## Disseminated cutaneous gout: A rare clinical presentation

Sir.

Gout is a common systemic disorder caused due to abnormal uric acid metabolism. Uric acid crystallizes and gets deposited in the joints resulting in recurrent arthritis. Chronic cutaneous gout is characterized by firm, erythematous nodules called tophi which are present intradermally or in the subcutis. They usually occur in avascular tissue over the ears, olecranon and pre-patellar bursae, or over acral areas around the joints. [1,2] We report a case of severe, disseminated cutaneous gout that involved non-articular sites and was associated with sepsis.

A 58-year-old Korean male presented with tender, vellowish-red nodules on both upper and lower extremities since one week, along with fever and malaise. He had a history of chronic gouty arthritis for twenty years. Cutaneous examination revealed multiple, firm, yellowish to red nodules over an erythematous base along both sides of his forearms and legs [Figure 1]. The lesions were warm and swollen. There was no sign of venous insufficiency. The patient was drowsy but his neurological examination was within normal limits. Laboratory investigations revealed an elevation of all of the following: total leukocyte count with band forms, blood urea nitrogen, serum creatinine and serum potassium levels (suggestive of renal failure). The serum uric acid level was within normal limits. His chest radiograph revealed cardiomegaly. Two deep punch biopsies of the nodules over the left thigh were taken. Gross examination revealed whitish material similar to chalk. Histopathology revealed amorphous, pinkish, crystalline material in the dermis, surrounded by granulomatous inflammation [Figure 2]. Polarized light microscopy revealed negatively birefringent urate crystals with typical needle - like shapes [Figure 3]. These findings were consistent with intradermal tophaceous gout. Pseudogout was excluded by the absence of characteristic blue refractive crystals. The patient's renal function worsened over the next two days despite emergency hemodialysis. Blood culture revealed the presence of *Staphylococcus haemolyticus* and *Staphylococcus epidermidis*. He developed sepsis and expired due to ventricular tachycardia.

There are four stages of gout: asymptomatic hyperuricemia, gouty attack, intercritical period and chronic gouty arthritis. Skin biopsy and appearance of chalk-like material on the open skin and around joints is the gold standard for diagnosis of gout in the chronic stage. Treatment with allopurinol alone, or in combination with colchicine, is effective for the cutaneous and articular manifestations of gout. [1,2] Atypical forms of tophaceous gout include bullous, fungating and ulcerative gout. Gouty panniculitis has been described in six cases. [3] A variant of cutaneous gout with widespread milia-like eruptions of intradermal tophi is termed as miliarial gout. [4,5] Miliarial gout is characterized by multiple tiny, painless, white to yellow coloured papules on erythematous areas.

In our patient, the lesions were broadly distributed over both the arms and legs, including non-articular sites. A striking peculiarity of our case is that the lesions resembled nodules rather than tiny milia-like papules. It had a clinical presentation more typical of Sweet's syndrome or furunculosis. Serum urate levels are usually elevated in gout. However, gout can even occur in the absence of hyperuricemia limiting the diagnostic utility of measuring serum uric acid levels. Although the exact cause of sepsis was unclear in our patient, there is a possibility that cutaneous infection of the gouty nodules



Figure 1: (a-d) Multiple yellowish to red nodules with underlying erythema over both forearms and lower legs, involving the peri-articular and non-articular regions

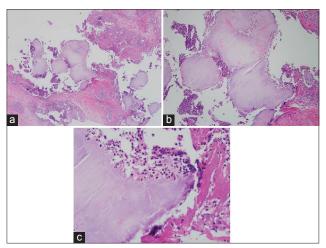


Figure 2: (a-c) Amorphous crystalline material in the dermis, surrounded by granulomatous inflammation (H and E). (a)  $\times 40$ , (b)  $\times 100$ , (c)  $\times 400$ 

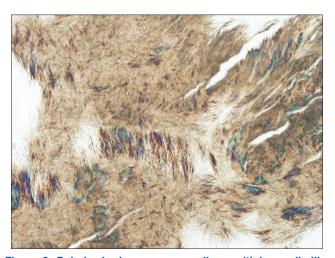


Figure 3: Polarized microscopy revealing multiple needle-like crystals which were negatively birefringent

may have played a role. The histopathological findings of acute inflammatory cells and abscess formation around the pink amorphous material support this theory.

Financial support and sponsorship Nil.

## Conflicts of interest

There are no conflicts of interest.

## Hwa Young Jung, Dong Soo Yu, Jin Wou Kim, Eun Duk Jang<sup>1</sup>

Department of Dermatology, <sup>1</sup>Department of Clinical Pathology, Uijeongbu St. Mary's Hospital, College of Medicine, The Catholic University of Korea, Seoul, Korea

Address for correspondence: Prof. Eun Duk Jang,
Department of Clinical Pathology, Uijeongbu
St. Mary's Hospital, College of Medicine,
The Catholic University of Korea, Seoul, Korea.
E-mail: changed@catholic.ac.kr

## **REFERENCES**

- Falasca GF. Metabolic diseases: Gout. Clin Dermatol 2006;24:498-508.
- Fam AG, Assaad D. Intradermal urate tophi. J Rheumatol1997;24:1126-31.
- Snider AA, Barsky S. Gouty panniculitis: A case report and review of the literature. Cutis 2005;76:54-6.
- 4. Aguayo RS, Baradad M, Soria X, Abal L, Sanmartín V, Egido R, et al. Unilateral milia-type intradermal tophi associated with underlying urate subcutaneous deposition: An uncommon cutaneous presentation of gout. Clin Exp Dermatol 2013;38:622-5.
- Mireku KA, Burgy JR, Davis LS. Miliarial gout: A rare clinical presentation. J Am Acad Dermatol 2014;71:e17-8.

This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 3.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as the author is credited and the new creations are licensed under the identical terms.

| Access this article online |                                  |
|----------------------------|----------------------------------|
| Quick Response Code:       | Website:<br>www.ijdvl.com        |
|                            | DOI:<br>10.4103/0378-6323.164220 |
|                            |                                  |

**How to cite this article:** Jung HY, Yu DS, Kim JW, Jang ED. Disseminated cutaneous gout: A rare clinical presentation. Indian J Dermatol Venereol Leprol 2016;82:204-5.

Received: September, 2014. Accepted: January, 2015.