

Giant atria in a patient with systemic lupus erythematosus

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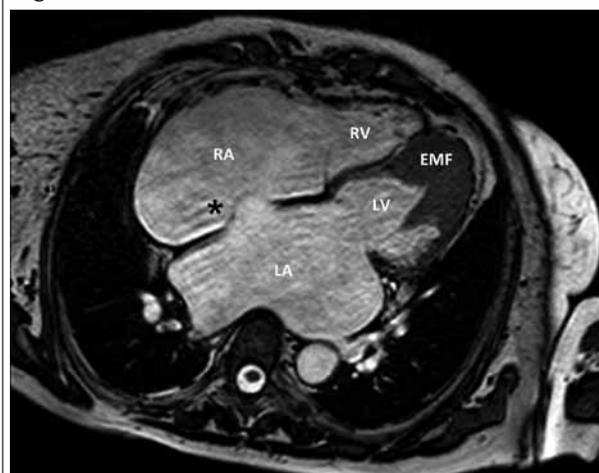
Cardiac disease is common among patients with systemic lupus erythematosus (SLE) and includes endomyocardial fibrosis (EMF).¹ However, the far-reaching consequences due to diastolic dysfunction are relatively unknown. Early cardiac evaluation and subsequent treatment in SLE patients may improve outcome.²⁻⁴

A 56-year-old Caucasian woman with SLE presented with atrial fibrillation (AF) and symptoms of progressive heart failure. Her chest X-ray revealed severe cardiomegaly and an obtuse carinal angle. The electrocardiogram showed AF and inverted T waves in the precordial leads. Cardiovascular magnetic resonance imaging (*figure 1*)

revealed massive biatrial enlargement with intra-atrial septal aneurysm, obliteration of the apex, decreased LV ejection fraction (0.37) and severe tricuspid regurgitation. The late gadolinium enhanced images were non-diagnostic due to arrhythmia artifacts.

Long-term treatment included conventional heart failure therapy, anticoagulation and adequate rate control for AF. Because of refractory congestive heart failure symptoms, she was operated on. EMF was confirmed and a tricuspid valvuloplasty, atrial reduction and manual dissection of the obliterated ventricle was performed. Afterwards her cardiac condition stabilised for three years. Unexpectedly, she recently died because of a complicated infection as a consequence of long-term immunosuppressive therapy.

Figure 1. MRI scan



LA = left atrium; RA = right atrium; LV = left ventricle; RV = right ventricle; EMF = endomyocardial fibrosis; (*) = intra-atrial septal aneurysm.

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