

Large atrial myxoma mimicking severe mitral stenosis associated with right heart enlargement and severe pulmonary hypertension

Sunnar Leo, Kan Yang, Chunyan Weng, Zhongshu Liang

Department of Cardiology, Third Xiangya Hospital, Central South University, Changsha, Hunan 410013, P.R. China

Corresponding to: Kan Yang, MD, PhD, Professor. Tongzipo 172, Third Xiangya Hospital of Central South University, Changsha, Hunan 410013, P.R. China. Email: yangkanxy3@gmail.com.



Submitted Jan 22, 2013. Accepted for publication Feb 20, 2013.

doi: 10.3978/j.issn.2223-3652.2013.02.02

Scan to your mobile device or view this article at: <http://www.thecdt.org/article/view/1545/2260>

Introduction

Cardiac myxomas are rare tumors of the heart and are easily misdiagnosed. We report a case of large left atrial myxoma with clinical presentation mimicking severe mitral stenosis.

Case report

A 58 year-old woman was referred after complaining of dyspnoea and chest pain on exertion for 3 months. Dyspnoea had progressed to orthopnea and paroxysmal nocturnal dyspnoea. She denied experiencing fever, weight loss, or syncope. Blood pressure was 110/70 mmHg with a pulse rate of 79 bpm.

Cardiac auscultation revealed a loud S1 with a high-pitched, grade III/VI systolic murmur at the cardiac apex, which was thought to be a sign of significant mitral regurgitation, commonly found in congestive heart failure. An additional low-pitched, grade III/VI rumbling diastolic murmur suggested mitral stenosis. Mild oedema was found in lower extremities.

Brain Natriuretic Peptide (BNP) was increased to a level of 5,613.42 pg/mL. Creatinine Kinase MB was slightly increased but Cardiac Troponin I was normal. ECG showed sinus rhythm with 0.1 mV ST segment depression in lead II, III, and AVF.

Chest radiograph revealed cardiomegaly. Transthoracic echocardiography was notable for enlargement of both atrium and the right ventricle with a large (58 mm × 30 mm), elliptical, mobile hyperechoic mass inside left atrium that prolapsed into the left ventricle during diastole. Color Doppler revealed increased velocity of blood flow across the mitral valve, moderate mitral regurgitation

and severe tricuspid regurgitation with pulmonary artery systolic pressure estimation of 88 mmHg. The M-mode echocardiogram of mitral valve in the parasternal long axis showed prolongation of the ejection fraction slope similar to those seen in mitral stenosis (*Figure 1A,B,C*).

The patient was referred for open heart surgery, which was performed with cardio-pulmonary bypass. A large, 6 cm × 5 cm × 4 cm mass was identified inside left atrium, attached via a stalk to the postero-superior side of foramen ovale. The mass was successfully excised, and histopathological examination showed typical features of myxoma (*Figure 1D*).

Postoperatively, the symptoms of heart failure improved and the patient remained free from angina.

Discussion

Myxoma is the most common primary cardiac tumor, and accounts for 83% of primary tumors (1). The age of presentation varies from 1 month to 81 years with predilection of female gender (65%) (2). Most cardiac myxomas are sporadic, but a mutation of chromosome 17 (17q24) in protein kinase A type I-A can lead to Carney complex, a familial cardiac myxoma (3).

Due to a variety of nonspecific findings at first presentation, myxoma is initially suspected in only 5.7% of patients (4). Clinical manifestations can be classified into cardiac (67%), embolic (29%), and systemic (34%) symptoms (5).

Symptoms of left ventricular failure, as in the current case, are typically caused by partial obstruction of the mitral valve orifice by the myxoma. As in our case, myxomas might be initially misdiagnosed as mitral stenosis, but severe mitral stenosis, as seen in our patient has been described in only

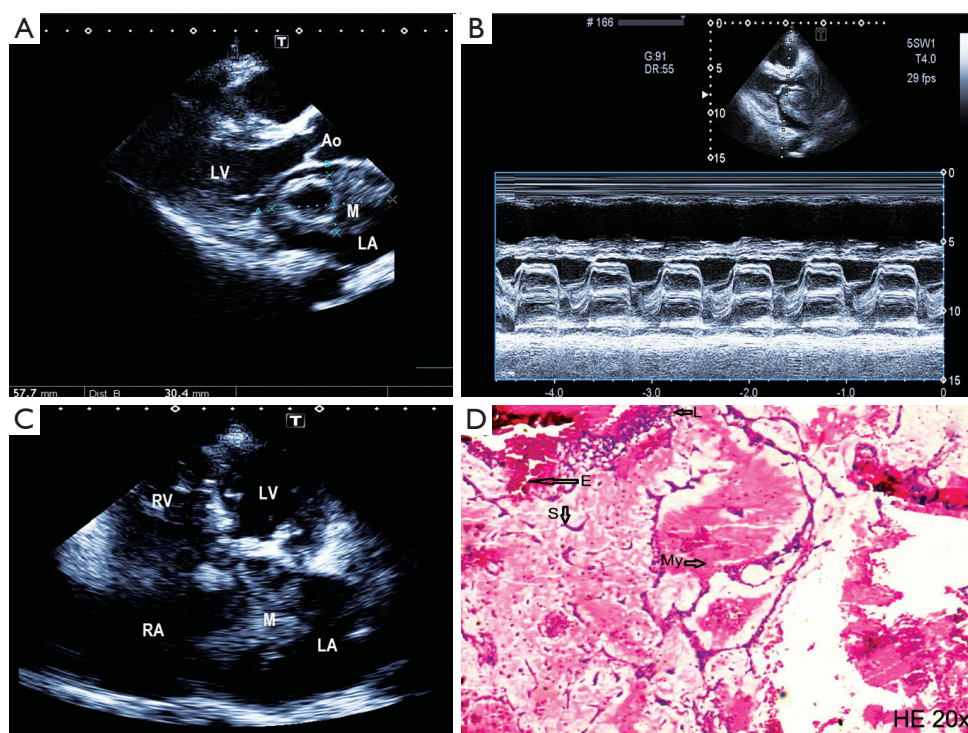


Figure 1 A. Echocardiography showed an elliptical and mobile mass attached to the interatrial septum; B. Prolongation of the ejection fraction slope (M-mode); C. The mass prolapsed into the left ventricle during diastole. Right atrium and ventricle were found enlarged; D. Haematoxylin and eosin stain (20 \times) showed a myxoid matrix with stellate or elongated polygonal (lepidic) cell, scattered lymphocytes and red blood cells. LA = left atrium; LV = left ventricle; Ao = Aorta; RV = right ventricle; RA = right atrium; M = myxoma; My = myxoid degeneration; S = stellate cell; E = erythrocyte; L = lymphocyte

14% of cases (6).

Presentation with chest pain is seen in 28% of cases (7). Typical angina with ischemic changes in electrocardiogram, as noted in our patient might easily misdiagnosed as coronary heart disease. Some authors have described coronary steal phenomenon as the possible underlying mechanism for the chest pain (8).

A diastolic murmur can be found in 64-67% of cases (5,7). 'Tumor plop' sound is more specific for diagnosis but only found in 15% of cases (5). Left atrial enlargement is a common finding in echocardiography but right heart enlargement, as noted in our patient, was found in only 4% of the cases (9).

Myxomas are benign histologically and have excellent long term prognosis with low recurrence rate after surgical resection (5).

In summary, myxomas are easily misdiagnosed and should be considered in the differential diagnosis in patients with suspected mitral valve disease. Echocardiography should be performed as early as possible to establish a

prompt diagnosis and management.

Acknowledgements

Disclosure: All procedures followed were in accordance with the ethical standards of the responsible committee on human experimentation (institutional and national) and with the Declaration of Helsinki, as revised in 2000. For the purpose of publication, written informed-consent has been signed by the patient. This paper has not been published elsewhere in whole or in part. All Authors declared that they have no conflict of interest.

References

1. Blondeau P. Primary cardiac tumors--French studies of 533 cases. *Thorac Cardiovasc Surg* 1990;38 Suppl 2:192-5.
2. Kuon E, Kreplin M, Weiss W, et al. The challenge presented by right atrial myxoma. *Herz* 2004;29:702-9.
3. Yin Z, Kirschner LS. The Carney complex gene PRKAR1A

- plays an essential role in cardiac development and myxomagenesis. *Trends Cardiovasc Med* 2009;19:44-9.
4. Goswami KC, Shrivastava S, Bahl VK, et al. Cardiac myxomas: clinical and echocardiographic profile. *Int J Cardiol* 1998;63:251-9.
 5. Pinede L, Duhaut P, Loire R. Clinical presentation of left atrial cardiac myxoma. A series of 112 consecutive cases. *Medicine (Baltimore)* 2001;80:159-72.
 6. Panidis IP, Mintz GS, McAllister M. Hemodynamic consequences of left atrial myxomas as assessed by Doppler ultrasound. *Am Heart J* 1986;111:927-31.
 7. Aggarwal SK, Barik R, Sarma TC, et al. Clinical presentation and investigation findings in cardiac myxomas: new insights from the developing world. *Am Heart J* 2007;154:1102-7.
 8. Alvarez JR, Quiroga JS, Cereijo JM, et al. Coronary steal by left atrial myxoma: a case report. *Cases J* 2009;2:6741.
 9. Oliveira R, Branco L, Galrinho A, et al. Cardiac myxoma: a 13-year experience in echocardiographic diagnosis. *Rev Port Cardiol* 2010;29:1087-100.

Cite this article as: Leo S, Yang K, Weng C, Liang Z. Large atrial myxoma mimicking severe mitral stenosis associated with right heart enlargement and severe pulmonary hypertension. *Cardiovasc Diagn Ther* 2013;3(1):52-54. doi: 10.3978/j.issn.2223-3652.2013.02.02