Extensive aortic valve ring abscess formation: a rare complication of Q fever endocarditis

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Summary: We report the successful management and 2 year follow up of a young patient with Q fever endocarditis on a congenitally bicuspid aortic valve complicated by extensive abscess formation in the aortic valve ring and interventricular septum. Aortic root abscess formation complicating Q fever endocarditis has been reported in only one previous patient. Serological tests may thus be indicated in patients with aortic abscesses.

Despite extensive aortic and intramyocardial abscess formation it proved possible to control the progression of disease by open drainage of the abscess and aortic valve replacement. Although the requirement for aortic root replacement was anticipated in this patient, it has not been required.

Introduction

Q fever is a systemic zoonosis caused by *Coxiella burnetii*. Endocarditis is one of the most important of its many clinical manifestations, and predominantly affects the aortic valve. Prolonged antibiotic treatment remains the first line therapy of Q fever endocarditis, with surgery reserved for severe haemodynamic complications. We report a case of Q fever endocarditis complicated by extensive suppuration affecting the aortic valve ring and myocardium.

Case report

A 21 year old male presented with a 2-week history of lethargy, perspiration, rigors, night sweats, weight loss, palpitations and breathlessness. Twelve months previously his general practitioner had detected a heart murmur and hypertension (160/110 mmHg). Coarctation was considered because of reduced femoral pulses, but aortography demonstrated a bicuspid aortic valve and pseudo-coarctation. There was no evidence of infective endocarditis and the aortic root was normal.

Five months prior to the presentation with endocarditis, the patient suffered a 'flu-like' illness with fever, rigors, pleuritic chest pain and haemoptysis. The chest radiograph demonstrated left basal consolidation, but the causative organism was never identified.

On presentation to the cardiac unit he was pyrexial (39°C) and he had finger clubbing, a single splinter haemorrhage and two Janeway lesions. His blood pressure was 140/70 mmHg. He had a loud ejection systolic murmur, as before, but in addition a new and long early diastolic murmur of aortic regurgitation.

A chest radiograph demonstrated a normal heart size, with no evidence of pulmonary oedema. An electrocardiogram demonstrated first degree heart block (PR interval 0.26 seconds) and right bundle branch block. The serum haemoglobin was 11.5 g/dl with a white cell count of 10.3 x 10⁹/l, and the blood film demonstrated early iron deficiency changes. The erythrocyte sedimentation rate was 38 mm/h. Six sets of blood cultures, a mid-stream specimen of urine, and nasal and throat swabs were all negative. There was no microscopic haematuria. Immunoglobulin G levels were elevated to 24.1 g/l.

On a presumptive diagnosis of infective endocarditis he was treated with high doses of benzyl penicillin, flucloxacillin and gentamicin intravenously, but he remained febrile. Antimicrobial treatment was changed to tetracycline and co-trimoxazole based upon the results of serology. The Phase I and Phase II *Coxiella burnetii* titres were elevated to dilutions of 1:1024 and 1:8192 respectively indicating the diagnosis of Q fever. *Coxiella burnetii* was subsequently isolated from guinea pigs inoculated with material from the excised aortic valve. On detailed enquiry the patient admitted to a single episode of exposure to farm animals.

Echocardiography demonstrated a calcified bicuspid aortic valve with a large vegetation on the right (or anterior) coronary cusp measuring 1.5 x 2.2 cm. There was an adjacent abscess formation in the aortic root which extended into the interventricular septum (see Figures 1 and 2). Doppler echocardiography did...
Figure 1  Parasternal long axis echocardiogram demonstrating abscess in anterior aortic root (arrow) which extends into the upper interventricular septum, and a large vegetation on the right coronary cusp of the aortic valve.

Figure 2  Short axis echocardiogram through the aortic root. An abscess is seen anteriorly (arrow), bulging into the right ventricular outflow tract below the pulmonary valve.
not demonstrate any significant aortic stenosis, but he had considerable aortic regurgitation. The left ventricular dimensions were 6.7 cm and 3.9 cm in diastole and systole respectively, with functional shortening 42%. There was also a moderate pericardial effusion (1 cm). The patient therefore underwent urgent aortic valve replacement. At operation two abscesses were found adjacent to the right coronary cusp; these were spherical and 2.0 cm and 1.5 cm in diameter, and they extended into the interventricular septum, and towards the main pulmonary artery respectively. The abscesses were curetted but left open, and the valve was replaced with a 13A Starr Edwards prosthesis inserted below the abscess cavities.

Following surgery he had complete heart block, for which an A-V sequential pacemaker was inserted. His post-operative course was marked by a swinging pyrexia for 18 days, but no focus of infection was found. Since then he has remained afebrile on treatment with doxycycline and warfarin. At review 24 months after surgery he was well with no evidence of recurrent Q fever or endocarditis.

Discussion

This patient is unusual because he had aortic valve ring abscess formation, with intra-myocardial extension and heart block, complicating Q fever endocarditis. The abscesses were detected pre-operatively by echocardiography, and were the presumptive cause of his first degree AV block and right bundle branch block. Although the need for aortic root replacement was considered it proved possible to control the spread of infection in this man by aortic valve replacement and open drainage of the aortic and intramyocardial abscesses.

Necropsy studies suggest that valve ring abscesses are a frequent complication of infective endocarditis, but pre-operative diagnosis can be difficult. The sensitivity of echocardiography can be improved considerably by oesophageal and intra-operative studies, but even then, when abscess formation is recognized it is usually associated with infections with virulent bacterial organisms such as Streptococcus, Staphylococcus or Pneumococcus. Abscess formation in the aortic root was detected intra-operatively in one patient from a series of 10 patients with Q fever endocarditis. However, valve ring and intramyocardial abscess formation in Q fever endocarditis, to our knowledge, has not been previously described.

Early surgical intervention with extensive excision of infected material is recommended in cases of sepsis involving the aortic valve ring and aortic root. Without this, and in spite of appropriate antibiotic therapy, mortality is high because of continuing infection, aortic regurgitation and heart failure. Since it may be difficult to control infection in Q fever and to prevent subsequent prosthetic valve endocarditis, there is a temptation to postpone valve replacement. In patients with abscesses this would clearly be inappropriate, and the experience of our patient confirms that the result of early surgery in Q fever may be excellent. Antibiotic treatment is recommended for at least 12 months and probably for life, but this has not been the subject of controlled studies.

The clinical features of Q fever are protean, but endocarditis is the most serious complication. This can occur between one and 20 years after the acute infection, and it is associated with a mortality of 31–56%. This report demonstrates that a diagnosis of Q fever endocarditis should be considered in patients with aortic valve ring or aortic root abscess formation and that such patients can be managed by open drainage of the abscess and aortic valve replacement.

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References

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