Imaging of Elastofibroma: A Rare Forgotten Tumor of Chest Wall

Abstract
Elastofibroma is a rare benign tumor of chest wall occurring commonly in elderly female and showing abundant elastin fibers. The lesion is usually large at the time of presentation and requires surgical removal only in symptomatic patients. The lesion has a characteristic appearance on imaging especially on computed tomography (CT) scan and usually requires no further intervention. The present article discusses this rare tumor in young adult male with its characteristic CT scan findings, differential diagnosis and review of literature as this tumor is benign and less often recognized by radiologists.

Keywords: Elastofibroma, computed tomography

Introduction
Soft tissue tumors of the chest wall including elastofibroma [also known as elastofibroma dorsi] are rare [1]. Elastofibroma is most frequently encountered in middle aged to elderly females in infrascapular location, more commonly on right side but may be bilateral. As imaging and microscopy [FNAC or biopsy] can achieve the diagnosis in majority of cases, hence surgical intervention should be reserved for symptomatic cases [1,2].

Case Report
A 25 years old male presented to us with complaint of soft tissue swelling and pain in right infrascapular region. Local examination revealed palpable soft tissue mass with freely mobile overlying skin and slight pain only on performing scapular movements. No evidence of any other external abnormality or tenderness was noted. Laboratory examination, chest radiograph and ultrasonography (USG) of abdomen were unremarkable. However, high resolution USG of the mass revealed predominantly solid-appearing, fairly-defined, smooth-margined, echogenic mass interspersed with some linear hypoechoic areas. No evidence of any internal calcification or frank necrosis was noted. Contrast-enhanced, computed tomography (CT) scan of the thorax revealed a nonenhancing, soft-tissue attenuating, solid mass lesion along the posterolateral aspect of right thoracic wall in infrascapular location (Figure 1). The tumor was lying between the chest wall and serratus anterior and latissimus dorsi muscles. The tumor revealed hypodense internal areas of fat attenuation (Figure 2). The lesion measured approximately up to 135*80*65mm corresponding to approximately up to 350-400 ml in volume. However, no evidence of any obvious internal calcification or necrosis was noted. No evidence of any obvious adjacent osseous involvement or intrathoracic extension was noted.

Based on the clinicoradiologic findings, the diagnosis of elastofibroma of chest wall was suggested and follow up was suggested. Six months follow up of patient did not reveal any significant change in the clinical symptoms, clinical appearance and size of the tumor. However, as the patient was apprehensive, hence open biopsy was planned to exclude the sarcomatous lesion and to reassure the patient, that surgical excision is not indicated (Figure 3).

Histopathology revealed fibrous tissue containing clusters of elastin and collagen fibers interspersed with fatty areas confirming the diagnosis of elastofibroma (Figure 4).

Discussion
Elastofibroma is a rare, benign tumor of chest wall arising from the connective tissues of dermis, occurring commonly in elderly females. It was first reported by Jarvi and Saxen in 1959. Majority of them are bilateral while in unilateral cases, right side is commoner than left side. The characteristic location is infrascapular or subscapular regions between the serratus anterior, rhomboid & latissimus dorsi muscles and chest wall. Uncommonly, the tumor may occur in the deltoid muscle and region of ischial tuberosity, greater trochanter, olecranon process.
of ulna, foot and mediastinum [3]. The tumor grows very slowly and is more prevalent in persons involved in manual labor involving hands and arms.

Though the etiology is unknown yet it is considered by some to represent a reactive process or a fibroblastic pseudotumor as it is characterized by accumulation of abnormal elastin fibers [4]. An abnormal gene on chromosome 1 and 2 has been described attributing some of the lesions as genetic [5].

Majority of patients are asymptomatic with few of them being palpable initially. When large, they are associated with clicking/snapping sensation or pain during scapular movement. Rarely, it may involve the overlying skin which may ulcerate [6].

Imaging features are characteristic with high-resolution ultrasonography showing solid, echogenic, well-defined, fibroelastic tissue interspersed with multiple, hypoechoic, linear areas of fatty tissue without internal necrosis or calcification or vascularity on color Doppler imaging [7]. Computed tomography (CT) scans show fairly-defined, nonenhancing or minimally-enhancing, mass often lentiform or semilunar-shaped, appearing isodense to adjacent muscles in characteristic infrascapular or subscapular location that shows internal linear hypodensities of fatty attenuation [7,8]. No evidence of any obvious sign of internal necrosis or calcification is noted. They are rarely attached to the underlying periosteum.

Magnetic resonance imaging (MRI) shows non-capsulated, solid mass in characteristic location appearing isointense to adjacent muscles on T1W and T2W images showing internal linear hyperintensities corresponding to fat on both T1W and T2W images without obvious restriction on DW images. No evidence of any obvious internal calcification or necrosis is noted. Post-Gadolinium images show mild, near-homogeneous enhancement [7-9].

FDG PET-CT shows uptake on tracer allowing incidental detection of clinically silent small elastofibroma [10]. No evidence of malignant transformation has been described in the literature.

Few cases of elastofibroma have been described in literature with majority of them in elderly females unlike our index case in young male [11].
Histopathological diagnosis is needed in equivocal cases or when the patient is highly-apprehensive of cancer by performing a surgically biopsy. It reveals tumor-like masses of collagen and elastin fibers with scattered areas of fat on microscopy. Complete excision of the tumor is indicated in symptomatic and equivocal cases.

Differential diagnoses that need to be considered in the characteristic location of elastofibroma include lipoma, fibromatosis, sarcoma and metastases. Lipoma is an encapsulated lesion showing homogeneous internal fat without obvious postcontrast enhancement. Fibromatosis appears as solid, soft-tissue tumor with irregular margins without internal striations or fat and showing mild to moderate postcontrast enhancement. Sarcoma is fast-growing, infiltrative tumor showing moderate to intense postcontrast enhancement and internal necrosis with invasion/destruction of adjacent structures. Confident diagnosis of metastases requires a known primary tumor.

To summarize, elastofibroma is a rare, benign, soft tissue tumor of chest wall that can be confidently diagnosed on imaging obviating the need for excision biopsy or major surgery in asymptomatic patients. Small elastofibroma may be incidentally detected on PET-CT where they need differentiation from metastases.
References


