Symptomatic fibrous dysplasia of the right first rib excised via a posterolateral thoracotomy

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Primary tumours of the first rib remain a rarity, but their diagnosis and management present difficult clinical problems. We describe the management of one such case.

Case report

A 35 year old housewife presented to her general practitioner complaining of discomfort in her right supraclavicular fossa for some weeks past. No abnormality was found on examination at this time, but a chest radiograph (fig 1) showed a 3.5 cm calcified expansion of the anterior end of the first rib. Radiologically this was considered to be a chondroma, although malignancy could not be excluded. On examination at this time a smooth mass could be palpated just posteriorly to the clavicle. There was no evidence of a neurovascular lesion and no regional lymphadenopathy. A mass radiograph taken five years previously showed no evidence of this lesion. In view of the recent history of pain, excision was deemed necessary to confirm the pathological nature of the mass. The anterior three quarters of the first rib, including the tumour, was excised through a posterolateral approach after division of the second rib (fig 2). Postoperative recovery was rapid and uneventful. Histological examination showed the lesion to be fibrous dysplasia. The cosmetic result was good and the patient remains well.

Discussion

Primary rib neoplasms are rare, and with neoplasms of the sternum account for about 5–10% of bony tumours.1 They present management problems because of the difficulty of assessing clinically and radiologically whether they are benign or malignant. Histological assessment is difficult with needle biopsy or incision biopsy as considerable histological variation can occur throughout the tumour. Rapid increase in volume, the development of pain, and the size itself have been suggested as indicating malignancy. From 33%2 to 66%3 of rib neoplasms are found to be malignant. The possible causes of a lesion such as this are many and varied and include osteochondroma, chondroma, chondrosarcoma, osteosarcoma, chondromyxoid sarcoma, fibrous dysplasia, and haemangioma.4,5 Fibrous dysplasia and osteochondroma are the most common benign tumours of ribs in most series.5,6 The treatment for all tumours is complete excision because even some of those tumours considered to be benign have been shown to recur and undergo malignant degeneration.6 Primary malignant lesions should be widely excised so that a region including the ribs above and below the tumour is removed.2,6

In view of the unknown nature of the lesion in our patient, excision of the mass was considered necessary. Neoplastic lesions of the first ribs are very rarely reported,4 and surgical access to the first rib is most commonly indicated as part of the treatment for relief of the "thoracic outlet syndrome." Three approaches have been described for excision of the first rib for relief of this syndrome—the anterior,7 the transaxillary,8 and the posterolateral approach.9 The aims of treatment of the thoracic outlet syndrome are removal of enough first rib to allow decompression of the outlet and if possible to give a good cosmetic result. All three procedures have their advocates but most would agree that anterior and transaxillary approaches give a good cosmetic result. The posterolateral approach, however, has the advantage of excellent exposure of the entire first rib, especially in the obese or muscular individual. In addition, any complication of the procedure can be corrected adequately through this.
incision, especially after osteotomy of the second rib.\textsuperscript{10} In view of this a posterolateral approach was preferred for our patient and first rib resection was carried out via this approach without difficulty. Postoperative recovery was uneventful and the cosmetic result was also satisfactory.

References


Book notices


This volume contains the proceedings of a conference held in June 1985 at the American College of Cardiology. About 50 papers are included, many of which amount to little more than speculative expressions of belief and hope. Other papers offer interesting descriptions of experience in educating patients, varying from the problems of educating illiterate patients in the third world on the one hand to descriptions of the design of interactive videodisc programmes for educating individual North American patients on the other. Although most of the papers are inspired by the problem of improving understanding of the treatment and prevention of cardiac disease, some of them have wider relevance and could interest workers in other medical disciplines. The book offers an unusual concentration of the relatively few references to published work on one particular aspect of medical education—education of the patient.—AB


This book is intended for a very narrow readership of respiratory therapists. It is an attempt to produce a fully comprehensive learning system that encompasses everything from the cell to the use of drugs in cardiopulmonary resuscitation, pulmonary rehabilitation, and neonatal and paediatric respiratory care. Its comprehensive nature may be demonstrated by the fact that it includes sections on how to answer the telephone and how to sit a patient up in bed. There is a long section on the medicolegal consequences of accepting responsibility referred to members of the medical profession. Each chapter starts with a list of learning objectives, and ends with a series of tabulated procedures. It is profusely illustrated. Discussion with a senior physiotherapist supports the view that it is unlikely that such a book will find a market in the United Kingdom, where there are no equivalents of the respiratory therapist.—JCS