Brucellosis Reactivation after 28 Years

To the Editor: Approximately 10% of patients with brucellosis experience a relapse, 90% of which occur within a year after discontinuation of antimicrobial drug therapy (1,2). Here we report a patient who had brucellosis in a disease-endemic area, emigrated to a brucellosis-free region, and experienced focal reactivation in her gallbladder 28 years later. To our knowledge, this interval is among the longest reported asymptomatic intervals between a first brucellosis episode and reactivation. The case suggests that physicians should not disregard remote histories of brucellosis and past residence in brucellosis-endemic areas when confronted with possible reactivation disease.

A woman, born in 1955, had prolonged fever without focal symptoms in 1981 and had received a diagnosis of brucellosis while living near Valencia, Spain. The brucellosis was attributed to an episode of eating unpasteurized cheese from the local dairy and was successfully treated. She immigrated to Switzerland in 1990 and was well until March 2009, when malaise and right upper quadrant pain developed, without fever. She was otherwise healthy and did not take any medication. Computed tomography (CT) scan showed a mass contiguous with the gallbladder, extending intrahepatically, with concentric calcifications and multiple gallstones (Figure). Gallbladder cancer was suspected, but when a laparotomy was performed, an acutely inflamed gallbladder with a surrounding inflammatory mass was found and excised. Gallbladder cultures on standard media (Columbia agar with 5% sheep’s blood, chocolate agar, Brucella blood agar, and brain-heart infusion broth) were discarded when they remained sterile after 5 days of incubation. Histopathologic examination showed granulomatous cholecystitis, and the patient was referred for infectious disease consultation. Formalin-fixed gallbladder tissue was negative for Mycobacterium tuberculosis complex DNA but positive for Brucella melitensis by PCR (3). Blood cultures (BACTEC Plus Aerobic/F and Anaerobic/F [Becton Dickinson, Allschwil, Switzerland], incubated for 10 days) remained sterile. No Brucella DNA was detected in blood and serum (1,2), and a rose bengal serum agglutination test was negative for anti-Brucella antibodies. Because of the rarity of the manifestation (late reactivation) and location (gallbladder), plus a residual abscess shown on CT scan 8 weeks after surgery, prolonged treatment with doxycycline and rifampin was administered for 3 months, with gentamicin added during the initial 2 weeks (4). Nine months after antimicrobial drug therapy was discontinued, the patient remains well.

Switzerland reported the elimination of animal brucellosis in 1963 (5) and has officially been brucellosis-free since 1998, according to article 14.1.2. of the Terrestrial Animal Health Code (www.oie.int/eng/normes/mcode/en_chapitre_1.14.1.htm). The last case of B. melitensis infection in a sheep or goat in Switzerland was reported in 1985. The annual number of human brucellosis cases reported in
Switzerland remains low but seems to be increasing; 3, 5, and 13 cases were recorded in 2007, 2008, and 2009, respectively (6).

Most of these cases were linked to international travel or ingestion of food products imported from disease-endemic areas, e.g., Turkey. Our patient traveled annually to her family’s home town in Spain, but she vigorously denied exposure to any dairy or meat products imported from an area with known brucellosis endemicity or to any unpasteurized dairy products in general since she left Spain in 1990. Therefore, the most likely explanation was brucellosis reactivation, 28 years after the initial episode in Spain, where brucellosis was endemic. In the absence of a positive culture, brucellosis remains formally unproven. However, the granulomatous inflammation in the resected gallbladder tissue, the absence of another identified cause (e.g., tuberculosis), and the patient’s clinical and radiologic response to specific anti-Brucella treatment argue against the positive PCR result being merely attributable to long-term, persistent, but clinically latent, brucellosis.

Ariza (7) and Diaz (8) have reported patients with chronic hepatosplenic brucellosis after remote episodes of brucellosis (2–40 years previously). Because these patients lived in disease-endemic areas, reinfection cannot be excluded. Brucella organisms persist intracellularly, and their DNA has been detected in the peripheral blood of asymptomatic patients, even years after the diagnosis of clinical brucellosis and its clinically successful cure. These facts suggest that the outcome of treated brucellosis may be a chronic, persistent, asymptomatic infection, rather than complete bacterial eradication (9).

The pathogenesis of unusually late brucellosis reactivation in our patient remains obscure. Reactivation has only rarely been linked to underlying illnesses or immunosuppression (2). Rare cases of reactivation have been reported during pregnancy. Brucella persistence has been recorded inside calcified lesions in chronic hepatosplenic brucellosis (7), and adherence to foreign bodies such as joint prostheses or prosthetic heart valves and other cardiac devices may occur, perhaps in association with episodes of transient Brucella bacteremia (10). Unlike the situation with typhoid fever, no evidence links the presence of gallstones to the persistence of Brucella organisms in humans.

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DOI: 10.3201/eid1612.100678

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