A rare case of mobile ball-like mass in right atrium associated with dilated cardiomyopathy

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Abstract
Mobile thrombus in the Right Atrium (RA) is a rare condition. Here, we described a case of a 59-year-old male who presented in emergency room with complaints of progressively worsening dyspnea for 7 years. He was normotensive, had elevated jugular venous pressure and clear lungs. ECG revealed ventricular tachycardia. After successfully electrical defibrillation, an urgent 2D echocardiogram was obtained and revealed the enlargement of all heart chambers and a ball-like, mobile mass in the right atrium. An emergent CT scan revealed no sign of pulmonary embolism. Based on morphological features on echocardiogram, we highly suspected it as RA thrombus. The patient was treated with anticoagulant therapy comprising of low molecular heparin and warfarin since he rejected surgical treatment. Repeat echocardiogram was performed 7 days after anticoagulant therapy and showed disappearance of the RA mass. He was subsequently maintained on warfarin (INR=2.4-3.0) and other medications for heart failure. The patient was followed up as outpatient and kept in good clinical condition. We concluded that early recognition of this rare situation and prompt management can help in preventing life-threatening complications, and anticoagulant therapy was a therapeutic option for RA thrombus.

Keywords: right atrial mass; thrombus; echocardiogram; anticoagulation.

Introduction
Although intracardiac masses are commonly encountered in clinical setting, a mass in the Right Atrial (RA) cavity is relatively rare. The masses usually include thrombi, vegetations, metastatic tumors, myxomas, inflammatory granuloma, etc [1,2]. It’s very challenging to differentiate the nature of masses therefore provide appropriate management to prevent the possible complications. The definite diagnosis can be only established by surgical resection and pathology, however certain morphological features on imaging studies can help in differential diagnosis. We hereby describe a case of a mobile ball-like RA thrombus diagnosed with Transthoracic Echocardiography (TTE) and the patient was treated successfully with anticoagulation therapy without any complications.

Case presentation
A 59-year-old male presented in emergency room with complaints of progressive dyspnea for 7 years and worsening for 3 days. His medical history was significant for hypertension but well-controlled with anti-hypertensive drugs. Denied any known liver, kidney diseases and diabetes. On physical examination, he was normotensive, had elevated jugular venous pressure and clear lungs. Emergency ECG revealed ventricular tachycardia (HR=135bpm) without ST-T changes. Electrical defibrillation was performed and restored as sinus rhythm (HR=95 bpm). Pitting lower limb edema was noticed. His blood analysis was as follows: hemoglobin 13.5 g/dl, leukocyte count 7.9×10⁹/uL, platelet count 59,000/uL, high-sensitivity troponin protein 0.303ng/ml, CK-MB 7.3ng/ml, Lactate dehydrogenase (LDH) 10750U/L,
NT-proBNP 22702pg/ml, serum creatinine 438.1mmol/L, BUN 22.54mmol/L, eGFR=12.12ml/min*1.73m². Liver biochemistry tests were as followings: Alanine aminotransferase (ALT) 3518 U/L, aspartate Aminotransferase (AST) 4267 U/L, cholinesterase 3406.7 U/L, HBsAg was negative. Blood gas analysis: PH=7.072, PO₉ 99.70 mmHg, PCO₂ 19.60 mmHg, actual bicarbonate 5.6 mmol/L, standard bicarbonate 8.6 mmol/L. Coagulation tests: PT 57.1s, APTT 58.0s, D-Dimer 7.97 µg/ml, FDP 94.5 µg/ml. Serum potassium 6.6 mmol/L, serum sodium 130.7 mmol/L, serum chloride 89.4 mmol/L. An urgent 2D echocardiogram was obtained and revealed the enlargement of all heart chambers (left ventricular diastolic diameter 75 mm). Left ventricular ejection fraction (LV EF) 25%. A mobile ball-like mass was noticed in the right atrium (Figure 1). An emergent CT scan was ordered and revealed no sign of pulmonary emboli. Ultrasound-Doppler didn’t detect any sign of bilateral lower limb Deep Venous Thrombus (DVT). Patient hospitalized and received treatment for heart failure. Since the high risk of pulmonary embolism, surgical management of RA mass was recommended but rejected by the patient. Given the morphological features of this mobile RA mass, we highly suspected it as RA thrombus. The patient was then treated with anticoagulant therapy consisting of low molecular heparin and warfarin after obtaining written informed consent from the patient. Repeat TTE was performed 7 days later and showed disappearance of the mobile RA mass, suggesting complete thrombus resolution (Figure 2). He was subsequently maintained on warfarin (maintaining INR 2.4~3.0) and other medications for heart failure with significant improvement of symptoms. The patient did well during the follow-up period.

Written consent was obtained from the patient prior to disclose his information and images. The case report was conducted in accordance with the principles of the Declaration of Helsinki.

**Figure 1:** Transthoracic two-dimensional echocardiogram showing a mobile ball-like thrombus in the right atrium; LV: Left Ventricle; RA: Right Atrium; RV: Right Ventricle; A: Apical four chamber views; B: Subcostal view.

**Figure 2:** Transthoracic two-dimensional echocardiogram performed at 7 days after anticoagulation therapy, showing no residual thrombus in the right atrium; LV: Left Ventricle; RA: Right Atrium; RV: Right Ventricle; A: Apical four chamber views; B: Subcostal view.

**Discussion**

Atrial thrombi are more commonly seen in left atrium (including left atrial appendage) and left ventricle, and usually accompanied by structural heart diseases or other predisposing conditions, such as acute myocardial infarction, left ventricular aneurysm, cardiomyopathies and myocarditis, valve disease, atrial fibrillation/atrial flutter [3-5]. Right atrial thrombi are relatively rare but if present, there are significant morbidity and mortality due to massive pulmonary embolism or paradoxical embolism [6,7]. The differential diagnosis of right atrial thrombi includes tumors (most are metastatic tumors), myxoma, vegetations, inflammatory granuloma, et al [1,2,8,9]. 2D echocardiography (TTE or Transesophageal Echocardiogram (TEE)) is the most easy-use and feasible tool for detection of RA masses in clinical setting, which can provide the essential information about the mass including morphology, echogenicity, mobility, with/without stalk. However, sometimes it is difficult to differentiate one mass from another. In our case, the major differentiation is myoma, they usually have similar echocardiographic features. Given the RA mass appears as a mobile, ball-like, medium echogenicity and no stalk in this case, we highly recommended the diagnosis of RA thrombus, taking consideration with the patient’s history.

The early and accurate diagnosis of RA thrombi is very important because of high risk of pulmonary embolism. Study has indicated that it is associated with increased mortality of more than 40% if mobile RA thrombus is present [7]. Current therapeutic options for RA thrombi include thrombolysis, surgical embolectomy or percutaneous embolectomy, but the most effective therapy remains unknown. In our case, we promptly started anticoagulation therapy with low molecular heparin and warfarin since the patient rejected surgical treatment. The patient was monitored during anticoagulation therapy. The RA mass was found to disappear at 7 days after treatment without any complications, which indirectly confirmed our diagnosis. To our knowledge, this is the first report of successful anticoagulation therapy with warfarin for RA thrombus in such short treatment period. Based on the case described in the present report, anticoagulation therapy can be one of reasonable options for the treatment of RA thrombi. Our conclusion is that early recognition of RA thrombus is very important due to extreme instability of mobile thrombus and prompt treatment can prevent life-threatening complications. Anticoagulation therapy is a reasonable therapeutic option.

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References

1. Felner JM, Churchwell AL, Murphy DA. Right atrial thromboem-
boli: Clinical, echocardiographic and pathophysiologic manifes-
6. Basarici I, Yilmaz H, Demir I, et al. Imminent pulmonary embo-

7. Chapoutot L, Tassigny C, Nazeyrollas P, et al. Pulmonary embo-

9. Martha Alehli Rangel-Herna´ndez, Alberto Aranda-Fraustro, Ga-

briela Melendez-Ramirez, Nilda Espi´nola-Zavaleta. Misdiagno-