## **QUIZ SECTION**

## **Three Cephalic Plaques: A Quiz**

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A 74-year-old male presented with 3 infiltrated angiomatous plaques on his vertex and forehead for 6 months (Fig. 1), which were asymptomatic. He had a medical history of prostate cancer in 2016. He had controlled plaque psoriasis and gout, a drug rash following allopurinol introduction, but no other notable history. Clinically, the patient had neither other similar plaques nor adenopathy. Biopsy showed a neutrophilic and eosinophilic infiltration in the superficial

Fig. 1. (A) Two angiomatous plaques of the vertex. (B) Lesion of the forehead.

and mid-dermis, a Grenz zone, rich vascularization and multiple capillaries, a minimal perivascular fibrosis and an intact epidermis (Fig. 2). Laboratory tests revealed no eosinophilia or any other anomalies.

What is vour diagnosis? See next page for answer.

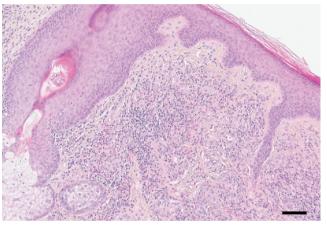


Fig. 2. Mixed dermal infiltrate formed by lymphocytes, histiocytes, plasma cells and neutrophils and a grenz zone (haematoxylin and eosin, original magnification ×400). Black bar; 100 µm.

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# **ANSWERS TO QUIZ**

### **Three Cephalic Plaques: A Commentary**

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### Diagnosis: Extrafacial granuloma faciale

Extrafacial granuloma faciale (GF) is a limited cutaneous disease of unknown aetiology (1). Middle-aged lightskinned men are commonly affected. GF is expressed by the presence of 1 or multiple plaque/nodule lesions of redbrown to violaceous colour with follicular accentuation and telangiectasia. Isolated facial presentation is commonly reported in the literature, but some patients also have associated extrafacial lesions. Approximately one-third of patients (31%) had isolated extrafacial lesions in a reported series of 32 patients (2). The trunk is usually the most frequently affected zone, followed by the upper extremities and the scalp (2). The pathophysiology of GF remains unknown, but it is probably related to sun exposure, as the lesions are generally found in photo-exposed areas. As diagnosis is based on histopathology, any well-demarcated infiltrated plaque should be biopsied (3). GF is frequently resistant to conventional therapy. Topical and injectable corticosteroids or tacrolimus and dapsone have been proposed, and combination with cryotherapy can also be used to obtain a better sustainable therapeutic response (2-4).

#### ACKNOWLEDGEMENTS

*Ethics approval:* The patient provided written informed consent for publication of his case details.

#### REFERENCES

- 1. Deen J, Moloney TP, Muir J. Extrafacial granuloma faciale: a case report and brief review. Case Rep Dermatol 2017; 9: 79–85.
- Gupta L, Naik H, Kumar NM, Kar HK. Granuloma faciale with extrafacial involvement and response to tacrolimus. J Cutan Aesthet Surg 2012; 5: 150–152.
- De D, Kanwar AJ, Radotra BD, Gupta S. Extrafacial granuloma faciale: report of a case. J Eur Acad Dermatol Venereol 2007; 21: 1284–1286.
- Mookadam M, Mesinkovska N, Bridges AG. Evaluating the clinical and demographic features of extrafacial granuloma faciale. Cutis 2017; 100: E18–E22.