed along Blaschko lines [1]. There is currently no clear consensus among authors about whether to consider blaschkitis and lichen striatus different clinical entities or a spectrum of the same disease [2].

The most common clinical entities of BLAISE — blaschkitis and lichen striatus — are generally distinguished in literature by the age of onset, lesion distribution, and histopathology. Although blaschkitis is characterized by papules and vesicles generally distributed in a spiral arrangement along multiple ipsilateral blaschko lines of the trunk of adult patients, lichen striatus lesions generally present with lichenoid papules, which tend to follow a linear pattern along blaschko lines on the extremities of pediatric patients. Whereas the former clinical entity exhibits predominantly spongiotic changes on histopathology, lichen striatus displays lichenoid changes [3].

Case Synopsis
A 21-year-old man presented with unilateral pruritic cutaneous lesions that had appeared 1 week before. There was no history of chronic disease or medication. Dermatologic examination (Figure 1, 2) revealed erythematous papules and vesicles distributed on the left side of the abdomen and upper left limb, grouped along these locations in a whorl and streak
configuration, respectively. A punch biopsy revealed hyperkeratosis, acanthosis, spongiosis, and lichenoid infiltrates in the dermis (Figure 3). The patient was treated with betamethasone cream with complete resolution of the cutaneous lesions in 3 weeks.

Discussion

In both blaschkitis and lichen striatus an external acquired stimulus (e.g. infection, trauma) or autoimmune phenomenon (e.g. triggered by somatic mutations in keratinocytes) may lead to T-cell mediated inflammatory reaction directed against human keratinocyte mosaics present along Blaschko lines [4, 5]. Although the same theoretical triggers are reported, the reason why the cutaneous reaction pattern has such differing clinical and histological characteristics in pediatric and adult patients remains unsolved [1, 6].

Conclusion

We describe a man with a clinical presentation and histopathology with overlap features consistent with both blaschkitis and lichen striatus. Our case seems to support the opinion of some authors who suggest that these clinical entities may in fact represent a spectrum of the same pathological process.

References


Figure 2. Clinical aspect of the lesions: erythematous papules and vesicles distributed on the upper left limb, forming a linear configuration.

Figure 3. A) Hyperkeratosis, acanthosis, spongiosis, and lichenoid infiltrates in the dermis. H&E, 10%. B) Spongiosis, lichenoid infiltrates in the dermis, and apoptotic keratinocytes. H&E, 40%.