Chronic human brucellosis with pancytopenia in a young girl from a camp for Internally Displaced Persons

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Case Report

A 19-year old girl was transferred to our hospital from Base hospital Vavuniya in June 2009 for investigation and management of abdominal distention. She lived in a camp for the Internally Displaced People (IDP) in Mannar and had developed a low grade fever and abdominal distention for one month duration. Apart from mild fatigue, there were no other significant symptoms. She had consumed unpasteurised cow's milk in the war area. She had no past history or a contact history of tuberculosis.

On examination she was pale, anicteric, afebrile and had no lymphadenopathy. She had non-tender mild splenomegaly but liver was not palpable. Her respiratory, cardiovascular and neurological systems were clinically normal. Her Hb was 7.6 g/dL while White cell and platelet counts were 2.0x10^9/L and 25x10^9/L, respectively. Blood picture showed pancytopenia and possible iron deficiency anaemia. Her initial ESR was 45 mm. Serum AST and ALT were 41 and 30 U/L, respectively, while INR was 1.49. Renal functions were normal but CRP was repeatedly elevated. Serum protein electrophoresis showed normal albumin band with polyclonal increase in γ globulin. Abdominal ultrasonography showed splenomegaly with multiple echogenic spots in both spleen and liver. Bone marrow biopsy showed no evidence of haematological or non-haematological malignancy and marrow iron stores were completely depleted. Her liver biopsy showed focal nodular hyperplasia but no granulomas were detected.

At this stage, the possibility of either chronic disseminated TB or chronic brucellosis was considered. Bone marrow culture for mycobacterium tuberculosis was negative and PCR for TB was negative. However brucella PCR was positive for 10 copies.

Contrast CT abdomen revealed focal nodular hyperplasia (a benign condition seen in females) in the right lobe of the liver with multiple hyper-echoic lesions. With the history of consumption of unpasteurised cow's milk and positive PCR for brucella we confirmed the diagnosis of chronic brucellosis associated with bone marrow suppression causing pancytopenia.

She was started on the combination of oral ciprofloxacin 500 mg bid and doxycycline 100 mg bid for 6 weeks. Haematinics were given to replenish iron stores. At the end of treatment she was asymptomatic and her blood count returned to normal.

Discussion

Brucellosis is a chronic granulomatous zoonotic infection which can affect every organ of the human body. Among the four species found, a vast majority is caused by B.melitensis (1). It is transmitted to human by consumption of unpasteurised animal milk products and as an occupational disease in shepherds, veterinarians and dairy-industry professionals. Brucella species have a unique ability of invading both phagocytic and non-phagocytic cells and surviving in the intracellular environment by avoiding the immune system in different ways, explaining why brucellosis is a systemic disease and can involve almost every organ (2). Our patient has consumed unpasteurised cow's milk during the war and developed brucellosis complicated by pancytopenia. The frequency of pancytopenia ranges from 3% to 20% (3). Hematological abnormalities usually resolve promptly with treatment of the primary disease (3). Pancytopenia could be due to hypersplenism, bone marrow involvement, haemophagocytosis or immune-mediated cell destruction (4,5). The gold standard for diagnosing
brucellosis is bone marrow culture which is difficult in Sri Lanka. PCR is fast, could be done in any body tissue and becomes positive after 10 days of inoculation (1). Our patient was successfully treated with the combination of oral ciprofloxacin and doxycycline for 6 weeks.

Keeping in mind the possibility of brucellosis in people who are living in temporary camps, mainly in the North, East provinces in Sri Lanka, health care workers should be more alert on early diagnosis and treatment of brucellosis to prevent further spreading.

References
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Fatal haemorrhage due to a fish bone in the oesophagus; a case report
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Introduction
Foreign bodies in the upper aero-digestive tract are a common problem. Rupture of the oesophagus is one of the rare yet more dramatic conditions and is an emergency. Studies show that 10% to 20% of foreign body ingestion in the gastrointestinal tract requires endoscopic treatment and its retention in the esophagus may represent risk of severe complications, with rare fatalities. In addition to endoscopic treatment, surgical procedure might be necessary in approximately 1% (1).

Case Report
A 56-year old male was admitted to a surgical ward with a three day history of odynophagia. He had no dysphagia or haematemesis during the admission. There was no history of ingestion of a foreign body. The initial upper gastrointestinal endoscopy and chest radiograph did not reveal any significant changes and the patient was discharged. A week later, he was readmitted due to an episode of haematemesis. The endoscopic examination was repeated and there was a resistance at the upper third of oesophagus due to spasm of the oesophageal wall. Biopsies were taken. The patient suddenly collapsed and died on the next day.

Autopsy revealed a 750 ml of clotted fresh blood in the thoracic cavity, which originated from a point between the oesophagus and the aorta. Further sectioning showed a perforation at the level of the arch of the aorta with a 3.5 cm long fish bone (Fig. 1) impacted in the ulcerating aorto-oesophageal fistula (Fig. 2, Fig. 3). Tarry black stools were found in the lower gastrointestinal tract.