



CASE REPORT

Multiple left ventricular aneurysms in a young female



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Received 18 November 2014; accepted 2 September 2015

Available online 3 February 2016

KEYWORDS

Chest radiograph;
Coronary angiogram;
Cardiac
catheterization;
Ventriculogram;
Cardiac magnetic
resonance imaging;
Aneurysm

Abstract Multiple left ventricular aneurysms (LVAs) are rare, especially in a young female. A 29-year-old woman presented vague symptoms. Multiple LVAs were revealed and confirmed on different imaging modalities, including chest radiography, echocardiography, contrast ventriculography and cardiac magnetic resonance imaging. Detailed work-up for probable etiologies including ischemic, infectious, inflammatory and autoimmune causes was negative. In the absence of angina, decompensated congestive heart failure, arrhythmias and embolism, the patient was managed conservatively, with excellent mid-term outcome.

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PALAVRAS-CHAVE

Radiografia de tórax;
Angiografia
coronária;
Cateterismo
cardíaco;
Ecografia ventricular;
Ressonância
magnética cardíaca;
Aneurisma

Múltiplos aneurismas do ventrículo esquerdo numa jovem

Resumo Múltiplos aneurismas no ventrículo esquerdo (MAVE) são uma situação rara em especial numa jovem que apresentou queixas vagas. Vários MAVE foram revelados e confirmados em diferentes modalidades de imagem. A doente submeteu-se a radiografia de tórax, ecocardiografia, ecografia de contraste e ressonância magnética cardíaca. A investigação minuciosa de etiologias isquémica, infecciosa, inflamatória e autoimune foi negativa. Dada a ausência de angina, insuficiência cardíaca congestiva, arritmias e embolia, a doente foi tratada de forma conservadora apresentando um excelente prognóstico a médio prazo.

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Introduction

Left ventricular aneurysm (LVA) is usually defined as a segment of the ventricular wall that exhibits paradoxical systolic expansion; the term is generally reserved for a discrete, dyskinetic area of the left ventricular (LV) wall with a broad neck.^{1,2} It most frequently develops after myocardial infarction (MI).³ However, other causes may be infectious, inflammatory, metabolic or autoimmune in nature. Multiple idiopathic LVAs are distinctly unusual and rarely reported. The clinical spectrum ranges from asymptomatic to congestive heart failure (CHF) to life-threatening ventricular arrhythmias and cardiac arrest.^{3,4} Herein, we report the case of a young female with multiple large LVAs, managed conservatively in the absence of overt CHF, arrhythmias and embolism, with excellent outcome.

Case report

A 29-year-old woman presented with easy fatigability and chest discomfort for several months. These symptoms were insidious in onset and progressed little. There was no history of palpitations, syncope or dyspnea. She was hemodynamically stable with diffuse apex beat shifted downward and outward. The chest radiograph showed calcified LV apical and basal aneurysms. The electrocardiogram showed no pathological Q waves and normal precordial R-wave progression (Figure 1). Two-dimensional transthoracic echocardiography (iE33 xMATRIX, Philips Healthcare, Andover, MA, USA) showed moderate LV dysfunction; multiple LVAs (large apical aneurysm, moderate sized submitral aneurysm, and basal anterior aneurysm); normal contractility of the rest of the LV wall; and normal valves, other chambers and great vessels (Figure 2). The echocardiogram showed continuity of the myocardium of the aneurysms with the rest of the myocardium. Color flow Doppler showed flow in and out of the aneurysms.

The coronary angiogram (Philips Medical Systems, Nederland B.V.) showed normal epicardial coronaries. Catheterization showed LV pressure of 114/3 mmHg. Systemic and pulmonary arterial, right atrial and ventricular pressures were also normal. The left ventriculogram showed a large LV apical aneurysm with a heavily calcified thrombosed distal part, a large aneurysm of the basal anterior wall with a smooth and calcified distal part, and a moderate sized submitral aneurysm. All the aneurysms had large necks, poor contractility and paradoxical systolic filling. Contrast clearance was slightly delayed by a beat (Figure 3).

To further evaluate LV function and anatomy and the shape of the aneurysms and to differentiate between true and pseudoaneurysms, magnetic resonance imaging (MRI) (Achieva 1.5 T system, Philips Healthcare, Andover, USA) was performed. Using a dedicated cardiac coil, images were acquired with electrocardiographic gating and balanced turbo field echo/fast field echo in short-axis, vertical long-axis, 4-chamber and transverse planes, followed by perfusion studies and delayed contrast-enhanced imaging. These showed LV ejection fraction of 39.5% and stroke volume of 55.4 ml. Images were analyzed for the location of aneurysms, maximal internal width of the orifice, and maximal parallel internal diameter. The largest outpouching on the anterolateral wall of the LV showed thinning of the overlying myocardium (thickness 3.5 mm). The neck and maximum diameter were both 24 mm and the ratio of maximal internal width of the orifice and maximal parallel internal diameter was 1.0. There was no delayed pericardial enhancement. All these findings suggested that this was a true aneurysm. Other aneurysms on the anterior wall and submitral area were smaller and had similar characteristics (Figure 4). MRI ruled out inflammatory and infiltrative disorders.

The patient denied a history of rheumatic or systemic diseases, toxin exposure or chest trauma. There was no involvement of skin, mucosa or other organ systems. Family history was insignificant for CHF, cardiomyopathy,

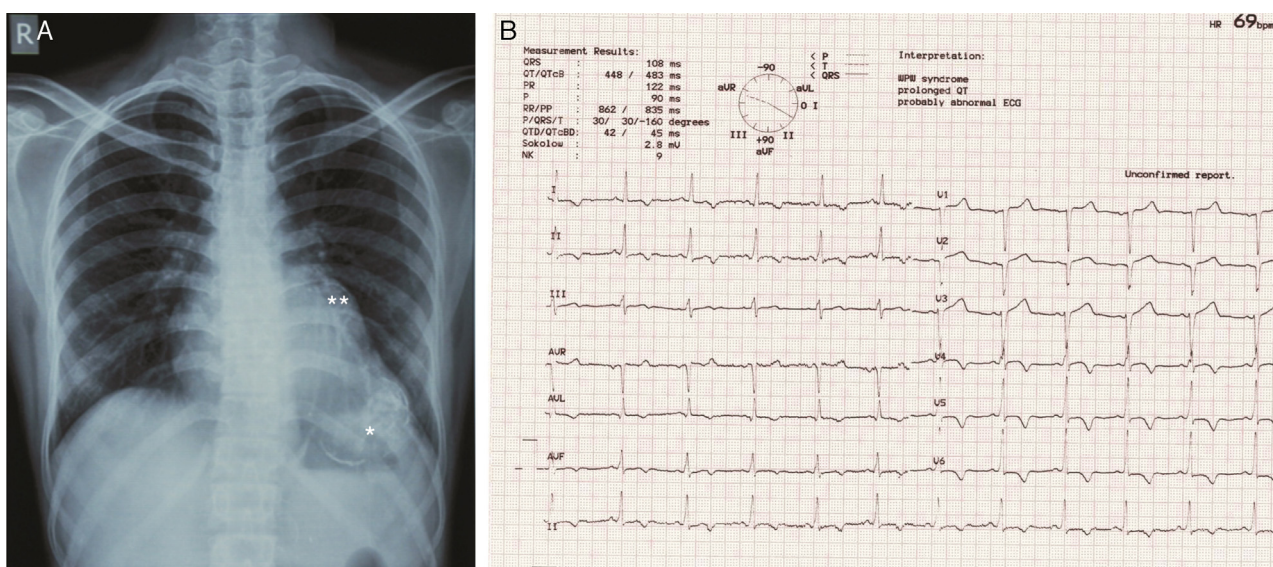


Figure 1 (A) Chest radiograph in posteroanterior view showing large calcified left ventricular apical aneurysm (*) and basal aneurysm (**); (B) electrocardiogram showing no pathological Q waves and normal progression of precordial R wave.

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