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A 59-year-old woman with visible precordial pulsations

CLINICAL INTRODUCTION

A retired 59-year-old woman presented to the cardiology clinic concerned with cardiac pulsations that were visible on her chest wall. These were not associated with dyspnoea, syncope or chest discomfort.

Of note, 8 years previously, she complained of recurrent nocturnal diaphoresis and 5 kg weight loss. Blood sampling at that time revealed a microcytic anaemia, reactive thrombocytosis and raised inflammatory markers (erythrocyte sedimentation rate 99 mm/hour, C-reactive protein 161 mg/L). Following an episode of transient diplopia, ophthalmoscopy demonstrated

a cotton wool spot in the left inferotemporal retinal arcade. She commenced a 2-year tapering course of 1 mg/kg prednisolone.

On examination, she had a lean physique with a supine blood pressure of 162/60 mm Hg and palpable Corrigan's pulse. She had a prominent apical pulsation and a loud early diastolic murmur was present at the left sternal edge radiating to the apex. Echocardiography showed severe central aortic regurgitation and a dilated aortic root (see online supplementary figure 1). Cardiac CT was performed to clarify the diagnosis (figure 1).

QUESTION

Which of the following diagnoses best explains this presentation?

- A. Ankylosing spondylitis
- B. Takayasu arteritis
- C. Salmonellosis
- D. IgG4-related aortitis
- E. Giant cell aortitis

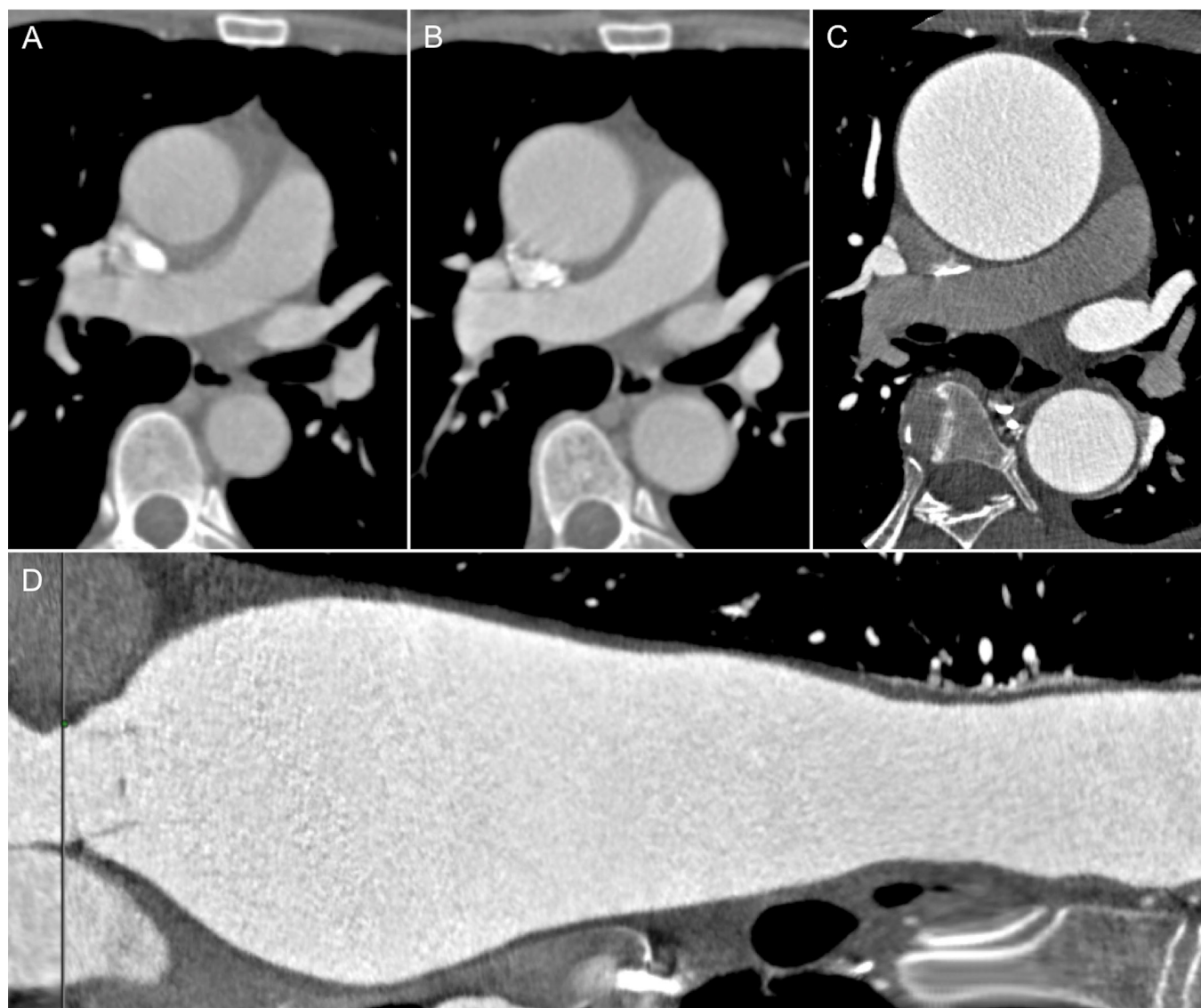


Figure 1 Contrast-enhanced CT of the thorax at index presentation (A) and 6 months (B). Prospective ECG-gated cardiac CT angiogram (75% R-R interval) performed at 8 years from index presentation (C) with a stretched multiplanar reconstruction of the aortic annulus, aortic root and thoracic aorta (D).

Image challenge

CORRECT ANSWER: E

The correct answer is giant cell aortitis. Progressive aortic ectasia is noted on serial imaging. The ECG-gated CT aortogram demonstrates an aortic annulus within normal limits, but there is severe dilatation of the aortic root. The proximal ascending aorta is severely aneurysmal (maximum diameter 77.5 mm). The aortic arch and descending thoracic aorta show circumferential 4 mm intramural thickening in the absence of atheroma and with contrast enhancement of the adventitia (figure 2). The history of polymyalgia rheumatic and the CT pattern of annuloaortic ectasia in the presence of intramural thickening (≥ 2 mm) with adventitial contrast enhancement are compatible with giant cell aortitis.¹

Takayasu arteritis typically manifests as concentric aortic wall thickening with a 'double ring' appearance on contrast CT, due to inflammation of the adventitia. A later occlusive stage is often characterised by branch vessel stenoses.² Seronegative arthropathies are associated with subvalvular and cusp thickening, which are not present in the case.³ Infections of the aorta are rare, however, *Salmonella* aortitis can occur in immunocompromised patients with coexistent osteomyelitis and an atherosclerotic aortopathy which are not features of this case. IgG4-related disease is a fibroinflammatory condition associated with lymphoplasmacytic infiltrates and retroperitoneal fibrosis.⁴

The patient underwent valve-sparing surgery with replacement of the aortic root, ascending aorta and hemiarch (see online supplementary figure 2). Aortitis is an important diagnosis to consider in the management of suspected vasculitis syndromes.

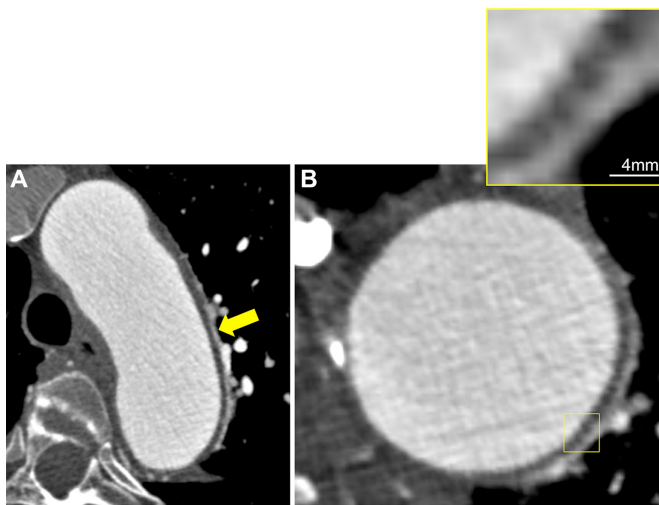


Figure 2 ECG-gated cardiac CT angiogram reconstructions of the (A) aortic arch and (B) descending thoracic aorta. Circumferential intramural thickening (4 mm) with adventitial contrast enhancement of the descending thoracic aorta (inset).

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