

## **Clinical vignette**

# Calcinosis cutis: need for early and aggressive treatment

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A 57-year-old woman with oesophageal dysmotility and complaint of 'crystals' emerging from wounds presented to the clinic. She brought crystals (Fig. 1) that had grown and fallen from the wounds. Examination was notable for sclerodactyly distal to the wrists, in addition to ulcerated, painful nodules extruding a whitish-yellow chalk-like matter on her bilateral shins (Fig. 1) and the ulnar border of the left forearm. Blood work revealed ANA 1:640 (centromere pattern), positive anticentromere antibody and negative anti-RNA polymerase III, anti-Scl-70, anti-Smith, anti-dsDNA, anti-SSA, anti-SSB and anti-RNP. Serum calcium, phosphorus, vitamin D and parathyroid hormone were normal. A diagnosis of limited SSc with advanced calcinosis cutis was made. Treatment was initiated with MMF 1.5g twice daily, diltiazem 120 mg twice daily and colchicine 0.6 mg daily. After 6 months without im-



Figure 1. Calcinosis cutis of left shin with patient's collection of calcium crystals that had fallen out

provement of the calcinosis, the colchicine was stopped, and minocycline was started at 100 mg daily with follow-up in 6 months. Treatment for calcinosis cutis remains challenging, especially in advanced disease. Intralesional sodium thiosulphate, IVIG, minocycline, colchicine, CYC and rituximab have shown some promise; however, there is a dearth of large evidence-based studies [1]. Surgical excision can provide cosmetic benefits, but it does not prevent recurrence [2]. Treatment of the underlying disease early and aggressively to prevent morbidity is the current consensus.

#### Data availability statement

Data are available upon reasonable request by any qualified researchers who engage in rigorous, independent scientific research, and will be provided following review and approval of a research proposal and Statistical Analysis Plan (SAP) and execution of a Data Sharing Agreement (DSA). All data relevant to the study are included in the article.

#### Funding

No specific funding was received from any bodies in the public, commercial or not-for-profit sectors to carry out the work described in this article.

Disclosure statement: The authors have declared no conflicts of interest.

### References

- 1. Traineau H, Aggarwal R, Monfort JB et al. Treatment of calcinosis cutis in systemic sclerosis and dermatomyositis: a review of the literature. J Am Acad Dermatol 2020;82:317-25.
- 2. Balin SJ, Wetter DA, Andersen LK, Davis MD. Calcinosis cutis occurring in association with autoimmune connective tissue disease: the Mayo Clinic experience with 78 patients, 1996-2009. Arch Dermatol 2012;148:455-62.

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