Primary cardiac tumour – An unusual incidental finding with significant implications

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Abstract
Primary cardiac tumours are rarely seen in clinical practice and often present with symptoms like shortness of breath, syncope and palpitations [1]. None the less, even more rarely, cardiac tumours can be incidentally found on imaging intended for un-related clinical indications. These lesions carry a significant risk of serious complications even in asymptomatic individuals hence require urgent attention for further investigations and treatment. We describe a case of an otherwise fit and well woman who was found to have a left atrial mass on the CT scan intended to rule out a gastrointestinal malignancy. The left atrial mass was further evaluated by transthoracic echocardiogram and was considered to arise from the inter-atrial septum. She underwent a successful resection of this lesion and histological examination confirmed it to be a myxoma. We present a brief literature review on epidemiology, pathophysiology, clinical features, diagnosis and treatment options for cardiac myxomas. Any suspected primary cardiac mass should be further evaluated as a matter of urgency; surgery is usually the only effective treatment option available in most cases.

Introduction
Primary cardiac tumours are rare in clinical practice with an incidence ranging from 0.0017 – 0.19% in general population [2]. Due to the rarity of this condition, medical literature on pathophysiology, characterisation and treatment modalities is limited to case series and reports. We describe a case of incidentally discovered primary cardiac tumour and present a brief review on epidemiology, pathophysiology, clinical features, and diagnosis and treatment options.

Case
A 61 years old woman with no notable past medical history presented with per-rectal bleeding and weight loss. She denied any other symptoms including chest pain, shortness of breath, palpitations and syncope. Initial routine blood tests and ECG were within normal range. A CT scan of thorax, abdomen and pelvis was performed, mainly to rule out a GI malignancy. The scan showed an incidental 3.5 × 3 cm large left atrial mass attached to the inter-atrial septum (IAS) (Figure 1). No extra cardiac lesions or lymphadenopathy were present on the scan. A transthoracic echocardiogram was carried out urgently that revealed a large mobile mass arising from IAS, abutting the mitral valve in systole (Figure 2). Systolic and diastolic functions of the heart were preserved. Cardiac catheterisation showed normal coronary arteries. Diagnosis of a primary cardiac tumour was made, and she was urgently referred to cardiothoracic centre for surgical resection. The tumour was excised in its entirety using a median sternotomy approach under general anaesthesia following which she made uneventful recovery. The excised mass was confirmed to be a myxoma on histology.

Discussion
In adults, over 75% of primary cardiac tumours are benign. Myxomas are the commonest cardiac tumours, followed by lipomas, fibromas, haemangiomas and neurofibromas. Highest incident of these tumours is reported between third to sixth decade of life and most of these occur sporadically, although familial cases have been described in the literature too [3]. Majority of Myxomas arise from the inter-atrial septum and are found in the left atrium. The symptoms correlate poorly with the histological type of the tumour and mainly depend upon the size, consistency and the site of tumour. Clinical features can be broadly classified into three categories: embolization, intra-cardiac obstruction and constitutional features. Approximately, one third of the myxomas
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Dyspnoea, syncope and sudden cardiac death are the manifestations of intra-cardiac obstruction arising from these tumours. Transthoracic echocardiogram is a useful bedside investigation for initial assessment. Trans-oesophageal echocardiogram and cardiac MRI scan can help to further characterise these lesions by offering useful information about their consistency and anatomical relationship [5]. Coronary angiogram is reserved for those who are suspected to have concurrent coronary artery disease. Surgical resection in the only available effective treatment for primary cardiac tumours and should be performed promptly after the diagnosis is made in order to avoid complications [6].

Conclusion

Primary cardiac tumours are very rare and usually present with symptoms due to heart failure, embolization, arrhythmias and syncope. When found incidentally, they carry significant implications for further investigations and treatment. Untreated cardiac tumours can lead to serious complications including systemic embolization, stroke and sudden cardiac death.

References

2. Reynen K (1996) Frequency of primary tumours of the heart. Am J Cardiol 77: 107. [Crossref]