HYPERMETABOLIC BILATERAL INFRASCAPULAR LESIONS IN PET-CT IMAGING: ELASTOFIBROMA DORSI

Case Report

PET-BT GÖRÜNTÜLEMEDE HİPERMETABOLİK BİLATERAL İNFRASKAPULER LEZYONLAR: ELASTOFİBROMA DORSİ

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ABSTRACT

Elastofibromas are benign pseudotumoral fibroelastic lesions commonly localized in the periscapular region in elderly women. The etiology of these lesions are not known. They were first reported in 1961. These lesions are well recognized in the pathology literature and have also received some attention in the radiology literature recently. They are often confused with neoplasms on CT (Computed Tomography) or MRI (Magnetic Resonance Imaging). We present a patient with infrascapluar elastofibroma dorsi. Bilateral subscapuler 18-Deoxy-Glucose) FDG (Flour hypermetabolism PET in (Positron Emission Tomography) were detected and these images were compared to CT or PET-CT fusion images, after which we elastofibroma diagnosed dorsi. This technique may prevent patients undergo unnecessary biopsies and surgical procedures.

Keywords: Elastofibroma; pseudotumor; positron emission tomography; hypermetabolism.

ÖZET

Elastofibromlar genellikle orta-ileri vastaki kadınlarda, skapula cevresindeki dokularda yerleşmiş olan benign natürde, psödotümöral fibroelastik lezvonlardır. Etiyolojisi hala belli değildir. Patoloji literatüründe çok iyi bilinen ve son yıllarda radvoloji literatüründe ilgi ile karsılanan bu lezyon ilk olarak 1961 yılında tarif edilmistir. Bu patoloji bilgisayarlı tomografi (BT) veya manyetik rezonans görüntüleme (MRG) sırasında tespit edildiklerinde tümörler ile karıştırılabilirler. calışmada infraskapülar yerleşimli Bu elastofibroma dorsi tespit edilen bir hasta bildirilmiştir. Hastanın positron emisyon tomografi (PET) görüntüleme sırasında izlenen FDG hipermetabolizması, BT veva PET-BT füzyon görüntüleri ile karşılaştırılaraık elastofibroma dorsi tanısı kondu. Bu yöntem hastaları gereksiz cerrahi girişim veya biopsi işlemlerine maruziyetten koruyabilir.

Anahtar kelimeler: Elastofibroma; psödotümör; positron emisyontomografi; hipermetabolizma.

INTRODUCTION

Elastofibroma dorsi (ED) is a benign pseudotumoral lesion first described by Saxen in 1961. Jarvi ve It is characteristically derived from abnormal fibroelastic tissue proliferation in subscapular localization. (1,2) It is reported in 11-24% of autopsy reports (3). It's detection rate on thorax computerized tomography is 2%. The reason for this difference is milimetric elastofibroma lesions cannot be detected on CT. ED is usually asymptomatic and sometimes can cause symptoms like shoulder pain, shoulder motion limitation, or growing soft tissue mass (4). ED is seen bilaterally in 10% of pateints. Most frequent localization of ED is serratus anterior, latissimus dorsi and rhomboid muscles. ED lesions are smooth limited and have similar density with muscle. They may include band style hypodense streaks of fat tissue (5). Similary, MRI appearance of ED is similar to muscle tissue (hypointense on T1A and T2A images with linear hyperintense streaks (6). To our best knowledge, a few studies have reported hypermetabolic activity on PET analysis of ED lesions (7-9).

CASE REPORT

A 78 year old female patient, who had been operated for carcinoma of sigmoid colon two years ago, was operated again for a relapse lesion 6 months ago. She had defecation difficulty and pelvic pain. She had the same complaints 2 years ago when she was diagnosed of cancer. PET-CT was ordered for advanced evaluation of the patient.

PET-CT examination was performed with a 16-detector device composed of CT and

PET fusion (GE Healthcare Discovery ST Hpower 60 PET-CT System). PET-CT was performed 45 min after giving patient 575

MBq FDG intravenously. Patient had not consumed any food or drink except water for five hours before process. The section thickness was 3.75 mm in study. After the procedure, PET-CT fusion images were evaluated by processing images on the work station (GE Advantage Workstation Release 4.3 Software).

Minimal thickening of pelvic intestinal loops' walls and density increase that heterogeneous showing no contrast enhancement in presacral fat-marked plans described in previously applied contrast-enhanced abdominal CT patient. FDG examinations of the accumulation was not detected at a pathologic level during evaluation of PET-CT images. However, in fusiform softtissue masses which deep-seated in both infrascapular and symmetrically followed, has similiar density with surrounding muscle tissue in CT images, homogeneous FDG uptake that can be distinguished more demarkable than environment plans was observed. FDG uptake which also homogeneous in the coronal (Figure 1) and axial (Figure 2) sections, the findings

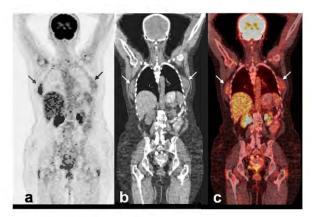


Figure 1. 78 year old female with elastofibroma dorsi. In view of PET (a), CT (b) and fusion (c) images obtained in the coronal plane, in fusiform soft-tissue masses which deep-seated in both infrascapular and symmetrically followed, has similiar density with surrounding muscle tissue in CT images, homogeneous FDG uptake that can be distinguished more demarkable than environment plans was observed.

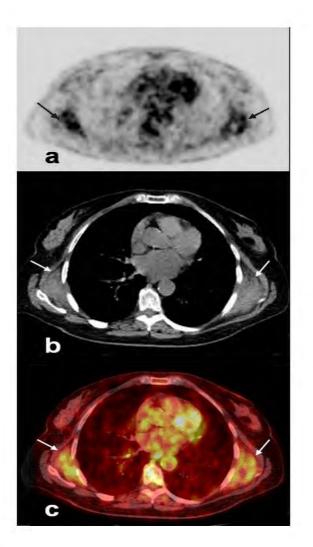


Figure 2.78 year old female with elastofibroma dorsi. In view of PET (a), CT (b) and fusion (c) images obtained in the axialplane, orientation to localization of the lesion is also easier, FDG uptake is observed as deep-located in serratus anterior muscle in infrascapular localization.

CT and PET-CT fusion images and have SUVmax value of 3.69 in right and 3.24 in left showed the activity in these lesions that located deep in serratus anterior muscle in infrascapular localization.

In our case who is without focus of FDG uptake supporting residual-recurrent malignancy in other areas of the body, without a pain, restriction during each of two upper extremity movement and without respiratory distress, in CT of calcification, heterogeneity, invasion osseous destruction were not identified. Because of it features like location, fat content in the form of striae in CT sections it is bilateral symmetric, and only suggested the follow-up of patients for this lesions that is characteristic for elastofibroma dorsi.

DISCUSSION

Although elastofibroma dorsi was described as rare after it was identified firstly in 1961, it was proved that in fact it is a pseudotumoral mass (1-3). It is usually asymptomatic and effects middle-advanced age female population. It is bilateral in 10-60 % of cases and located in deep of posterolateral chest wall muscles (10).

It has same density with muscle tissue and shows same features with signal characteristics in CT and MRI exhibitions and facilitates the recognition of contained fat striae (5,6).Gocmen et al reported that CT has a low sensitivity than MRI in elastofibroma diagnosis, but thanks to multi-slice CTs used in recent years this difference has disappeared. They reported that elastofibrom diagnosis in 63 year old female patient was made with a clarity that gives the impression of looking pathology specimen with 64-detector CT (11). In our case, diagnosis was made by using a device composed of a 16-detector CT and PET fusion. Again, the appearance multi-layer composed of typical hypoechoic-hyperechoic bands in ultrasonography is typical for also elastofibroma (12).

Elastofibroma dorsi rarely has symptoms like shoulder pain, shoulder motilite limitation, or growing soft tissue mass (4). This lesion which is sometimes a cause of click during shoulder movement (snapping scapula), and sometimes imitating pathology of the rotator muscles in literature is curable with operation in symptomatic cases and have rarely reported in relapse cases (13). The presence of fat cells located between dense fibrous tissue bands in histopathological examination and the

absence of capsule is also getting along with the imaging findings (14).

In elastofibroma dorsi cases that identified in PET-CT studies reported in the literature, FDG uptake (SUVmax values) levels have been ranged from 1.52-2.8 (7-9). In our cases, we determined the FDG uptake at 3,69 right, 3,24 left SUVmax values for elastofibroma dorsi cases.

PET-CT imaging is a relatively new technology, a useful examination method in staging and especially in detecting of the identification and spread of malignancy. Its application is rapidly spreaded by development of CT and its fusion. However, it has lots of mistake point because of FDG affinity in physiological activity and benign lesions. When infrascapuler localized low-level FDG uptake is determined in PET-CT imaging, elastofibroma dorsi can be recognized with the help of CT sections and thus unnecessary forth evaluation, biopsy, surgery can be avoided.

REFERENCES

1)Jarvi O, Saxen AE. Elastofibroma dorsi. Acta Pathol Microbiol Scand 1961;51(suppl 144):83–84. PMID:13789598.

2)Ochsner JE, Sewall SA, Brooks GN, Agni R. Best cases from the AFIP: Elastofibroma dorsi. Radiographics. 2006;26:1873-6. PMID:17102057.

3)Brandser EA, Goree JC, El-Khoury GY. Elastofibroma dorsi: prevalence in an elderly patient population as revealed by CT. AJR 1998;171:977 – 80. PMID:9762978.

4)Vastamaki M. Elastofibroma scapulae. Clin Orthop 2001;392:404–8. PMID:11716415.

5)Naylor MF, Nascimento AG, Sherrick AD, Mc Leod RA. Elastofibroma dorsi: radiologic findings in 12 patients. AJR1996; 167:683–7. PMID:8751681.

6)*Kransdorf MJ, Meis JM, Montgomery E. Elastofibroma: MR and CT appearances with radiologic-pathologic correlation. AJR 1992;159:575– 9. PMID:1503030.*

7)Wasyliw CW, Caride VJ. Incidental detection of bilateral elastofibroma dorsi with F-18 FDG PET/CT. Clin Nucl Med. 2005;30:700-1. PMID:16166850.

8)Patrikeos A, Breidahl W, Robins P. F-18 FDG uptake associated with Elastofibroma dorsi. Clin Nucl Med. 2005;30:617-8. PMID:16100483.

9)Pierce JC 3rd, Henderson R. Hypermetabolism of elastofibroma dorsi on PET-CT. AJR Am J Roentgenol. 2004;183:35-7. PMID:15208104.

10)Nagamine N, Nohara Y, Ito E. Elastofibroma in Okinawa: a clinicopathologic study of 170 cases. Cancer 1982;50:1794–1805. PMID:7116305.

11)Gocmen R, Yesilkaya Y. 64-Channel Multi-Detector Computed Tomography Findings of a Case with Bilateral Elastifibroma Dorsi and Review of Literature. Turkiye Klinikleri J Med Sci 2011;31:471-4.

12)Bianchi S, Martinoli C, Abdelwahab IF, Gandolfo N, Derchi LE, Damiani S. Elastofibroma dorsi: sonographic findings. AJR Am J Roentgenol 1997; 169:1113–5. PMID:9308474.

13)Majo J, Gracia I, Doncel A, Valera M, Nunez A, Guix M. Elastofibroma dorsi as a cause of shoulder painor snapping scapula. Clin Orthop Relat Res 2001;388:200–4. PMID:11451120.

14)Domanski HA, Carlen B, Sloth M, Rydholm A. Elastofibroma dorsi has distinct cytomorphologic features, making diagnostic surgical biopsy unnecessary. Diagn Cytopathol 2003;29:327–33. PMID:14648789.