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[PICTURES IN CLINICAL MEDICINE]

Idiopathic Granulomatous Mastitis, Erythema Nodosum, and Arthritis

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Picture 1.



Picture 3.

A 27-year-old woman presented with a 1-month history of right breast pain, arthralgia, and painful rashes on her legs. A physical examination revealed a reddish, firm, and



Picture 2.



Picture 4.

painful lump on her right breast (Picture 1, arrows); ankle swelling and tenderness (Picture 2); and multiple tender, reddish nodules on both legs (Picture 3). A skin lesion biopsy showed panniculitis, indicating erythema nodosum (EN). Right breast ultrasonography showed an ill-defined, irregular, heterogeneous, hypoechoic lesion with multiple tentacles 1.8 cm in diameter (Picture 4), which are typical find-

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ings of granulomatous mastitis (GM) (1). Aspiration culture (including for mycobacterium) of the breast lump was negative. All blood tests, including serum calcium, angiotensinconverting-enzyme, and lysozyme for diagnosis of sarcoidosis, were normal. Chest computed tomography showed no bilateral hilar lymphadenopathy or pulmonary findings. A diagnosis of idiopathic GM, EN, and arthritis was made. After 2 weeks of prednisolone (60 mg per day) administration, her symptoms resolved promptly. Idiopathic GM is a rare disease and is a differential diagnosis of EN and arthritis because 30% of cases of idiopathic GM cause EN and arthritis (2). Corticosteroids are an effective treatment option (3).

The patient's family provided their consent for the publication of this case with the removal of all identifying information to ensure anonymity and retain the patient's privacy.

The authors state that they have no Conflict of Interest (COI).

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References

- 1. Pluguez-Turull WC, Nanyes EJ, Quintero JC, et al. Idiopathic granulomatous mastitis: manifestations at multimodality imaging and pitfalls. Radiographics **38**: 330-356, 2018.
- 2. Parperis K, Achilleos S, Costi E, Vardas M. Granulomatous mastitis, erythema nodosum and arthritis syndrome: case-based review. Rheumatol Int 41: 1175-1181, 2021.
- Sheybani F, Sarvghad M, Naderi H, Gharib M. Treatment for and clinical characteristics of granulomatous mastitis. Obstet Gynecol 125: 801-807, 2015.

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