A massive pericardial effusion in South Africa is not always due to tuberculosis

**CASE REPORT**

Sipho Ntshalintshali, Nontembiso Mhlana, Farzana Moosajee, Riyad Abousriwiel, Kiran George and Riette du Toit

Radiographs of involved joints were normal while the chest radiograph showed an increased cardiothoracic ratio (CTR), while the echocardiogram revealed a massive pericardial effusion, and pericardiocentesis was done. The abdominal ultrasonography was normal. The blood test results are summarised in Table I. A decision to start empiric ATT was made considering the local high incidence of tuberculosis. A week later he had no clinical improvement but developed a drug-induced liver injury and ATT was discontinued. SJIA was considered as a more likely alternative diagnosis since the patient fulfilled the Yamaguchi criteria, and was given a 750mg intravenous pulse of solumedrol daily over a 3-day period, followed by a remarkable clinical response. Within 2 weeks he was discharged on prednisone at 0.5mg/kg/day and azathioprine 50mg daily as a steroid sparing agent.

The second case is that of a 28-year-old male with a background of dengue fever 3 years prior, who now presented with a 1-week history of a pharyngitis, polyarthralgia, and chest pain with shortness of breath. He presented with Beck's triad for a pericardial tamponade with a blood pressure of 65/45, heart rate of 130 beats per minute, and a respiratory rate of 28 breaths per minute. He had a quotient pattern fever, a non-palpable, non-blanching vasculitic type rash on both shins, and small cervical lymphadenopathy. He also had bilateral small knee effusions. Other organ-system findings were unremarkable.

**ABSTRACT**

South Africa (SA) has a high incidence of tuberculosis. Medical conditions mimicking tuberculosis often result in erroneous treatment with antitubercular therapy (ATT) before a definitive diagnosis is made. We report on 2 cases presenting with massive pericardial effusions secondary to Still’s disease (SJIA) and Adult onset Still’s disease (AOSD). Both cases were treated with ATT, with an alternative diagnosis only considered upon poor response to therapy and the development of ATT-associated side effects. Our objective is to remind clinicians of other potential differential diagnoses in the clinical scenario of massive effusive pericarditis in a tuberculosis endemic region. SAHeart, 2020;17:204-207

**INTRODUCTION**

Massive pericardial effusions can be due to multiple etiologies. In South Africa, tuberculosis is a common cause of massive pericardial effusion, SJIA and AOSD, which usually present in 5% - 10% of patients as pyrexia or fever of unknown origin (FUO) accompanied by systemic manifestations, can be associated with a massive pericardial effusion and mimic tuberculosis.

**CASE REPORTS**

We report on a 15-year-old male, HIV uninfected, who presented with a 3-week history of a productive cough, dyspnoea, fever, weight loss and multiple large joint arthralgia. On examination he was tachycardic, normotensive, had a respiratory rate of 22 breaths per minute, and had a fever with a quotient pattern (Figure 1). He had cervical, axillary and inguinal lymphadenopathy of approximately 1cm. On cardiovascular and respiratory examination he had a small volume and tachycardic pulse, with distended neck veins, audible normal heart sounds with no pericardial rub, and bilateral coarse crackles. He had inflammatory polyarthritis involving the wrists, elbows, knees, and ankles. No skin rash was detected and other organ system findings were unremarkable.

Radiographs of involved joints were normal while the chest radiograph showed an increased cardiothoracic ratio (CTR), while the echocardiogram revealed a massive pericardial effusion, and pericardiocentesis was done. The abdominal ultrasonography was normal. The blood test results are summarised in Table I. A decision to start empiric ATT was made considering the local high incidence of tuberculosis. A week later he had no clinical improvement but developed a drug-induced liver injury and ATT was discontinued. SJIA was considered as a more likely alternative diagnosis since the patient fulfilled the Yamaguchi criteria, and was given a 750mg intravenous pulse of solumedrol daily over a 3-day period, followed by a remarkable clinical response. Within 2 weeks he was discharged on prednisone at 0.5mg/kg/day and azathioprine 50mg daily as a steroid sparing agent.

The second case is that of a 28-year-old male with a background of dengue fever 3 years prior, who now presented with a 1-week history of a pharyngitis, polyarthralgia, and chest pain with shortness of breath. He presented with Beck’s triad for a pericardial tamponade with a blood pressure of 65/45, heart rate of 130 beats per minute, and a respiratory rate of 28 breaths per minute. He had a quotient pattern fever, a non-palpable, non-blanching vasculitic type rash on both shins, and small cervical lymphadenopathy. He also had bilateral small knee effusions. Other organ-system findings were unremarkable.

The electrocardiogram showed a sinus tachycardia with microvoltage. The chest X-ray revealed an increased CTR (Figure 2), and on echocardiogram had evidence of a cardiac tamponade.

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without stranding. The abdominal ultrasonography was normal. The initial diagnostic impression was tuberculous pericarditis with Poncet’s disease. A therapeutic and diagnostic pericardiocentesis was performed, and he was started on empiric ATT. A week later, he had ongoing temperature spikes while his rash and arthritis improved. Since he fulfilled the Yamaguchi criteria for AOSD and was not responding to ATT as expected, he was treated with an intravenous steroid pulse of 750mg over a 3-day period, followed by a remarkable response. He was later started on oral prednisone 30mg per day and azathioprine 50mg and was discharged a week later for review as an outpatient.

**DISCUSSION**

Tuberculosis is a granulomatous disease secondary to *Mycobacterium tuberculosis* infection. It is a significant public health issue in South Africa – 3% of the world’s tuberculosis cases are from SA and it is the eighth leading cause of death in this country. In 2018, a total of 63 000 people died of tuberculosis of which two-thirds were infected with HIV. It is estimated that about 301 000 people were diagnosed with tuberculosis in South Africa in 2018. However, according to the 2019 WHO report, both tuberculosis incidence and mortality rate are decreasing each year.

Pulmonary tuberculosis is the commonest form of presentation. Extra-pulmonary tuberculosis is not uncommon, however,
A MASSIVE PERICARDIAL EFFUSION IN SOUTH AFRICA IS NOT ALWAYS DUE TO TB

and more so in immunocompromised patients. Tuberculous pericarditis is not infrequent in HIV-1 infected patients, and it is the most life-threatening form of extrapulmonary tuberculosis. Tuberculosis causes inflammation of the pericardium leading to pericardial effusion, potentially causing acute circulatory collapse. One of the long-term complications of tuberculous pericarditis is constriction of the pericardium.\(^2\)

The treatment for TB pericarditis is the standard ATT consisting of rifampicin, isoniazid, pyrazinamide and ethambutol. Pericardial drainage is also performed in cases of massive pericardial effusion, and subsequently pericardiectomy if constriction develops. Adjunctive therapies to ATT have been evaluated. The Investigation of the Management of Pericarditis (IMPI), a large series of clinical trials that evaluated the effect of prednisolone or Mycobacterium indicus pranii or both, added to the standard ATT regimen, showed no mortality benefit with either of the adjunctive therapies. However, the prednisolone group showed a lower rate of constriction and hospitalisation compared to the placebo group. The increased incidence of malignancy with both interventions, and a marked increase in HIV-related malignancies in the prednisolone group, were matters of concern.\(^3\)

Fever is one of the cardinal features of tuberculosis, and cases of disseminated or extrapulmonary forms of tuberculosis are usually associated with high fevers.\(^4\) Tuberculosis-associated fever pattern can be intermittent or recurrent, or present in a double quotidian form, which is 2 temperature spikes per day.\(^5\) SJIA/AOsd are often included in the differential diagnosis of patients presenting with high fevers more than 39°C, usually as a quotidian (daily) pattern, and often in the afternoon or early evening.\(^6\) Tuberculosis with reactive joint involvement referred to as Poncet’s disease can mimic SJIA/AOsd disease due to similar clinical findings.\(^7\) The fever pattern may occasionally assist in differentiating the 2 conditions.

Over and above the clinical features, a serum globulin level of more than 40g/L and a leukocyte count less than 10 x 10^9/L are highly predictive of tuberculous pericarditis in the appropriate setting.\(^7\) Pericardial fluid investigational variables that can be used for a diagnostic model for tuberculous pericarditis include interferon gamma (IFN gamma) and adenosine deaminase (ADA) (see Table II). The combination of fluid ADA, fluid lymphocyte/neutrophil ratio, HIV status and blood leukocyte count have a diagnostic sensitivity of 96% and specificity of 97%.\(^7\)

SJIA is a systemic autoinflammatory disease mainly affecting the joints of patients below the age of 16 and is responsible for approximately 10% - 15% of all cases of JIAs. Similarly, AOsd is a systemic autoinflammatory disease in the population older than 16 years.\(^8\) Multiple diagnostic criteria have been formulated for the diagnosis of AOsd, and among these the most prominent and widely utilised are the Yamaguchi and Fautrel’s criteria (Tables III and IV respectively).\(^8\)
The combination of a 5-fold increase in serum ferritin and the total glycosylated ferritin of less than 20% increases the odds for AOSD diagnosis (sensitivity of 43.2%, and specificity of 92.9%).\(^9\) AOSD-associated massive pericardial effusions resulting in pericardial tamponade have been described in the literature. Therapeutic pericardiocentesis and immunomodulatory therapy have been the cornerstone of therapy.\(^10\)

**CONCLUSIONS**

Many similarities exist between the clinical presentation of extrapulmonary tuberculosis and that of sJIA/AOSD. While exclusion of an underlying infection is essential in the diagnostic workup of any patient with sJIA/AOSD, the consideration of an auto-inflammatory condition is important in any patient with a presumed diagnosis of tuberculosis. This is true in the absence of an appropriate clinical response to ATT, without microbiological confirmation of tuberculosis.

**LEARNING POINTS**

- Tuberculosis remains the most common cause of massive effusive pericarditis in South Africa; however, alternative diagnoses must be considered in the appropriate setting.

- The quotidian fever pattern and glycosylated ferritin levels below 20% may assist in the differentiation of tuberculosis in favour of sJIA/AOSD.

- SJIA/AOSD is associated with markedly elevated ferritin concentration in 70% of patients. This correlates with disease activity and can be used as a marker for clinical response.

**Conflict of interest**: none declared.

**REFERENCES**


**TABLE III: Yamaguchi classification criteria for AOSD.\(^8\)**

<table>
<thead>
<tr>
<th>Major criteria</th>
<th>Minor criteria</th>
<th>Exclusion criteria</th>
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<tbody>
<tr>
<td>Spiking fever ≥39° C lasting 1 week or longer</td>
<td>Maculopapular rash</td>
<td>Infections</td>
</tr>
<tr>
<td>Arthralgia or arthritis lasting 2 weeks or longer</td>
<td>Leukocytosis ≥ 10 000/mm³</td>
<td>Recent development of significant lymphadenopathy</td>
</tr>
<tr>
<td>Transient erythema</td>
<td>Polymorphonuclear cells ≥80%</td>
<td>Malignancies (mainly malignant lymphoma)</td>
</tr>
<tr>
<td>Pharyngitis</td>
<td>Glycosylated ferritin ≤20%</td>
<td>Other rheumatic disease (mainly systemic vasculitides)</td>
</tr>
<tr>
<td>Leucocytosis ≥10 000/mm³ with ≥80% polymorphonuclear cells</td>
<td>Abnormal liver function tests</td>
<td></td>
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<td></td>
<td>Negative tests for antinuclear antibody (AN) and rheumatoid factor (RF)</td>
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<td>Five or more criteria required, of which 2 or more must be major.</td>
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**TABLE IV: Fautrel classification criteria for AOSD.\(^8\)**

<table>
<thead>
<tr>
<th>Major criteria</th>
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<td>Glycosylated ferritin ≤20%</td>
</tr>
<tr>
<td></td>
<td>Four or more major criteria required or 3 major + 2 minor criteria.</td>
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