

CASE STUDY

Staphylococcus capitis endocarditis: two cases and review of the literature

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Abstract
Coagulase negative staphylococci are the principal cause of prosthetic valve endocarditis but are a rare cause of native valve infections. However, the incidence of native valve endocarditis is increasing. Staphylococcus capitis is a coagulase negative staphylococcus with the capacity to cause endocarditis on native heart valves. Two cases of native valve endocarditis caused by S capitis are presented; both in patients with aortic valve disease. The patients were cured with prolonged intravenous vancomycin and rifampicin and did not need surgery during the acute phase of the illness. Five of the six previously described cases of endocarditis caused by this organism occurred on native valves and responded to medical treatment alone. (Heart 1999;82:e1)

Keywords: Staphylococcus capitis; endocarditis; valvar disease; coagulase negative staphylococci

Coagulase negative staphylococci (CoNS) are the most common cause of prosthetic valve endocarditis but are responsible for only 10% of native valve infections. However, the incidence of CoNS native valve infection seems to be increasing. Staphylococcus epidermidis is the species most frequently associated with native valve endocarditis; one recent series reported it as the cause of 91% of cases studied. Nevertheless, endocarditis associated with other species of CoNS is well documented. We report two cases of native valve endocarditis caused by Staphylococcus capitis in patients with aortic valve disease, and review six previously published cases of both native and prosthetic valve infection with particular reference to medical and surgical management and outcome.

Case 1
A 46 year old man presented with acute ischaemia of the right foot. He was in end stage renal failure, secondary to chronic glomerulonephritis, and was being treated with continuous ambulatory peritoneal dialysis. The patient was also known to have mild aortic stenosis and paroxysmal atrial fibrillation. On examination, he had an intermittent low grade fever but was haemodynamically stable (pulse 80 beats/min, sinus rhythm; blood pressure 140/100 mm Hg). Auscultation revealed an ejection systolic murmur which was heard throughout the precordium. There were multiple splinter haemorrhages and digital ischaemia of the left hand and a pregangrenous appearance in the right forefoot. Haemoglobin was 144 g/l; white cells 9.1 × 10⁹/l; platelets 296 × 10⁹/l; and C reactive protein (CRP) 329 mg/l. Blood urea and creatinine were 18.1 mmol/l and 1094 µmol/l, and serum sodium was 131 mmol/l, potassium 2.8 mmol/l, chloride 90 mmol/l, and bicarbonate 28 mmol/l. Blood cultures (two of six sets) grew a Gram positive coccus, identified as S capitis by the Staphylococcus Reference Laboratory of the Central Public Health Laboratory. Disc diffusion testing revealed the isolate to be susceptible to vancomycin, flucloxacillin, and rifampicin. Echocardiography revealed a large (2 × 1 cm) vegetation on the non-coronary cusp of the aortic valve with no significant aortic incompetence. There was a 40 mm Hg gradient across a calcified aortic valve with hypertrophy but good left ventricular function. Endocarditis was diagnosed and treatment was started with intravenous vancomycin hydrochloride and oral rifampicin. We discussed the possibility of early aortic valve surgery because of the vegetation and previous embolic event but we decided to treat medically. On day 6, repeat echocardiography showed the vegetation to have vanished. This was not associated with a clinical embolic event.

Despite prostaglandin infusion the patient developed increasing pain in his right foot, which subsequently became gangrenous and required amputation. After a six week course of antimicrobial agents he had no fever, CRP was 16 mg/l, and repeat echocardiography showed no vegetation on the aortic valve and no change in valve appearance. There was no evidence of recurrent infection 10 months after discharge.

Case 2
A previously healthy 35 year old man presented with pleuritic-type retrosternal chest pain radiating to the left scapular region. In the preceding two weeks he had increasing arthralgia, night sweats, and lethargy. On examination he was afebrile but pale and in obvious discomfort. Two splinter haemorrhages on his left index finger were noted, but no other stigmata
Table 1  Summary of the data on published episodes of Staphylococcus capitis endocarditis

<table>
<thead>
<tr>
<th>Case (reference)</th>
<th>Age (years)</th>
<th>Previous cardiac pathology</th>
<th>Associated conditions</th>
<th>Valve(s) involved</th>
<th>Preceding dental treatment</th>
<th>Surgery during acute phase</th>
<th>Antibiotics</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>72</td>
<td>NS</td>
<td>Peripheral vascular disease, CVA</td>
<td>Mitral NS</td>
<td>No</td>
<td>28 days iv vancomycin + iv gentamicin</td>
<td>Survived</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>53</td>
<td>Mitral valve prolapse</td>
<td>None</td>
<td>Mitral No</td>
<td>No</td>
<td>14 days iv amoxicillin + iv netilmicin followed by 30 days iv ceftazidime Cloxacillin (duration not stated)</td>
<td>Survived</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>63</td>
<td>Ventricular septal defect, previous endocarditis</td>
<td>None</td>
<td>Tricuspid Yes</td>
<td>No</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>29</td>
<td>Mitral valve prolapse with insufficiency</td>
<td>None</td>
<td>Mitral NS</td>
<td>No</td>
<td>40 days iv benzylpenicillin + iv gentamicin</td>
<td>Survived</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>62</td>
<td>None</td>
<td>Mitral Yes</td>
<td>No</td>
<td></td>
<td>Initially iv vancomycin + gentamicin, followed by 21 days iv benzylpenicillin + gentamicin, followed by 14 days po pefloxacin + rifampicin</td>
<td>Survived</td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>65</td>
<td>Mitral valve replacement (× 3)</td>
<td>NS</td>
<td>Mitral No</td>
<td>Yes</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>46</td>
<td>Aortic stenosis</td>
<td>ESRF</td>
<td>Aortic No</td>
<td>No</td>
<td>42 days iv imipenem + iv vancomycin</td>
<td>Survived</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>35</td>
<td>Bicuspid aortic valve</td>
<td>None</td>
<td>Aortic No</td>
<td>No</td>
<td>28 days iv vancomycin + po rifampicin</td>
<td>Survived</td>
<td></td>
</tr>
</tbody>
</table>

NS, not stated; CVA, cerebrovascular accident; ESRF, end stage renal failure.

ears. As with other CoNS it is known to cause infection around prosthetic devices, but is rarely recognised as causing endocarditis; our cases bring the total number of reported cases to eight. An underlying cardiac abnormality which may have predisposed the patient to infection was reported in seven of the eight cases (table 1). The patient in case 2 had a congenitally bicuspid valve which was only diagnosed after the patient presented with endocarditis. Although the mitral valve was most commonly infected (five of eight cases) it is difficult to infer a tropism for a particular valve because of the small number of cases described. All but one case involved native valves, which contrasts with the widely held view that CoNS endocarditis is primarily an infection involving prosthetic valves. Data from this small series suggest that, in order to cause disease, S capitii requires the presence of an underlying valvar anomaly. In our laboratory, we have found that S capitii produces DNase with extended incubation (JAT Sandoe, unpublished data, 1997). This enzyme is a recognised virulence factor of Staphylococcus aureus and could play a role in the pathogenesis of cardiac infection. It is noteworthy that in two of the six cases where the history was recorded, the patients had had dental work before the development of endocarditis. However, as S capitii is not known to be part of the normal oral microflora, these may be coincidental findings.

Methicillin resistance (and thus fluoroacillin resistance) is uncommon among S capitii isolates but has been reported in one community acquired case of native valve endocarditis and one hospital acquired case of prosthetic valve endocarditis. A variety of antibiotic regimens has been used successfully in the management of S capitii endocarditis. In the five cases in which duration of treatment is given, antimicrobial agents were administered for between four and six weeks. All patients survived, although one individual required valve replacement during the acute phase of the illness. Of particular interest are reports of two...
cases which, despite major embolic events, were managed successfully with medical treatment, without recourse to valve replacement. Commercially available kits for the identification of CoNS are available but they are relatively expensive and time consuming to use. Although such bacteria are frequently isolated in the diagnostic microbiology laboratory, many of the isolates are thought to be contaminants which have been introduced during specimen collection or processing in the laboratory. Accordingly, it has not been routine laboratory practice to identify CoNS to species level and, therefore, these isolates may be labelled as *Staphylococcus epidermidis* (*sensu lato*). Consequently, infection which is associated with non-epidermidis species may be underrecognised and underreported.

This review highlights the potential for *S. capitis* to cause serious native heart valve infections in immune competent patients with valvar disease. Although the number of cases in this series is small, the data suggest that a four week course of antimicrobial agents may be sufficient to treat *S. capitis* endocarditis provided that the patient has a sustained clinical response and that inflammatory markers return to within normal limits. Native valve endocarditis due to *S. capitis* has a favourable prognosis and may respond to medical treatment alone even in the presence of a major embolic phenomenon. Species identification of CoNS is important to establish the natural history of endocarditis caused by individual species, and thus help in the decision to manage the disease with medical or combined medical and surgical treatment.