Takayasu’s arteritis and an elevated antistreptolysin O titre – a potentially expensive diagnostic conundrum

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Lesson
Takayasu’s arteritis is a chronic large vessel vasculitis which may be associated with a false positive antistreptolysin O titre.

Keywords
Antistreptolysin O Titre, Takayasu’s arteritis

Case report
A 19-year-old Caucasian woman was referred with a four-month history of exertional chest tightness radiating to the left arm. Occasionally this had been associated with palpitations and dizziness. She took no medications, was a non-smoker and consumed moderate alcohol. There was no history of illicit drug use. Her grandfather had suffered from ischaemic heart disease.

Physical examination yielded blood pressure 114/60 mmHg, a regular pulse, no pulse deficits, normal heart sounds and murmurs consistent with mixed aortic valve disease. The jugular venous pressure was normal with clear lung fields. There was no evidence of arthritis or rashes.

A resting 12-lead electrocardiogram showed sinus rhythm meeting voltage criteria for left ventricular hypertrophy. A subsequent exercise tolerance test was electrically and symptomatically positive with significant widespread ST-segment depression.

Trans-thoracic echocardiography revealed a dilated ascending aorta with severe aortic regurgitation. Magnetic resonance angiography confirmed a dilated aortic root with significant mural thickening from the aortic root to the arch.

Laboratory investigations revealed a markedly elevated erythrocyte sedimentation rate (ESR) 92 mm (at only 25 min), C-reactive protein (CRP) 48 mg/L, WCC 11.0 x 10⁹/L, IgG 24.27 g/L and antistreptolysin O titre (ASOT) 3840 U/mL. Urea and electrolytes, liver function tests, anti-nuclear antibody, anti-neutrophil cytoplasmic antibody, C3 and C4 were normal. HIV, syphilis and toxoplasma gondii antibodies were negative. Serology was consistent with previous Epstein–Barr virus and cytomegalovirus infection.

18F-FDG PET/CT revealed increased metabolic activity in the wall of the aorta extending from root to arch, showing an active large vessel arteritis consistent with Takayasu’s arteritis (TA) (Figure 1). The patient was treated with oral Prednisolone and Mycophenolate Mofetil. Concurrent intravenous Benzylpenicillin was given to cover the possibility of rheumatic fever. A repeat ASOT was persistently elevated but additional serum testing for anti-DNase B and Streptococcal emm gene was negative, whereupon Benzylpenicillin was stopped.

After two weeks of immunosuppressant therapy, there was a significant fall in inflammatory markers, with ESR 10 mm/h and CRP <4 mg/L. Subsequent coronary angiography and aortography revealed critical left main stem ostial stenosis and severe aortic regurgitation.

Following referral to a quaternary cardiothoracic centre, the patient underwent aortic root and mechanical aortic valve replacement, left main stem arterioplasty with re-implantation and precautionary coronary artery bypass grafting. Histopathology of the aortic root supported the clinical and radiological diagnosis of TA.

Discussion
TA is a chronic large vessel vasculitis affecting the aorta and its main branches. It is a rare disease in the UK, with a reported mean prevalence of 4.7 per million in primary care and 7.1 per million in a secondary care setting. A study from Italy suggests coronary artery involvement is an uncommon finding in the West.

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There are many causes of aortitis (Table 1) and a significantly elevated ASOT in our patient raised the possibility of rheumatic fever, creating diagnostic uncertainty. Before continuing with immunosuppressant therapy alone for suspected active large vessel vasculitis, it was therefore important to exclude group A Streptococcal infection.

Serum antibody testing against more than one group A Streptococcal antigen increases the diagnostic sensitivity for Streptococcal infection. DNA amplification of the Streptococcal emm gene can also provide evidence of Streptococcal infection. The combined negative results for anti-DNase B antibody and serum emm gene in our patient confirmed the
elevated ASOT was a false positive result due to cross reactivity. Current in-house testing for ASOT costs approximately £8 and testing for anti-DNAse B and emm gene, £45 and £136, respectively at the UK national reference laboratory. By contrast, a 10 day course of intravenous benzylpenicillin 1.2 g qds costs £75 and prophylactic oral penicillin V 250 mg bd until the age of 40 costs £400.7 Although additional testing for Streptococcal infection carries a cost, negative results remain cheaper than empiric long-term treatment.

The association between an elevated ASOT and TA has been described previously, with 19% of patients returning an ASOT twice the upper limit of normal in one study.8 However, searching PubMed with simple terms (antistreptolysin O antibody and Takayasu) does not yield relevant results regarding this association. We feel it is important to reiterate the potential for false positive ASOT in the context of TA, as misinterpretation could lead to incorrect long-term treatment.

At six months following operation, the patient remained in remission from vasculitis, with normal inflammatory markers and only residual uptake on PET/CT related to postoperative changes.

Declarations

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Guarantor: OL

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