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# Un exceptional cause of lymphocytic colitis: Brucellosis

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#### **Abstract**

Colitis is a rare but serious and life-threatening complication of brucellosis. The association of infective colitis and brucellosis has formerly been reported. However, to our knowledge, we report herein the first case of lymphocytic colitis due to brucellosis. A 44-year-old man presented with myalgia, chronic diarrhea and intermittent vomiting. Laboratory tests showed severe hypokalemia with malabsorption syndrome and positive serological test for brucellosis.

Colonoscopy revealed lymphocytic colitis. After treatment with rifampicin and doxycycline, the patient recovered.

# Introduction

Brucellosis is a common zoonosis which still remains as a major health problem in certain parts of the world and is transmitted from animals to man through the ingestion of un-pasteurized milk and milk products. Undifferentiated febrile illness and arthritis-dominant syndrome with hepatosplenomegaly and lymphadenopathy are the usual modes of presentation. The gastroenterological manifestations of human brucellosis are diverse relatively uncommon, ranging from the nonspecific, such as diarrhea and abdominal pain, to the pathologically distinct hepatic lesions, and to the rare colonic, pancreatic, and peritoneal involvement [1-4]. Infective colitis secondary to brucellosis has been reported. But, to our knowledge, we report herein the first case of lymphocytic colitis due to brucellosis with a brief review of the relevant literature.

## Case report

A 44-year-old man was admitted initially in department of neurology with rapidly evolving muscle pain in proximal limbs and arms over a few weeks. Shortly after that, generalized weakness developed, and he began to have an intense aching pain in the muscles of his arms and legs. The thighs were slightly tender to touch, and the patient was unsteady on his feet. Muscle strength was 5/5 in the arms and the legs. The deep tendon reflexes were normal, as was sensation.

Serum enzymes indicating myositis were markedly elevated (creatine kinase [CK] 6897 U/L, normal range 0-171 U/L; Lactate dehydrogenase was also elevated (285 U/L; normal range 50-248 U/L).

Electromyography of the right iliopsoas muscle showed low amplitude, polyphasic, myopathic units, the interference pattern indicated myopathy. Electroneurography was normal. The overall picture was suggestive of a polymyositis-like syndrome with rhabdomyolysis. The patient was transferred in our department of internal medicine.

He had neither recent history of muscular injuries, viral infections or heavy physical exercise nor a family history of hereditary neuromuscular diseases. He also denied drug intake, trauma, infection and exposure to any chemicals. He has lived in a rural area with a

livestock-raising family.

He also suffered from chronic diarrhea and described an intermittent vomiting without fever.

On physical examination, the abdomen was tender; the liver and spleen were not enlarged. There was no pathological finding on cardiac, pulmonary, head and neck examination. The cranial-nerve functions were preserved. Laboratory test results revealed: hemoglobin 11.6 g/dl; MCV: 96fL; white blood cells 3050 /mm³; platelets 368,000/mm³; reticulocyte count was 0.1%; ALT 553 U/L (normal 5-52); AST 250 U/L (normal 8-35); and erythrocyte sedimentation rate was 65 mm/hour.

A severe hypokalemia with malabsorption syndrome was showed. Parenteral potassium and hydration with intravenous fluids was immediately initiated.

Serological tests for antinuclear antibody, anti-DNA and rheumatoid factors, HBsAg and IgM for anti-HBc, anti-HAV, EBV VCA, and CMV were all negative.

A colonoscopy was performed. Macroscopic examination was normal. Microscopic examinations of biopsy showed intraepithelial lymphocytes (35 lymphocytes/100 surface epithelial cells) in the colonic mucosa. The gland architecture is within normal limits (Figure 1).

At this point, the patient tested positive for brucellosis by the serum agglutination test at titers of 1/160. Retrospectively, our patient reported his consumption of un-pasteurized milk.

The final diagnosis was lymphocytic colitis due to brucellosis. The patient was treated by rifampicin (600 mg PO OD) and doxycycline (200 mg PO BD). The two antibiotics were maintained for 3 months. One month after starting the treatment, the symptoms began to

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Table 1. Gastrointestinal	manifestations of brucellosis: re	eview of literature.	
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	1	2	3	4	5	6
	[5]	[6]	[7]	[8]	[9]	Our case
Sex	F	F	M	M	M	M
Age (years)	22	16	16	21	39	44
Digestives manifestations	Splenomegaly	Rectal bleeding	Abdominal pain, vomiting, diarrhoea abdominal tenderness splenomegaly	Vomiting Diarrhea	Mucosanguineous diarrheas	Vomiting Diarrhea
Other organs involvements	Fever Rash	osteomyelitis	skin rash fever.	Fever, chills, rigors	Fever low back pain arthritis	Fever Myalgia
Blood culture		++++	++++	++++	++++	
Pus culture		++++				
Brucella agglutinin titer			++++	++++	++++	++++
Colonoscopy	Edema; hemorrhage in the propria	Nonspecific colitis	NP		Infective colitis	Lymphocytic colitis

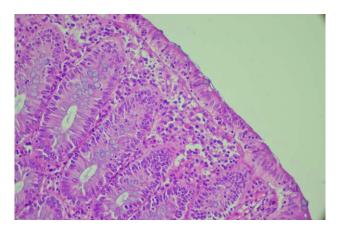


Figure 1. Presence of increased lymphocyte in surface epithelium with epithelial cell injury with epithelial cell injury HE 400.

improve. At the end of 3 months, all his complaints were resolved and the patient recovered. His physical and neurological examination and laboratory findings were all within normal ranges. CK decreased to the normal level.

During regular follow up till two years after the incident, the patient remained asymptomatic with no recurrences.

#### Discussion

Gastrointestinal manifestations of brucellosis in humans are relatively uncommon and may manifest as anorexia, nausea, vomiting, abdominal pain, diarrhea, or constipation, but systemic symptoms, such as artharlgia and myalgia, are more common than localized gastrointestinal symptoms. Brucellosis presenting as gastroenteritis has been reported in sporadic case reports (Table 1) from as early as 1934 [5].

Our case was very interesting as the presenting symptoms suggested gastroenteritis with musculoskeletal symptoms, and the laboratory findings were similar to that of gastroenteritis; only the positivity serum agglutination test of Brucella led to the diagnosis.

Diffuse myositis resulting from infectious etiology is presumed to be immune-mediated, although definitive evidence is lacking. It is reasonable to assume the hypokalemia, the mechanism that explain the pathogenesis of myositis in our patient. This hypokalemia was associated with malabsorption syndrome. These metabolic disorders were caused by chronic diarrhea. Gastroenteritis is an illness caused by many pathogens, including bacteria.

Adult acute brucellosis as a cause of colitis was reported in only few cases. Petrella and Young described a case of acute brucella ileitis in 1988. Stermer, *et al.* reported a case of brucellosis as the cause of severe colitis in 1991[5]. Locutura et al reported a case of diarrhea as the first manifestation of brucellosis in 1998 [6].

Other few cases were reported [7-10]. The histopathologic finding in our cases were different. While endoscopic examination was normal, microscopic examination showed lymphocytic colitis.

We would recommend that, in any case suggestive of gastroenteritis, the differential diagnosis of acute brucellosis should be considered in countries where brucellosis is endemic.

# **Conflict of interest**

None.

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