Placa Anular Única na Fronte

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PALAVRAS-CHAVE – Doenças da Pele; Granuloma; Testa

Single Annular Plaque on the Forehead

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QUAL O SEU DIAGNÓSTICO?

A 51-year-old caucasian female presented with an 18-month history of an asymptomatic well demarcated annular plaque with central atrophy and raised erythematous margins on her forehead (Fig. 1). The patient had no associated systemic manifestations and denied any preceding local trauma or exposure to previous medication. There was no relevant personal or family history. Laboratory studies were unremarkable. Histopathologic examination showed a granulomatous infiltrate with numerous multinucleated giant cells surrounding solar elastosis (Fig. 2A). Verhoeff-van Gieson stain revealed elastophagocytosis (Fig. 2B). No infectious agents were identified after staining with Ziehl-Neelsen, Grocott and periodic acid-Schiff (PAS).

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Figure 1 - Annular plaque on the forehead with an atrophic and hypopigmented center and raised erythematous margins.
**Qual o Seu Diagnóstico?**

**DIAGNOSIS**

**ACTINIC GRANULOMA**

The clinical and histopathological findings supported a diagnosis of actinic granuloma. O’Brien’s actinic granuloma, or annular elastolytic giant-cell granuloma, is an uncommon form of granulomatous dermatosis appearing on sun-exposed areas. Although the pathogenesis is unknown, solar elastosis resulting from sun damage is believed to be a triggering factor. The postulated mechanism behind this hypothesis is that solar-damaged elastic fibers are weakly antigenic and result in a cell-mediated immune response with a predominance of CD4 lymphocytes trying to repair the actinically damaged skin but resulting in granulomatous inflammation instead.¹ ²

The usual characteristics of actinic granuloma include asymptomatic annular or polycyclic lesions with centrifugal expansion, an erythematous elevated border and a hypopigmented atrophic center.³ Elastophagocytosis is the typical histopathologic feature, usually associated with an infiltrate of multinucleated giant cells that phagocytize the degenerated elastic fibers.⁴

Clinical differentials include facial lesions of granuloma annulare, necrobiosis lipoidica and annular sarcoid, but they can be readily distinguished by histopathology.¹ Treatment of actinic granuloma is difficult and lesions tend to persist with little response to therapy.² Several treatments have been proposed with varying results: topical or intralesional corticosteroids, acitretin, isotretinoin, cyclosporin, chloroquine, pentoxifylline, cryotherapy, PUVA phototherapy and methotrexate.² ⁵ The disease presents a chronic evolution, however some patients experience spontaneous resolution.¹ ² ⁴

In the present clinical case there was no significant improvement with topical betamethasone. Assured of the benign nature of the disease, the patient refused other therapies, despite being informed of a potential enlargement and/or appearance of new lesions. The patient was advised to reduce sun exposure and use sunscreen to prevent development of new lesions. The lesion has remained stable for the last 10 months and will be kept under clinical surveillance.

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