Tricuspid Endocarditis, in a 12 year old girl with a previously normal heart

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Case Report

A 12 year old girl was referred to QECH Paediatric and Child Health Department with a two week history of dry cough, fever and chills. Three days before admission she became pale and short of breath. At the onset of the symptoms, she was given antimalarial tablets (lumifanterine/artemether) at her local health centre but symptoms worsened. Three days prior to admission when pallor was noted, she was referred to a district hospital where diagnoses of severe anaemia and pneumonia were made. She was transfused 440mls of whole blood and started on intravenous penicillin. She was referred to QECH after 2 days due to recurrent severe anaemia and worsening pneumonia.

She was previously well with no history of shortness of breath on exercise, orthopnoea or chronic cough and no relevant past medical history. There was no family history of congenital heart disease.

On arrival, she looked unwell, in severe respiratory distress, with an oxygen saturation of 64% in air. She was wasted with a body weight of 27kg. Her axillary temperature was 38.6°C, pulse rate was 128b/min and respiratory rate 88 cycles/min. She was pale, had bilateral coarse crepitations in the chest, a gallop rhythm, a 4cm tender hepatomegaly, 3cm splenomegaly and normal looking skin. She had no subungual petechiae or Janeway lesions.

On admission differential diagnoses included severe anaemia (unknown cause), heart failure and severe pneumonia.

Initial investigations were: haemoglobin 5.3g/dl, WBC 23x10³/mm³ (lymphocytes 2.70x10³/mm³, monocytes 0.80x10³/mm³, granulocytes 12.5 x10³/mm³) and platelets 23 x 10³/mm³. Microscopy of a thick blood film for malaria parasites, HIV antibody test (Determine™ Trinity Biotech plc, Bray Ireland) and Mantoux,(Tuberculin, Purified Protein Derivative RT 23 SSI, Denmark) were all negative. She was started on IV penicillin, gentamicin and frusemide. She received a blood transfusion and was put on oxygen therapy.

On day 2, her respiratory distress and oxygen requirements increased. She remained in heart failure. Blood cultures were taken. Chest x-ray showed bilateral fluffy infiltrates with an enlarged cardiac shadow (Figure 1).

Staphylococcal pneumonia was suggested and IV cloxacillin was added to her treatments.

In view of her chest x-ray, an echocardiograph was done which showed a large oscillating intracardiac mass (vegetation) on the tricuspid valve and tricuspid valve regurgitation. There was no other cardiac abnormality, in particular no ventricular septal defect or left to right shunt producing a jet impacting on the tricuspid valve (Figures 2 and 3).
Her blood cultures subsequently grew Staphylococcus aureus sensitive to cloxacillin, gentamicin and erythromycin. Unfortunately, the blood culture was not repeated. Therefore, according to the modified Duke criteria (clinical criteria) for the diagnosis of infective endocarditis (1), she was classified as a possible infective endocarditis - with one major (endocardial involvement) and one minor (fever > 38°C) criteria.

Maintenance frusemide was added to her treatment. She required three further transfusions on days 5 and 10. Her full blood counts (FBCs) on days 5, 10 and 25 were as follows:

<table>
<thead>
<tr>
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<th>On Admission</th>
<th>Day 5</th>
<th>Day 10</th>
<th>Day 25</th>
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<tbody>
<tr>
<td>Hb</td>
<td>5.3g/dl</td>
<td>5.2g/dl</td>
<td>5.9g/dl</td>
<td>10g/dl</td>
</tr>
<tr>
<td>WBC</td>
<td>16x10^9/mm³</td>
<td>14x10^9/mm³</td>
<td>11x10^9/mm³</td>
<td>9x10^9/mm³</td>
</tr>
<tr>
<td>Platelets</td>
<td>23x10^9/mm³</td>
<td>122x10^9/mm³</td>
<td>127x10^9/mm³</td>
<td>300x10^9/mm³</td>
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On day 14 of admission she developed a grade 2/6 holosystolic murmur best heard on the left lower sternal edge which persisted until the day of discharge (day 30). She was weaned off oxygen by day 18 when she saturated well in room air.

She remained intermittently febrile until day 26 of treatment. She was discharged home after 30 days of treatment with cloxacinill and frusemide with medications for a further two weeks. She was reviewed in the paediatric cardiac clinic 2 weeks after discharge when she had no symptoms or signs of cardiac failure. She still had a tricuspid regurgitation murmur.

Discussion

Bacterial endocarditis is an elusive diagnosis. Its variable systemic manifestations often obscure the underlying presence of valvular infection. The Modified Duke Criteria are used in the diagnosis of infective endocarditis. There are three diagnostic categories: definite, possible and rejected. A diagnosis of "definite infective endocarditis" is made using either pathologic or clinical criteria. See table above:

A common error in diagnosis is when an extra-cardiac complication of endocarditis is mistaken as the cause of all the symptoms. In this case, the complication (pneumonia) was recognized initially, but the endocarditis which had caused it had not. The consequent delay in correct evaluation may delay antibiotic therapy and decrease the probability of a successful outcome. Our patient presented with a severe pneumonia and anaemia. She initially was labeled as a ‘respiratory’ patient and was treated as such. The echocardiographic findings of tricuspid regurgitation were unexpected and the diagnosis could easily have been missed. Tricuspid endocarditis accounts for only 5-10% of all cases of infective endocarditis. Most cases occur in intravenous drug users, patients who have had invasive instrumentation, or in children with high-velocity jet streams of blood such as a ventricular septal defect or dysplastic atrioventricular valve. Although it is recognized that infection of the tricuspid valve does occur in children with previously normal hearts and no other known risk factors, this is thought to be extremely rare.

Symptoms arise from pneumonia or septic pulmonary emboli from dislodged valvular vegetative material. Pulmonary manifestations are common in right sided endocarditis. This is because there is migration of septic material from the tricuspid valve into the pulmonary arteries. Septic material can be dispersed into a pulmonary vessel causing either pulmonary embolism or pneumonia. Radiographic findings may resemble a staphylococcal pneumonia the features of which include extensive fluffy alveolar infiltrates involving a whole lobe or several lobes. The valvular infection may go unnoticed because the cardiac murmur is absent in 45% of the patients with pulmonary
complications\(^{10}\). In our patient, the murmur was either missed or absent in the initial stages. Staphylococcus aureus is the most common bacterial isolate and requires prolonged and aggressive antibiotic therapy. This includes treatment with semisynthetic β-lactamase-resistant penicillins for a minimum of 6 weeks. Surgery is indicated for large vegetations (>10mm). The surgical options include simple excision of the vegetation, valvectomy, tricuspid valve repair, or valve replacement. Valvectomy without reconstruction would result in frank regurgitation postoperatively. If the pulmonary pressure is normal (in most cases, the heart was previously normal), this may be tolerated for years. Valve replacement gives better long-term haemodynamic results when a repair is unlikely to be satisfactory, as in cases of large or multiple leaflet destruction\(^ {10}\).

**Conclusion**

Isolated tricuspid valve endocarditis is rarely seen without intravenous drug use, invasive cardiac procedure or previous valvular disease. A high index of clinical suspicion is required to make the diagnosis. It should be considered in all patients with staphylococcal pneumonia.

**Summary**

A 12 years old, HIV non reactive girl developed tricuspid valve endocarditis and pneumonia caused by Staphylococcus aureus. She presented with symptoms and signs of pneumonia, anaemia and heart failure. The patient had no history of intravenous drug use nor had she been admitted to hospital prior to this illness. Echocardiography revealed no predisposing pre-existing abnormalities of the valve or heart. The underlying diagnosis could have been missed.

**References**