Case Report

Soft tissue giant cell tumour revisited - a case report

Debashis Sen, Atul Jain, Banashree Majumdar, Sumit Sen, Pranshu Mishra, Akhilesh Behra

Department of Dermatology, IPGMER and SSKM Hospital, Kolkata

Abstract

Giant cell tumour of soft tissue (GCT-ST) is an uncommon entity which histopathologically resembles bony GCT. We report a case of giant cell tumour in a 45-year-old male patient who presented to the dermatology department with a gradually enlarging painless, smooth swelling over second toe of left foot. Histopathology revealed it to be giant cell tumour of soft tissue.

Key words

Soft tissue tumour, giant cell tumour, osteoclast.

Introduction

Giant cell tumour of soft tissue (GCT-ST) clinically presents as painless, firm, well-defined mass without any fixity to underlying structure. The tumour is usually benign in nature. It can recur after surgical excision. Malignant transformation may occur rarely.

Case Report

A 45-year-old male patient presented with gradually increasing swelling over left second toe with overlying normal skin for 2 years. It was not painful. No history of trauma or similar swelling in any other part of body was present. Clinical examination revealed a nontender, firm, nonpulsatile soft tissue swelling of about 3 to 4 cm diameter without fixity to underlying bone (**Figure 1**). Overlying skin was unremarkable. No limitation in movement of the affected toe was present. On auscultation no bruit was heard. Patient was otherwise normal.

Address for correspondence

Dr. Atul Jain,
Department of Dermatology,
IPGMER and SSKM Hospital, Kolkata
Email: dr.atul86@gmail.com

Routine investigations were unremarkable. Digital X-ray of the affected foot showed homogeneous soft tissue opacity without any calcification or phlebolith. No expansile lytic lesion or cortical erosion was present (Figure 2). Histopathological examination revealed spindleshaped cells arranged in interlacing fascicles. Multinucleated giant cells were present in a scattered manner, diffusely throughout the lesion. Focal area of hyalinization was seen. No significant nuclear pleomorphism or mitotic activity was appreciated. The epidermis showed hyperkeratosis, acanthosis and irregular downward elongation of rete ridges (Figure 3, **4**).

Discussion

GCT-ST is a rare primary neoplasm of soft tissue affecting the age group between 5-84 years with no sex predilection.¹ The tumour commonly involves extremities, trunk, abdomen and pelvis.² However, few reports suggested the commonest site to be an extremity and the finger was mentioned as the most common site.³ Although a benign condition, uncommon malignant transformation has also been noted with rare metastasis to lung⁴ and parotid glands.^{3,5} Metastatic potential is mainly related



Figure 1 A swelling over second toe of left foot. (Inset, view from plantar aspect).



Figure 2 X-ray of left foot showing soft tissue opacity of the second toe (AP and lateral view).

to depth of invasion. Evolution of GCT-ST is not clear, it has been thought that giant cells which cytomorphologically resemble osteoclasts are the result of fusion of circulating monocytes. Histologically, GCT-ST shows well-circumscribed, nonencapsulated tumour of round to spindle-shaped cells admixed with osteoclast like multinucleated giant cells. 5

Differential diagnosis of GCT-ST includes soft tissue mesenchymal tumour which are rich in giant cell, villonodular sinuvitis, cellular dermatofibroma, atypical fibroxanthoma, plexiform fibrohistiocytic tumour, malignant fibrous histiocytoma, leiomyosarcoma, osteosarcoma extraskeletal and epithelioid sarcoma.6

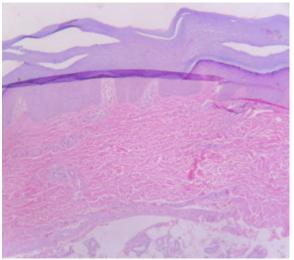


Figure 3 Photomicrograph revealing epidermal hyperkeratosis, acanthosis, and irregular downward elongation of rete ridges (H and E X100).

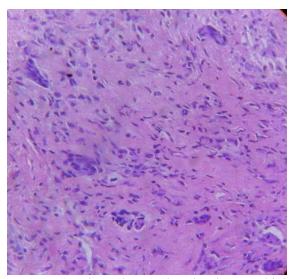


Figure 4 Photomicrograph showing multiple, diffusely scattered osteoclast like giant cells mixed with spindle cell stroma. (H and E X 400).

Immunohistochemistry can give added information regarding the tumour which shows positive stain for CD68, vimentin, tartarateresistant acid phosphatase, cytokeratin, smooth muscle actin.⁶ These investigations could not be performed in our case.

Optimal treatment of GCT-ST appears to be conservative surgical resection with tumour free margin which often provides a good prognosis. Recurrence rate after excision is very low.^{7,8} It is

necessary to follow up these patients carefully because of the rare possibility of malignancy. A dermatologist must have a high index of suspicion and consider GCT-ST in the differential diagnosis of swelling of a toe.

References

- 1. Tejera-Vaquerizo A, Ruiz-Molina I, González-Serrano T, Solís-García E. Primary giant cell tumor of soft tissue in the finger. *Dermatol Online J.* 2008;**14**:7.
- O'Connell JX, Wehrli BM, Nielsen GP, Rosenberg AE. Giant cell tumors of soft tissue: a clinicopathologic study of 18 benign and malignant tumors. Am J Surg Pathol. 2000;24:386-95.
- 3. Asuquo ME, Nwagbara VI, Agbor C *et al.* Giant cell tumour of soft tissue of the finger: a case report. *J Clin Exp Oncol.* 2013;2:3.

- 4. Guo H, Garcia RA, Perle MA *et al*. Giant cell tumor of soft tissue with pulmonary metastases: pathologic and cytogenetic study. *Pediatric Dev Pathol*. 2005;**8**:718-24.
- Angervall L, Hagmar B, Kindblom LG, Merck C. Malignant giant cell tumor of soft tissues: a clinicopathologic, cytologic, ultrastructural, angiographic, and microangiographic study. *Cancer*. 1981;47:736-47.
- 6. Hoang MP, Rogers BB, Albores-Saavedra J. Giant cell tumor of the skin: a morphologic and immunohistochemical study of five cases. *Ann Diagn Pathol.* 2002;**6**:288-93.
- 7. Oliveira AM, Dei Tos AP, Fletcher CD, Nascimento AG. Primary giant cell tumor of soft tissues: A study of 22 cases. *Am J Surg Pathol*. 2000;**24**:248-56.
- 8. Guccion JG, Enzinger FM. Malignant giant cell tumor of soft parts: an analysis of 32 cases. *Cancer*. 1972;**29**:1518-29.