Atrial Myxoma Presenting with Palpitations: A Case Report

George S. Prousi ¹, Joseph V. Moran ¹, Ross G. Biggs ²

1. Internal Medicine, Lehigh Valley Health Network, Allentown, USA 2. Cardiovascular Disease, Lehigh Valley Health Network, Allentown, USA

DOI: 10.7759/cureus.4093

☑ **Corresponding author:** George S. Prousi, gprousi@gmail.com Disclosures can be found in Additional Information at the end of the article

Abstract

Atrial myxomas are a rare phenomenon and although benign, primary neoplasms of the heart can be burdensome depending on their location. Clinical symptoms are caused through a variety of mechanisms including conduction disturbances, obstruction, and valvular interference. Size and symptom development are strongly correlated and can almost always be detected by the use of echocardiography, magnetic resonance imaging or computed tomography. This is a case of a 62-year-old female with no significant past medical history presented to our facility with complaints of palpitations and associated dizziness for three months.

Categories: Cardiac/Thoracic/Vascular Surgery, Cardiology

Keywords: myxoma, palpitations, transthoracic echocardiography, angiography, mitral valve, atrial septal defect

Introduction

Atrial myxomas are the most common benign primary neoplasm of the heart, accounting for close to 80% of all cardiac tumors [1]. Although quite rare, cardiac tumors in general can present with a multitude of symptoms that are closely correlated with their location [2]. When symptoms are present, echocardiography and other imaging modalities almost always detect a lesion [3]. When found in the left atria, manifestations include conduction disturbances, obstruction, and valvular interference owing to symptoms including dyspnea, orthopnea, cough, edema and fatigue. Given that clinical symptoms overlap with congestive heart failure and other cardiac abnormalities, accurate diagnosis is imperative for appropriate treatment and prognosis.

Received 02/11/2019 Review began 02/15/2019 Review ended 02/17/2019 Published 02/19/2019

© Copyright 2019

Prousi et al. This is an open access article distributed under the terms of the Creative Commons Attribution License CC-BY 3.0., which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Case Presentation

A 62-year-old female with no significant past medical history presented to our facility with complaints of palpitations and associated dizziness for three months. Prior work-up of her palpitations with Holter monitoring showed no irregularities. On arrival, she was in no acute distress and her palpitations had subsided. Vitals that were obtained were largely unremarkable except for a blood pressure of 142/77. Her EKG showed no acute irregularity and laboratory testing was within normal limits. On physical, a regular rate was observed, no murmurs, gallops or rubs were auscultated. She did not exhibit jugular venous distention or peripheral edema and other organ systems did not yield and irregularities.

The patient was admitted for further evaluation and a transthoracic echo was performed, revealing a $4.4\,\mathrm{x}$ 3.0-cm mass in the left atrium attached at the interatrial septum and

How to cite this article

Prousi G S, Moran J V, Biggs R G (February 19, 2019) Atrial Myxoma Presenting with Palpitations: A Case Report . Cureus 11(2): e4093. DOI 10.7759/cureus.4093

aortomitral intervalvular fibrosa. Additional imaging studies including cardiac magnetic resonance imaging (MRI) and transesophageal echocardiography (TEE) were obtained for further confirmation of the mass and its location (Figures 1, 2). Surgical resection was planned, and pre-operative cardiac catheterization was performed which revealed mild prolapse of the mass causing intermittent obstruction of the mitral valve.

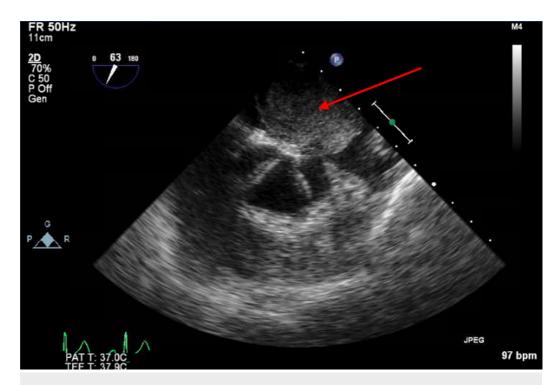


FIGURE 1: Transesophageal echocardiographic image showing left atrial mass (red arrow) seated above the aortic valve

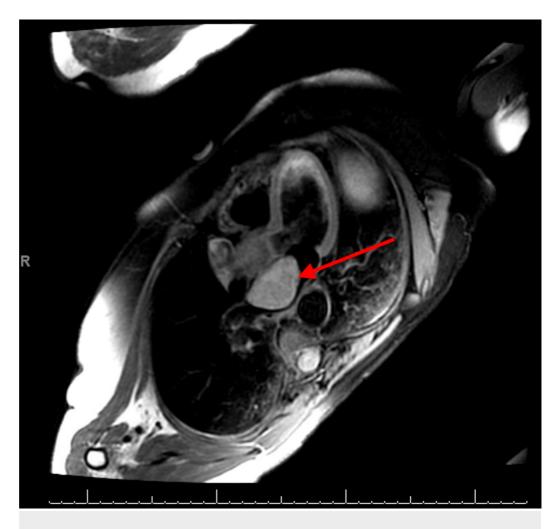


FIGURE 2: Cardiac magnetic resonance image depicting left atrial mass (red arrow)

The patient underwent full-thickness resection of the mass, resulting in an iatrogenic atrial septal defect which was closed with a Dacron patch. Her postoperative course was complicated by the development of intermittent complete heart block with junctional bradycardia and subsequent atrial fibrillation. She was evaluated by electrophysiology and a permanent pacemaker was implanted without complication. Discharge follow-up several weeks later noted the resolution of her prior symptoms, and repeat imaging showed no evidence of mass recurrence or mitral valve prolapse.

Discussion

Tumors of the heart commonly present with symptoms that correlate with the location, the degree of obstruction, and interference with heart valves or circulation. Clinical symptoms demonstrated in the literature include chest pain, dyspnea, and syncope but infrequently present with palpitations for an extended period of time [4-7]. Our patient had no past medical history and prior workup from an electrophysiological standpoint was within normal limits including EKG analysis and Holter monitoring. With the persistence of symptoms, echocardiography was obtained demonstrating a left atrial mass with further analysis showing intermittent mitral valve obstruction and resultant regurgitation. At present, evaluation of cardiac tumors aimed to determine size and location which is pertinent for further evaluation and eventual management. Given its wide availability, echocardiography continues to be the

mainstay of the initial evaluation, whereas CT and MRI are used to provide additional details by offering insight as to the type of tumor [8-10].

When the diagnosis is confirmed, prompt resection is suggested to avoid the possibility of further manifestations including embolization or sudden cardiac death [11-12]. Outcomes are largely favorable with a mortality rate under 5% [11-13]; however, post-operative development of arrhythmias and atrioventricular conduction abnormalities have been reported in approximately 25% of patients [12]. Recurrence is rare although possible and is strongly correlated with a family history of myxoma development [13].

Conclusions

This patient presented in an atypical fashion which lacked commonly associated symptoms including chest pain, dyspnea on exertion, and syncope. Imaging modalities including echocardiography, CT, and cardiac MRI are considered equally efficacious for the diagnosis of atrial tumors as they provide the location within the heart. Additional studies including angiography and biopsy are commonly required for further evaluation when surgery is planned. Surgical intervention consists of tumor resection and despite a low mortality rate, complications including arrhythmias and atrioventricular conduction delays are possible.

Additional Information

Disclosures

Human subjects: Consent was obtained by all participants in this study. **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following: **Payment/services info:** All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

- 1. Lam KY, Dickens P, Chan AC: Tumors of the heart. A 20-year experience with a review of 12,485 consecutive autopsies. Arch Pathol Lab Med. 1993, 117:1027.
- 2. Silvestri F, Bussani R, Pavletic N, Mannoe T: Metastases of the heart and pericardium. G Ital Cardiol. 1997, 27:1252.
- 3. Vander Salm TJ: Unusual primary tumors of the heart . Semin Thorac Cardiovascular Surg. 2000, 12:89.
- 4. Constantine G, Shan K, Flamm SD, Sivananthan MU: Role of MRI in clinical cardiology . Lancet. 2004, 363:2162. 10.1016/S0140-6736(04)16509-4
- 5. Gilkeson RC, Chiles C: MR evaluation of cardiac and pericardial malignancy . Magn Reson Imaging Clin N Am. 2003, 11:73.
- 6. Kaminaga T, Takeshita T, Kimura I: Role of magnetic resonance imaging for evaluation of tumors in the cardiac region. Eur Radiol. 2003, 13:
- 7. Gulati G, Sharma S, Kothari SS, Juneja R, Saxena A, Talwar KK: Comparison of echo and MRI in the imaging evaluation of intracardiac masses. Cardiovasc Intervent Radiol. 2004, 27:459. 10.1007/s00270-004-0123-4
- 8. Jelic J, Milici D, Alfirevi I, et al.: Cardiac myxoma: diagnostic approach, surgical treatment and follow-up. J Cardiovasc Surg. 1996, 13-6:113-117.
- 9. Keeling IM, Oberwalder P, Anelli-Monti M, et al.: Cardiac myxomas: 24 years of experience in 49 patients. Eur J Cardiothorac Surg. 2002, 22:971.
- 10. Selkane C, Amahzoune B, Chavanis N, Raisky O, Robin J, Ninet J, Obadia JF: Changing management of cardiac myxoma based on a series of 40 cases with long-term follow-up. Ann

- Thorac Surg. 2003, 76:1935.
- 11. Bhan A, Mehrotra R, Choudhary SK, et al.: Surgical experience with intracardiac myxomas: long-term follow-up. Ann Thorac Surg. 1998, 66:810.
- 12. Bakaeen FG, Reardon MJ, Coselli JS, et al.: Surgical outcome in 85 patients with primary cardiac tumors. Am J Surg. 2003, 186:641.
- 13. Pinede L, Duhaut P, Loire R: Clinical presentation of left atrial cardiac myxoma. A series of 112 consecutive cases. Medicine (Baltimore). 2001, 80:159.