CASE REPORT Open Access

A case report: a giant cardiac atypical lipoma associated with pericardium and right atrium



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Abstract

Background: Among primary cardiac tumors, atypical lipoma is very rare. In particular, cases with lipomas in both the pericardium and the atria are even rare.

Case presentation: We report the case of a 49-year-old male patient presented to our department because of chest pain. Echocardiography revealed two large masses in both the pericardium and the right atrium. Then the tumors were completely resected and the histopathological examination revealed atypical lipoma. The patient recovered well without any complication and discharged from hospital.

Conclusions: We report a very rare case of a huge atypical lipomas located on the pericardium and right atrium. These tumors were easily detected by echocardiography and final diagnosed after surgical resection and pathological examination.

Keywords: Atypical lipoma, Echocardiography, Cardiac tumor

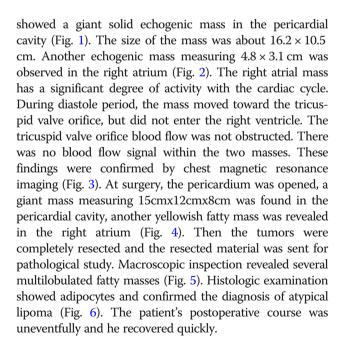
Background

Primary cardiac tumor are not conmen and the incidence range of 0.0017–0.28% as reported in autopsy studies [1]. One quarters of cardiac tumors can be categorized as malignant tumor on histology. The most common malignant tumors of the heart is angiosarcoma whereas atypical lipomas represent <1% [2, 3]. Cardiac atypical lipomas are usually found incidentally because of asymptomatic, and in most cases require treatment or surgical intervention. We report a case of a 49-year-old male patient was referred to our hospital because of chest pain. Echocardiography revealed a giant mass in the pericardial cavity, and another mass in the right atrium.

Case presentation

A 49-year-old male was admitted to our hospital because of chest pain over 20 days. Physical examination and electrocardiography were normal. Transthoracic echocardiography

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Discussion and conclusions

Liposarcoma-a mesenchymal malignant tumor that contains lipoblasts-is the most common soft tissue sarcoma



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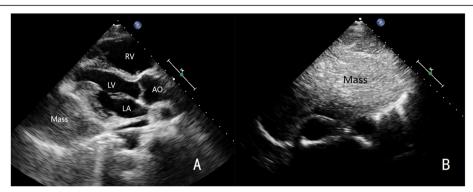


Fig. 1 a Left ventricular long axis view shows a large mass in the pericardium. b Subxiphoid view shows a giant mass in the pericardium. LA, left atrium; LV, left ventricle; RV, right ventricle; AO: aorta

and accounts for 20% of all mesenchymal tumors [4]. The tumor occurs most frequently in the limbs and retroperitoneum and rarely originates in the heart [5]. Peak incidence is between 40 and 60 years old, and there is a slight male predominance [6]. Liposarcoma of the heart is an extremely rare sarcoma, with less than 35 cases reported in the literature according to literature [7]. To our best knowledge, this is the first reported case of simultaneous atypical lipoma in the right atrium and pericardium.

Liposarcomas can be divided into four subtypes: Well-differentiated liposarcoma (or atypical lipoma), myxoid/round cell liposarcoma, pleomorphic liposarcoma, and dedifferentiated liposarcoma. Atypical lipoma was the most common subtype (50%), which had been observed in the atrium and the pericardium. The tumor usually grows slowly, insidious and remain asymptomatic until reach a giant size. Atypical lipomas can be asymptomatic or symptomatic depending on their size and location within the heart. They can invade adjacent organs, so they are more likely to be symptomatic than benign lipomas. Atypical lipomas have a mild biological behavior

and do not generally metastasis, but there is a possibility of focal recurrence.

Although pathological diagnosis serves as a gold standard of diagnosis of atypical lipoma, echocardiography is still essential. Echocardiography is an ideal and important diagnostic modality since it can provide information of the tumor on the size, location, boundary, modality, amount blood flow and so on. Although echocardiography offers a simple and non-invasive approach, it cannot conclusively differentiate between atypical lipomas and other primary tumors of the heart. Computed tomography or magnetic resonance imaging may be of additional value.

There are several diseases should be differentiated from atrial atypical lipomas, namely myxoma, thrombus, benign lipoma. Atrial myxoma is the most frequent primary cardiac tumor in adults. It usually arises in the area of the fossa ovalis, grows on pedicles, has the manner of soft and polyps. The tumor may move towards the ventricle during diastole. Atrial thrombus is usually associated with a history of atrial fibrillation or rheumatic heart disease, and ventricular thrombosis

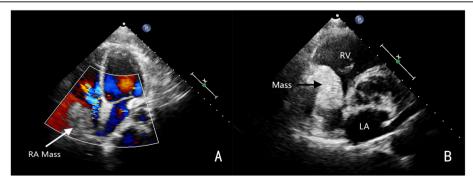


Fig. 2 a The apical four chamber view demonstrates the right atrial mass (arrow), there is no blood flow signal within the mass. b During diastolic period, the mass (arrow) moves toward the tricuspid valve orifice, but does not enter the right ventricle. LA, left atrium; RV, right ventricle

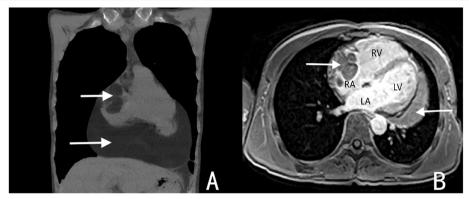


Fig. 3 Magnetic resonance imaging examination. **a** In the coronal section, shows the pericardium and atrial mass (arrow) **b** In the long axis four chamber view, shows the pericardium and atrial mass (arrow)

usually occurs on the basis of abnormal wall motion. The base of the thrombus is wide and can have activity. As time progresses, the echo of the thrombus can gradually change. The size of the thrombus may change after anticoagulant therapy. Cardiac benign lipomas can originate either from the subendocardium (50%), subpericardium (25%), or myocardium (25%) and maybe located more frequently in the left ventricle or right atrium [8]. Cardiac benign lipomas are most often detected incidentally and do not have specific ultrasound findings. The lipoma demonstrated typical loss of signal

RA Mass

Fig. 4 An intraoperative photograph shows a giant mass which locates in the pericardial cavity, another yellowish fatty mass is revealed in the right atrium

PE Mass

on fat suppressed magnetic resonance imaging sequence techniques.

Pericardial atypical lipomas may confuse with malignant mesothelioma, pericardial cyst, hemangiomas. The ultrasound manifestations of malignant mesothelioma usually include pericardial effusion, pericardial thickening, pericardial cavity mass. Pericardial cysts are usually congenial lesion or acquired [9]. Echocardiography showed an isolated, unilocular, smooth-walled cystic mass adjacent to the heart. There was no communicating tract between mass and heart cavity, and no blood flow signal within the mass. Cardiac hemangiomas are extremely rare and account for approximately 2% of all primary resected cardiac tumors. They can locate in atrium, ventricle, pericardium and endocardium [10]. Echocardiography usually presents as a hyperechoic mass with a well-defined boundary, and myocardial contrast echocardiography can demonstrate the vascularity of the mass.

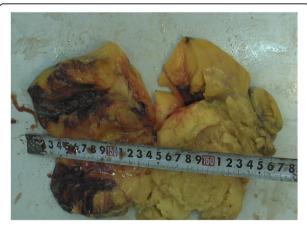


Fig. 5 Macroscopic inspection reveals several multilobulated fatty mass

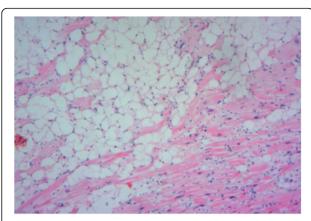


Fig. 6 Pathological examination shows a sheet-like arrangement of adipocyte. The nuclei of local lesion cells slightly enlarge (H&E staining; magnification, \times 20). H&E, haematoxylin and eosin

In conclusion, we report a very rare case of a huge atypical lipomas located on the pericardium and right atrium. These tumors were easily detected by echocardiography and final diagnosed after surgical resection and pathological examination. He is under follow-up for the possibility of recurrence.

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None.

Author's contributions

XW drafted the manuscript. XNY interpreted and revised the manuscript. WDR is involved in ultrasound diagnosis. DYL is the surgeon of the case. All authors read and approved the final manuscript.

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Availability of data and materials

The datasets used in the case are available from the corresponding author upon reasonable request.

Ethics approval and consent to participate

This study was approved by the Ethical Committee of Shengjing Hospital of China Medical University. Written informed consent was obtained from individual participant.

Consent for publication

Written informed consent was obtained from the patient for publication of this case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

Competing interests

The authors declare that they have no competing interests.

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