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Chest wall secondary chondrosarcoma arising from enchondroma in a young Asian female



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ABSTRACT

INTRODUCTION: Enchondromas are benign tumours derived from cartilaginous tissue and they rarely undergo malignant transformation unless they occur in multiple sites such as in the case of Ollier disease. **PRESENTATION OF CASE:** We report a rare case of a 36-year-old Chinese female, without any past medical history, who had a right chest wall tumour which was initially diagnosed to be benign, for which she was lost to follow up. The mass had been stable in size for the past 7 years until recently, when it gradually grew bigger. computed tomography (CT) scan of the thorax revealed that the mass had invaded the right chest wall suggestive of malignant change. She subsequently underwent an elective open right chest wall resection of the mass and chest wall reconstruction with mesh. A 3 cm by 3 cm tumour arising from the anterior aspect of the 4th rib was found intra-operatively. Post-operative course was uneventful. Histologically, the mass was confirmed to be chondrosarcoma and resection margins were negative. The patient was discharged on 4th post-operative day and her latest CT thorax during follow up 2 years after operation did not show any tumour recurrence.

DISCUSSION: The aim of chest wall reconstruction is to avoid instability and prevent thoracic organs from being exposed if the defect size is ≥ 10 cm or at least three ribs are resected. Polypropylene mesh may be relatively weak in terms of mechanical support for the chest wall, however, in this case, the defect is not too large and the laterally reflected pectoral muscle can provide enough support together with the mesh. **CONCLUSION:** Secondary chondrosarcoma in a young Asian female arising from a benign solitary enchondroma in the absence of multiple enchondromatosis, such as Ollier disease or Maffucci syndrome, is extremely rare. Regular follow up including physical assessment and radiological imaging can result in earlier detection of malignant transformation obviating the need for aggressive surgical treatment involving multiple rib resections.

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1. Introduction

Chondrosarcomas involve the trunk often, but rarely arise from the chest. Surgical excision is the mainstay of treatment because they are not very responsive to radiotherapy or chemotherapy. The surgeon must ensure complete excision of the lesion with adequate margins to avoid recurrence, whilst not to injury surrounding structures and provide chest wall reconstruction to cover the defect if necessary.

2. Case

We describe a case of malignant transformation of a benign right chest wall tumour to chondrosarcoma in a young Asian female and the management of her condition.

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A 36-year-old Chinese female presented with a seven-year history of a chest wall enchondroma, previously diagnosed via biopsy, that had enlarged in the recent months. She did not have any medical problems, neither did she smoke or drink. She was asymptomatic apart from the swelling. A chest radiograph was done at the clinic which showed a 3.3 cm \times 3.8 cm opacity in the right middle zone with a well-defined inferior border and a poorly defined superior border. There was a marked increase in size compared to the chest radiograph seven years ago (Fig. 1). Computed tomography scan of the thorax revealed an extra-pulmonary soft tissue mass, measuring 4.2 cm \times 3.8 cm, arising from anterior 3rd rib with some bony densities and specks of calcification, which was highly suggestive of a malignant lesion with invasion of the right chest wall (Figs. 2 and 3).

She was then admitted electively for excision of the right chest wall tumour. Upon admission, her vitals were stable. On examination, she was well. There was a palpable hard mass approximately 3 cm \times 3 cm over the right 3rd intercostal space which was mildly tender upon palpation and fixed to the chest wall. No other lump or axillary lymph node was palpable.

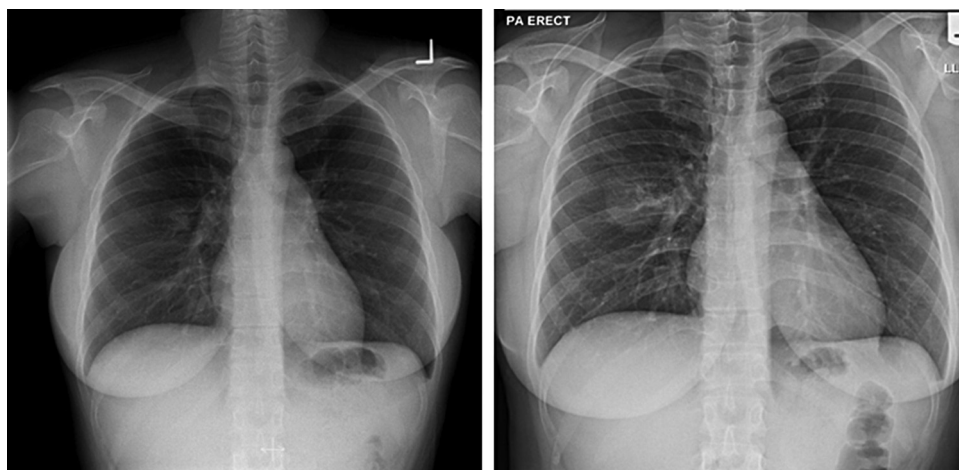


Fig. 1. Comparison of chest radiographs of patient in 2006 (left) and 2013 (right).

The patient subsequently underwent right chest wall tumour resection and reconstruction with mesh. Intra-operatively, a 3 cm × 3 cm tumour arising from the anterior aspect of the 4th rib was found which was removed en bloc including part of the 3rd to 5th ribs using diathermy and rib cutter with adequate margins of 3–4 cm (Fig. 4). The lungs, pleural and diaphragm were not involved by the tumour. The chest wall defect was about 6 cm × 8 cm and was reconstructed using a polypropylene mesh. Histologically, the mass was chondroid in nature, compatible with that of the diagnosis of grade 1–2 chondrosarcoma by the pathologist (Figs. 5 and 6). The margins of the specimen were negative of malignancy.

Her post-operative course was uneventful. Patient was subsequently discharged on 4th post-operative day and her latest CT scan on follow up 2 years after surgery did not show any recurrence.

3. Discussion

Chondrosarcomas generate chondroid matrix and are malignant. They can be primary, arising de novo, or secondary from pre-existing enchondroma or osteochondroma which underwent malignant transformation.^{1,2} They are the commonest primary chest wall tumour.³ The peak incidences for primary chondrosarcoma and secondary chondrosarcoma are 40–60 years of age and 25–45 years of age respectively.³ The usual presentation of a chest wall chondrosarcoma may include palpable mass or localised chest wall tenderness.⁴ Usual sites of secondary chondrosarcomas are anterior costochondral junction, costal angles, sternum and vertebral body.⁴ Computed tomography of the thorax depicts bony densities and specks of calcification.⁴ For this patient, the extensive

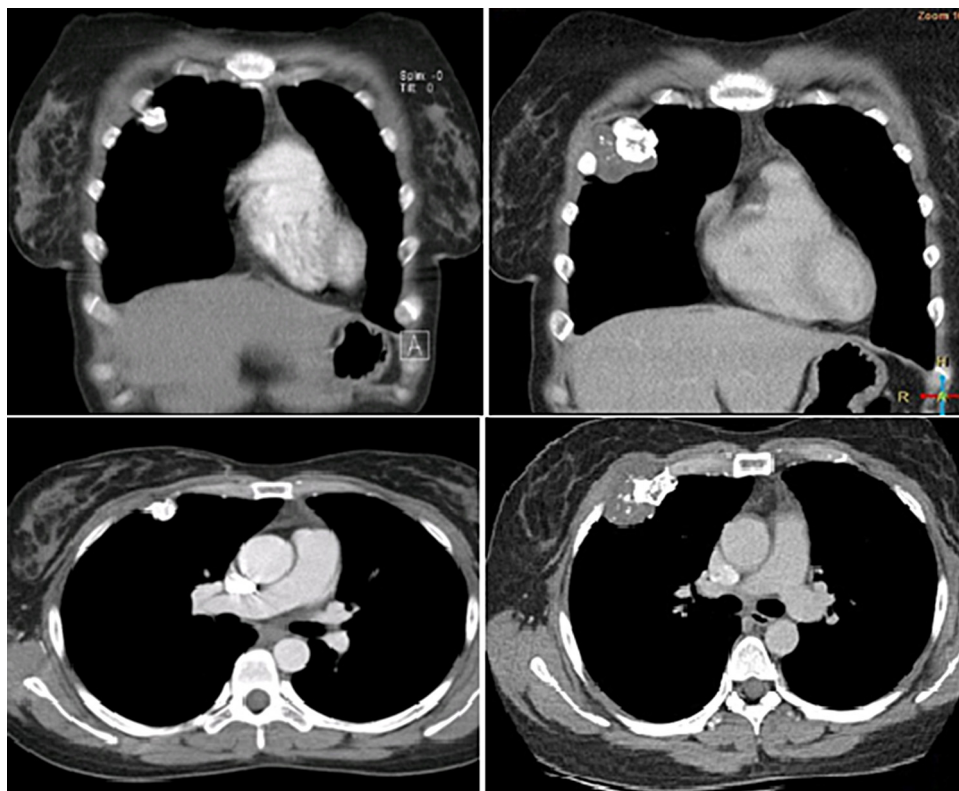


Fig. 2. Comparison of computed tomography of thorax of patient in 2006 (left) and 2013 (right), coronal cuts (above), axial cuts (below).

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