CASE REPORT

Case Report: Incidentally diagnosed hemangioma of the right atrioventricular groove in an athlete [version 1; peer review: 1 approved]

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Abstract
The purpose of this article is to illustrate a rare case of a pericardial hemangioma of the right atrioventricular groove of incidental discovery in a tennis player who presented with cough and dyspnea and was treated by surgical excision with a favorable outcome. We also report the role of cardiac magnetic resonance imaging (MRI) in the diagnosis and management of this pericardial tumor.

Keywords
Cavernous hemangioma, Cardiac tumors, Pericardium, Tamponade, Athletes

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Introduction
Cardiac hemangioma is a rare benign tumor and pericardial localization is extremely rare. It is usually asymptomatic, but it can be serious due to the risk of tamponade. We report the case of a pericardial hemangioma of the right atrioventricular groove in a young athletic patient who presented with cough and dyspnea and was diagnosed incidentally.

Case report
Patient information and initial presentation
A 31-year-old Caucasian female tennis player presented to the emergency department with dyspnea and dry cough for a few days. She had undergone surgery previously for a borderline ovarian tumor eight years ago. There was no history of cardiopulmonary disease, coronary artery disease, or other cardiovascular diseases. No abnormalities were found during the physical examination with no jugular venous distension.

Diagnostic assessment
A chest X-ray showed enlargement of the cardiac shadow suggestive of pericardial effusion (Figure 1). Transthoracic echocardiography confirmed a large circumferential pericardial effusion and showed a rounded, well defined pericardial hyperechoic lesion attached to the right atrioventricular groove. There was no right ventricular dysfunction.

A thoracic computed tomography (CT) scan was performed, which showed a large pericardial effusion and confirmed a pericardial mass with homogenous contrast enhancement within the right atrioventricular groove (Figure 2).

Cardiac magnetic resonance imaging (MRI) confirmed the large pericardial effusion with a pedunculated ill-defined homogeneous hypointense mass on T1 and a hyperintense mass in the right atrioventricular groove with progressive enhancement after contrast administration on T2 (Figure 3).

A coronary angiography was performed, which showed tumor blush.

Intervention
The patient was referred to a cardiovascular surgery center to be operated on by an experienced cardiac surgeon. General anesthesia was performed in supine position. Anaesthesia induction was performed by intravenous bolus of propofol (2mg/Kg), tracrium (0.5 mg/Kg) and fentanyl (2 mcg/Kg). Anaesthesia maintenance was performed by isoflurane 1.5% in oxygen and continuous intravenous infusion of tracrium (0.01 mg/Kg/min) and fentanyl (1 mcg/kg/hour). Surgery was initiated by a median sternotomy. Initial examination showed no extension of the mass into the cardiac chamber. A safety total excision of the mass was done using cutting diathermy. Vascular, pericardial and sternal sutures were performed by polypropylene, vicryl and wire, respectively. The anatomopathological examination of the mass revealed conjunctive tumor proliferation, vascular differentiated and concluded with a diagnosis of cavernous hemangioma. Post-procedural medication included antibiotic therapy with cefazolin (1 g intravenously, twice a day) for 48 hours, preventive anticoagulation by low molecular weight heparin (Enoxaparin 0.4 ml subcutaneously, once a day) and analgesic...
therapy by paracetamol (1 g intravenously, three times a day). Post-operative course was favorable and the patient was discharged after 72 hours.

Follow-up
Two months after surgery, the patient developed progressive dyspnea vomiting and precordial chest pain. CT scan found loculated left pleural effusion. Chest physiotherapy (one session a day) for two weeks and paracetamol (1 g orally, twice a day) for one week were prescribed with a favorable outcome. The patient remains well after two years of follow-up.

Discussion
Cardiac hemangiomas are rare benign vascular tumors and constitute only 2.8% of primary cardiac tumors. Pericardial localization is extremely rare. Histopathologically, hemangiomas are characterized by benign proliferation of the endothelial cell lining of the blood vessel with increasing vascularization.

Pericardial hemangioma is mostly asymptomatic. Clinical symptoms depend on location, size, and anatomic extension of the tumor. The most frequent symptoms are dyspnea, cardiac arrhythmia, murmurs, and heart failure. Tamponade due to pericardial effusion can also occur. Imaging is very useful for the diagnosis, localization, and extension of the tumor. CT scans with contrast can show enhancing foci at the arterial phase with diffuse or heterogeneous enhancement at the delayed phase. Small calcifications might be seen also. Cardiac MRI is a superior tool with a better contrast resolution. Hemangiomas have an intermediate T1 signal with the same intensity as myocardium and a high T2 signal. The dynamic postcontrast acquisition shows nodular enhancement with progressive fill-in on delayed images. Feeding vessel, tumor blush, and flow voids might be seen also. The tumors are usually ill-defined with no local invasion. Differential diagnoses can be made with solid pericardial masses such as mesothelioma, sarcoma, lymphoma, or paraganglioma. Surgical total excision is the treatment of choice for resectable tumors. The use of radiotherapy, corticosteroids, and beta blockers have been reported in some cases.

Conclusion
Pericardial hemangiomas are extremely rare benign vascular tumors whose prognosis depends on their location and size. Surgical excision constitutes the treatment of choice. Our case demonstrates the importance of cardiovascular MRI as a tool to evaluate the resectability of the tumor.

Data availability
All data underlying the results are available as part of the article and no additional source data are required.

Consent
Written informed consent for publication of their clinical details and clinical images was obtained from the patient.

References
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This is a nice case report of a rare condition that shows the importance of pre-operative imaging. I recommend indexing but a minor revision should be made regarding the description of MR findings.

In the "diagnostic assessment" section (MR part), the authors state that; "there is progressive enhancement after contrast administration on T2". This is misleading and should be corrected as usually contrast enhancement is not to be searched on T2-w imaging. I suppose that enhancement was detected either on post contrast T1-w imaging or on first pass perfusion (Saturation Recovery). This section should be re-written as well as the caption of Fig. 3 that is unclear in the present form.

I suggest the authors to re-use the excellent description from their discussion.

In the discussion please change the first sentence of the second chapter; the patient is asymptomatic, not the hemangioma.

Please delete the 's' at frequent (second chapter).

Is the background of the case's history and progression described in sufficient detail?
Yes

Are enough details provided of any physical examination and diagnostic tests, treatment given and outcomes?
Yes

Is sufficient discussion included of the importance of the findings and their relevance to future understanding of disease processes, diagnosis or treatment?
Is the case presented with sufficient detail to be useful for other practitioners?
Yes

**Competing Interests:** No competing interests were disclosed.

**Reviewer Expertise:** Radiology

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Author Response 09 Sep 2020

**Asma ACHOUR**, Monastir University of Medicine, Tunisia., Monastir, Tunisia

Dear master and colleague,

Thank you for the interest you have shown in our topic.

I thank you for your relevant comments which will improve the quality of our manuscript.

**Competing Interests:** No competing interests were disclosed.

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