

Case Report

Acute acalculus cholecystitis and hepatitis caused by *Brucella melitensis*

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Abstract

Acute cholecystitis is a very rare presentation of brucellosis. A case of acalculous cholecystitis caused by *Brucella melitensis* is reported with a review of previously reported cases.

Keywords:

J Infect Dev Ctries 2010; 4(7):464-467.

(Received 26 October 2009 – Accepted 16 January 2010)

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Introduction

Brucellosis is a zoonotic infection with a worldwide distribution. It is endemic in many parts of the world. Areas of high endemicity include the Mediterranean basin, the Arabian peninsula, the Indian subcontinent, parts of Mexico, and Central and South America [1]. Human brucellosis is a multi-systemic disease with a wide range of clinical manifestations [2]. Cholecystitis is a rare complication of brucellosis. In this report, a patient who presented with a classical picture of acute cholecystitis and from whom *Brucella melitensis* was cultured from blood is reported. Other previously reported cases of brucella cholecystitis are reviewed.

Case report

A 45-year-old previously healthy Saudi male presented to the emergency department of King Khalid University Hospital with a five-day history of fever, rigors and crampy abdominal pain. The abdominal pain started in the epigastric area and became localized to the right upper quadrant area. The patient had pruritis and produced dark urine. There was no history of vomiting or diarrhoea. On physical examination, the patient was febrile (temperature of 39°C), toxic, and jaundiced. Abdominal examination showed tender epigastric area and a positive Murphy's sign. Laboratory data at the time of admission were as follows: total bilirubin 109 µmol/L; direct bilirubin 91 µmol/L; alkaline phosphatase 322 U/L; alanine aminotrasferase

159U/L; aspartate aminotransferase 149 U/L; γ glutamyl transferase 331U/L; serum amylase 23U/L; leukocyte count $8 \times 10^9/L$ with 70% neutrophils, 64% lymphocyte, 16% monocyte; hemogloubin 9g/dl platelets $75 \times 10^9/L$; PT 16; PTT 37.9; and fibrinogen assay 4.12g/L. Ultrasound scan showed a thickened gallbladder wall with no evidence of stones and mild inflammation of the adjacent liver parenchyma (Figure 1). As the patient was septic with raised liver enzymes, computed tomography (CT-scan) of the abdomen was performed to rule out liver abscess; the CT-abdomen revealed no liver lesions. The patient was therefore managed as a case of acute cholecystitis and ascending cholangitis complicated with sepsis and reactive hepatitis. After three blood cultures were obtained, intravenous tazocin 4.5 g was

Figure 1. Abdominal ultrasound revealed thickened wall of the gallbladder

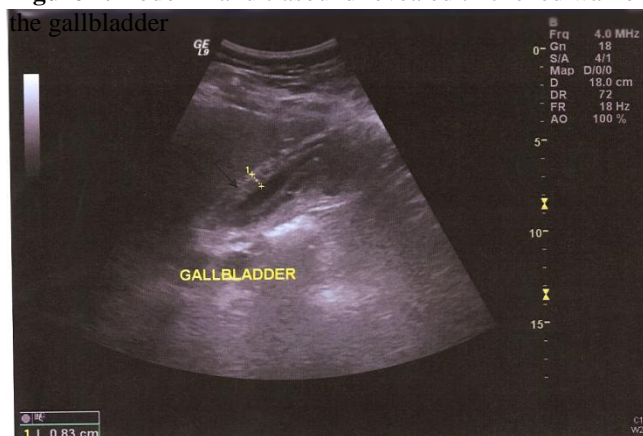


Table 1. Details of reported cases of acute cholecystitis associated with *brucella species*.

No	Author(ref)	Age/Sex	Risk factor	Blood culture	Bile culture	Gallstones	Treatment
1	Mettier ⁽¹²⁾	57/M	No	<i>B.melitensis</i>	<i>B.melitensis</i>	Absent	No
2	White ⁽¹⁶⁾	58/M	No	Negative	<i>B. abortus</i>	Present	Streptomycin
3	Valenzuela ⁽¹⁴⁾	56/M	Sheep and goat contact	NA	NA	Absent	Tetracycline +streptomycin
4	Morris ⁽¹¹⁾	34/M	Microbiologist	<i>B. suis</i>	<i>B. suis</i>	Present	Tetracycline +streptomycin
5	Berbegal ⁽¹⁷⁾	33/M	No	<i>B.melitensis</i>	Negative	Absent	Tetracycline +streptomycin
6	Shaheen ⁽¹⁸⁾	42/F	No	<i>B.melitensis</i>	<i>B.melitensis</i>	Absent	Tetracycline +streptomycin
7	Colmenero ⁽²⁾	58/M	Shepherd	<i>B.melitensis</i>	Negative	Present	Doxycycline +streptomycin
8	Fasquelle ⁽¹³⁾	72/F	Contaminated milk or dairy products	<i>B.melitensis</i>	<i>B.melitensis</i>	Present	Doxycycline + rifampin
9	Serrano ⁽¹⁹⁾	59/M	No	<i>B.melitensis</i>	Negative	Present	Doxycycline +streptomycin
10	Ashley ⁽²⁰⁾	6/M	No	<i>B. abortus</i>	NA	Absent	TMP/SMX +rifampin
11	Miranda ⁽¹⁵⁾	34/M	Sheep and goat contact	Negative	<i>B.melitensis</i>	Present	Doxycycline +rifampin
12	Andriopoulos ⁽¹⁰⁾	72/M	No	<i>B.melitensis</i>	<i>B.melitensis</i>	Absent	Doxycycline +streptomycin
13	Lopez-Prieto ⁽²¹⁾	56/F	No	Negative	<i>B.melitensis</i>	Present	Doxycycline +streptomycin
14	Kanafani ⁽²²⁾	55/M	No	<i>Brucella spp</i>	<i>Brucella spp</i>	Present	Doxycycline +rifampin
15	Kanafani ⁽²²⁾	29/F	No	Negative	<i>Brucella spp</i>	Present	Doxycycline + rifampin
16	Alotaibi (PR)	42/M	Raw milk ingestion	<i>B.melitensis</i>	NA	Absent	Doxycycline +streptomycin

commenced. The abdominal pain disappeared gradually and the patient's temperature returned to normal within three days. On day 9 after admission, one blood culture was positive. It grew small colonies of Gram-negative bacilli that were later identified as *B. melitensis*. Standard tube agglutination test (STA) and coombs anti-*Brucella* test were 1:80 and 1:2560, respectively. The patient was treated with a combination of doxycycline 200 mg twice daily for six weeks and streptomycin 1gm IM for two weeks. He remained asymptomatic during a six-week follow-up period with no evidence of relapse.

Discussion

Focal forms of brucellosis are present in approximately 30% of patients [3]. The most common gastrointestinal complication is reactive

hepatitis with or without granulomas [4]. Other less frequent complications include peritonitis [[5], intraabdominal abscesses [6], ileitis [7], colitis [8], pancreatitis [9], and appendicitis [10]. Cases of acute cholecystitis occurring as a complication of brucellosis are rare. In a review of literature (MEDLINE 1934-2005), only 16 cases have been reported to date (Table 1). As in our case, most reported cases are due to *B. melitensis* and describe both lithiastic and acalculous cholecystitis. Twelve of the sixteen cases were male patients and four were females. The average age was 49 years (range 6-72 years). All the patients had clinical symptoms of acute cholecystitis and had history of fever days or weeks prior to admission. Six patients had *B. melitensis* and one had *Brucella abortus* isolated in the bile. In one patient *Brucella suis* was isolated from both blood and bile cultures [11]. In six

patients, both blood and bile cultures were positive at the time of the infection. In five of the cases diagnosis was made on growth of the microorganism in the blood. Gallstones were present in nine patients. Thirteen patients underwent cholecystectomy. Histopathological examination of the gallbladder showed signs of acute and/or chronic inflammation. Three cases showed the presence of granulomas in the gallbladder wall [2,10,12]. Five patients had a brucellosis risk factor. Three patients had history of contact with animals [2,14,15] and one was a microbiologist [15]. In one case reported by Fasquelle [12], brucellosis was linked to contaminated milk and dairy products. In our patient brucellosis was not suspected at the time of admission. The diagnosis was reached only after blood cultures were performed due to the patient's fever. On further questioning of the patient, he gave a history of ingestion of raw milk one month prior to his illness.

The most commonly used antibiotic combination was doxycycline/streptomycin and tetracycline/streptomycin. The duration of therapy ranged from eight days to six months. All patients made uneventful recovery.

Brucella species, as in the case of *Salmonella* species, are usually associated with bacteremia and systemic infection. Both organisms are intracellular and may cause latent infection with subsequent clinical symptoms months or years after their onset. Involvement of the gallbladder in such systemic infections may occur via the lymphatic spread or as part of bacteremia. However, unlike *Salmonella* species in which the gallbladder acts as a reservoir for the microorganism, no reports on chronic carriage of *Brucella* in the gallbladder has been made. Localized brucellosis may result as a complication of bacteremia or may be the only manifestation of chronic infection [11]. In this case, it is unclear whether acute cholecystitis associated with brucellosis is a complication of a chronic latent infection or simply a localized form of acute brucellosis.

In conclusion, brucellosis should be considered in the differential diagnosis of acute cholecystitis in regions where brucellosis is an endemic disease.

References

1. Matyas Z and Fujikura T. Brucellosis as a world problem (1984) *Dev Biol Stand* 56: 3-20.
2. Colmenero JD, Reguera JM, Martos F, Sanchez-De-Mora D, Delgado M, Causse M, Martin-Farfan A, Juarez C (1996) Complications associated with *Brucella melitensis* infection: a study of 530 cases. *Medicine (Baltimore)* 75: 195-211.
3. Corbel MJ (1997) Brucellosis: an overview. *Emerg Infect Dis* 3: 213-221.
4. Cervantes F, Bruguera M, Carbonell J, Force L, Webb S (1982) Liver disease in brucellosis: A clinical and pathological study of 40 cases. *Postgrad Med J* 58: 346-350.
5. Hatipoglu CA, Yetkin A, Ertem GT, Tulek N (2004) Unusual clinical presentations of brucellosis. *Scand J Infect Dis* 36: 694-697.
6. Williams RK and Crossley K (1982) Acute and chronic hepatic involvement of brucellosis. *Gastroenterology* 83: 455-8.
7. Orte L, Teruel JL, Bellas C, Traver JA, Sanz-Guajardo D, Anaya A, Botella J (1979) Brucellosis of the kidney: description of three cases. *Rev Clin Esp* 152: 461-464.
8. Jorens PG, Michielsens PP, Van den Enden EJ, Bourgeois NH, Van Marck EA, Krueger GR, Ramon AM, Maercke YM (1991) A rare case of colitis-*Brucella melitensis*: Report of a case. *Dis Colon Rectum* 34: 194-196.
9. AL-Awadhi NZ, Ashkenani F, Khalaf ES (1989) Acute pancreatitis associated with brucellosis. *Am J Gastroenterol* 84: 1570-574.
10. Andriopoulos P, Tsironi M, Asimakopoulos G (2003) Acute abdomen due to *Brucella melitensis*. *Scand J Infect Dis* 35: 204-205.
11. Morris SJ, Greenwald RA, Turner RI, Tedesco FJ (1979) *Brucella*-induced cholecystitis. *Am J Gastroenterol* 71: 481-484.
12. Méttier SR and Kerr WJ (1934) Hepatitis and cholecystitis in the course of brucella infection. *Archives of Internal Medicine* 54: 702-705.
13. Fasquelle D, Charignon G, Rems M (1999) Acute calculous cholecystitis in a patient with brucellosis. *Eur J Clin Infect Dis* 8: 599-606.
14. Valenzuela Casas (1971) E. Colecistitis hemorrágica melitocócica. *Rev Esp Enf Ap Digest* 33: 723-726.
15. Miranda RT, Gimeno AE, Rodriguez TF, de Arriba JJ, Olmo DG, Solera J (2001) Acute cholecystitis caused by brucella melitensis: case report and review. *J Infect* 42: 77-78.
16. White CS (1934) Cholecystitis as a complication of brucellosis: report of a case. *Medicine Annals D.C.* 54: 702-705.
17. Berbegal Serra JM, Rodriguez Alfaro JJ, Royo Garcia G (1986) Brucellosis complicada con colecistitis aguda alitiasica *Revista de las Enfermedades del Aprato Digestivo* 70: 88.
18. Shaheen SE, EL Taweel AZ, AL Awadi NZ, Nassrula AY, Marzouk MM, Ghafoor MA (1989) Acute calculous cholecystitis associated with brucella melitensis. *American Journal of Gastroenterology* 84: 336-337.
19. Serrano FJ, Sanchez MA, Valdivieso P, Gonzalez P (1999) Colecistitis aguda en el curso de brucellosis sintamica. *Rev Clin Esp* 199: 621-662.
20. Ashley D, Vade A, Challapalli M, Brucellosis with acute acalculous cholecystitis(2000). *Pediatr Infect Dis J.* 19: 1112-1113.
21. Lopez-Prieto MD, Aller AI, Alcaraz S, Gutierrez de la Pena C (2003) *Enferm Infecc Microbiol Clin.* 21: 464-5.
22. Kanafani ZA, Sharara AI, Issa IA, Kanj SS (2005) Acute calculous cholecystitis associated with brucellosis: A report of two cases and review of the literature. *Scand J Infect Dis.* 37: 927-30.

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Conflict of interests: No conflict of interests is declared.