CASE REPORT

HIV infection associated with Strongyloides stercoralis colitis resulting in Streptococcus bovis bacteraemia and meningitis

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We report the case of an HIV infected patient with Streptococcus bovis bacteraemia and meningitis associated with gastrointestinal Strongyloides stercoralis infection. To our knowledge, this has been reported once previously and serves as a reminder to actively exclude asymptomatic S stercoralis infection in HIV infected individuals presenting with bacteraemia.

A 23 year old black African woman, presented with a 4 day history of fever and rigors, and 24 hours of severe frontotemporal headache, neck stiffness, and photophobia. She gave a 6 month history of weight loss and constipation, but denied diarrhoea. She reported previous episodes of malaria and a possible diagnosis of pulmonary tuberculosis, for which only 3 months treatment was given in Africa.

On examination her temperature was 39.7°C, pulse 126/minute, and blood pressure 133/73. She was alert, with mild neck stiffness and photophobia, but no focal neurological signs.

Funduscopy was unremarkable and there were no peripher al stigmata or murmurs suggestive of infective endocarditis. Lesions consistent with Kaposi's sarcoma were present on her shins.

Investigations showed haemoglobin 9.1 g/dl, white cell count (WCC) 4.68×10^6/l (neutrophils 2.67×10^6/l, lymphocytes 1.28×10^6/l, eosinophils 0.31×10^6/l), platelets 185×10^6/l, Na+ 122, other electrolytes and liver function tests normal, C reactive protein 8.6 mg/l, and malarial parasites not seen. Chest x ray was normal and a computed tomogram head scan was unremarkable. A lumbar puncture yielded turbid fluid with an opening pressure of 15.5 cmH2O. Microscopy revealed: red cells 155×10^6/l, WCC 1625×10^6/l, (neutrophils 2.67×10^6/l, lymphocytes 0.31×10^6/l, eosinophils 0.31×10^6/l, platelets 185×10^6/l). Histology from the colon showed marked infestation with S stercoralis and special stains for acid fast bacilli, fungi, and cytomegalovirus were all negative. She was treated with a 5 day course of ivermectin (12 mg/day). Her symptoms resolved and eosinophil count normalised. She remains well 4 months later, with an undetectable viral load (<50 copies/ml).

DISCUSSION

Streptococcus bovis is a part of the gastrointestinal tract flora in 10–16% of healthy people and is a well recognised cause of bacteraemia and endocarditis.1,2 The association between this organism and colonic neoplasms is also well documented and an estimated 40.3% of patients with S bovis bacteraemia or endocarditis are reported to have underlying gastrointestinal tumours.3 Meningitis due to S bovis is extremely rare and there are only 21 previously reported cases.

S stercoralis is an intestinal nematode endemic in many developing countries. Infection is frequently asymptomatic for long periods of time. Symptoms may consist of mild gastrointestinal discomfort and chronic diarrhoea. In those from endemic areas with diarrhoea, the incidence of S stercoralis in HIV infected patients is thought to be higher than in HIV negative individuals and is estimated to be around 5–6%.4 It is unclear whether a similar increase is present in asymptomatic cases. The occurrence of bacteraemia in S stercoralis infection is well documented and may result simply from bowel wall ulceration acting as a portal of entry for enteric organisms. There is some evidence that intestinal bacteria are also carried by invasive S stercoralis larvae on their outer surfaces.5 Commonly associated bacteria include Escherichia coli, Klebsiella spp, and Streptococcus fecalis, with only two previously reported cases of S bovis bacteraemia.6,7

Our case illustrates the importance of actively investigating the gastrointestinal tract in patients with S bovis bacteraemia. Furthermore, in HIV infected individuals with a history of travel to endemic areas one should exclude potentially fatal parasitic infections such as S stercoralis.

Key messages

- Individuals with enteric organisms in sterile sites should have appropriate gastrointestinal investigations; in such individuals with a risk factor for strongyloides this should be ruled out, in particular in HIV positive individuals, and individuals with strongyloides may not have diarrhoea.
CONTRIBUTORS
TdS, MR, and MP were involved in the care of this patient during admission; MP followed up the patient; TdS and MR carried out the literature review and wrote the initial manuscript; MP reviewed the initial manuscript before submission and revised the manuscript before resubmission.

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