Abstract A series of 8 cases of elastofibroma is reported, and the clinical, pathological and imaging features and different therapeutic modalities are reviewed. On this basis, we suggest an algorithm for the diagnosis and treatment of elastofibroma. Briefly, marginal excision is the treatment of choice in symptomatic patients, while follow-up appears to be a good solution in asymptomatic ones.

Keywords Elastofibroma · Dorsi · Scapula · Excision

Introduction

Elastofibroma dorsi is a rare pseudotumor developing in a highly characteristic anatomical location. In fact, it is usually localized anterior to the inferior pole of the scapula and the serratus anterior muscle over the ribs of the thoracic cage. Very rarely, elastofibroma is found in other sites, such as the tip of the elbow, near the ischial tuberosities, the deltoid muscle, foot, inguinal region, orbits, stomach, greater omentum and the intraspinal spaces [1–10]. It is relatively frequent among elderly women, with a mean age at onset of 70 years. It is bilateral in 10% of cases [1, 8, 10].

We reported a series of eight patients affected by elastofibroma. We discuss the clinical and radiological features and treatment modalities and review the literature concerned.

Case series

Seven women and one man of mean age 61 years (range, 47–82 year) were treated for elastofibroma during the period from 2000 to 2004 (Table 1). All had unilateral lesions except one woman who had bilateral masses. Every case was studied with radiography, ultrasound, computed tomography (CT) and gadolinium (MRI) enhanced magnetic resonance imaging. The patients were divided into 2 groups: asymptomatic (2 cases) and symptomatic (6 cases).

The asymptomatic patients were treated conservative-ly, and achieved a good range of motion without any increase of the mass’s dimensions 14 and 15 months after the diagnosis. The 6 symptomatic cases, complaining of snapping of the scapula, were treated surgically by mar-
original excision of the mass without performing a preliminary biopsy. Postoperatively, the operated shoulders were immobilized with a support for 3 weeks followed by rehabilitation and physical therapy. These patients were followed for a mean period of 27 months (range, 18–37 months). During this period, no recurrence of the elastofibroma was observed, and the patients were pain free with a completely normal shoulder range of motion.

Discussion

Elastofibroma is a rare lesion. In fact most publications [1–6, 8–10] reported either a single case or a small series of 2–7 cases, except for Nagamine et al. [7] who reported a study of 170 cases. These lesions tend to be slowly growing and asymptomatic in more than 50% of the cases. When symptoms are present, they are typical, consisting of local scapular swelling and a clunking sensation during abduction and adduction of the shoulder, with pain of moderate or, rarely, severe entity [2, 3, 5].

The location of elastofibromas and its incidence among older individuals may be related to the natural existence of fibro-elastoc tissue in this region and suggests a reactive process in response to friction of the scapula against the ribs [2, 3, 10]. Alternatively, the genesis of elastofibroma is thought to involve an overproduction of the collagenous connective tissue with a degeneration of the collagen fibres and an overproduction of immature elastic tissue, derived from fibroblasts, alternating with deposition of hyperplastic fat [2, 8, 10]. Genetic factors may also be involved [2, 8, 10].

Ultrasound examination shows an abnormal mass of tissue in the typical location of the elastofibroma with an alternating pattern of hypoechogenic and hypoechochogenic lines that are roughly parallel to the chest wall (Fig. 1). MRI is considered to be the investigation of choice because it shows the alternating pattern of fibrous and fatty tissues (Fig. 2). The lesions appear poorly circumscribed, heterogeneous with the margins that may be sharp and indistinct. On T1-weighted and T2-weighted sequences, fibrous tissue produces low-intensity signals identical to that produced by muscular tissue, while the fatty tissue is seen as a high-intensity signal on T1-weighted sequences and as an intermediate signal on T2-weighted sequences. Streaks of fatty tissue alternate with

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**Table 1 Clinical characteristics of 8 patients with elastofibroma**

| Patient | Sex | Age (years) | Location | Size (cm) | Treatment | Follow-up (months) |
|---------|-----|-------------|----------|-----------|-----------|-------------------|
| 1       | F   | 47          | Right    | 5.3       | Surgery   | 18                |
| 2       | F   | 63          | Right    | 4.9       | Surgery   | 25                |
| 3       | F   | 59          | Left     | 11.7      | Surgery   | 37                |
| 4       | F   | 82          | Right    | 3.8       | Conservative | 15            |
| 5       | M   | 58          | Right    | 6.18      | Surgery   | 29                |
| 6       | F   | 68          | Bilateral | 7.8–6.9  | Surgery   | 24                |
| 7       | F   | 54          | Left     | 3.5       | Conservative | 14            |
| 8       | F   | 57          | Left     | 6.9       | Surgery   | 27                |

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**Fig. 1 Ultrasound examination shows an abnormal mass in a location typical of elastofibroma, with an alternating pattern of hyperechogenic and hypoehogenic lines**
strands of fibrous tissue forming straight or curved lines. On STIR sequences, the mass is seen as a mosaic of low and high intensity areas. The elastofibroma appears as a large mass with ill-defined contours and marked gadolinium enhancement [6, 8–10]. CT is less sensitive than MRI in visualizing the fatty tissue, so that the elastofibroma may be seen as a homogeneous mass with a density inferior to that of muscles. Moreover, CT shows the absence of bone abnormalities.

Elastofibroma exhibits a characteristic structure where streaks of fatty tissue are alternated with strands of fibrous tissue. The hypertrophic fibrous tissue contains fibrillated material with identical staining affinities to that of necrotic fibrous tissue, muscle, and fat (Fig. 3).

Routine biopsy has been considered to rule out soft tissue sarcoma [4–9]. On the contrary, most of the recent publications indicate that biopsy is unnecessary when the MRI findings are sufficiently typical [1–3, 5, 6, 8]. In particular,
Hayes et al. [2] suggested a core biopsy only in case of suspicion of a soft tissue sarcoma. Turna et al. [9] considered a histological evaluation by intra-operative frozen section and performed complete resection in case of a benign histological results. On the other hand, Vastamaki [10] considered incisional biopsy unnecessary when the clinical findings are typical; thus biopsy is indicated only when the sub-scapular mass has enlarged very quickly within a few months.

Concerning the modalities of treatment, controversy is reported in the literature. Marginal excision has been sug-
gested because it carries a low recurrence risk [1, 5, 8–10], and no malignant transformation has been report-
ed. Guha and Reja [1] mentioned the marginal surgical
excision for these lesions and reported only one case of
recurrence after the excision; another case was success-
fully treated with radiotherapy. Malghem et al. [6] adop-
ted a wait and see approach in asymptomatic patients;
since these patients did not develop symptoms, the
authors concluded that elastofibroma is an unngrowing
lesion after the diagnosis.

Considering these reports, we suggest a diagnostic
and therapeutic algorithm for elastofibroma (Fig. 4). In
case of a scapular region mass in an elderly woman,
radiography, ultrasound and MRI with gadolinium
enhancement are required. If the patient is asympto-
matic and there is no enhancement on MRI, clinical fol-
low-up is sufficient. On the other hand, in symptomatic
patients with no enhancement, marginal excision of the
mass is necessary. Marginal excision is also indicated
in the absence of symptoms in the presence of enhance-
ment on MRI. We recommend biopsy to rule out sarco-
ma when both symptoms and MRI enhancement
are present.

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