A 15-year-old boy, who had immigrated to Canada from the Philippines at the age of 11, presented to the emergency department with fever and acute onset of periumbilical abdominal pain that migrated to the right lower quadrant over 24 hours. Before this, he had experienced four months of intermittent abdominal pain, which occurred about weekly, but no fever or weight loss. On examination, he was focally tender over the McBurney point. He had a normal hemoglobin level of 150 (normal 137–180) g/L, and elevated white blood count of 12.8 (normal 4.0–11.0) $\times 10^9$/L and neutrophil count of 10 (normal 2.0–8.0) $\times 10^9$/L. Ultrasonography of the abdomen showed an enlarged appendix leading into a bi-lobed, complex right lower quadrant collection. He underwent a laparoscopic appendectomy for acute perforated appendicitis. At the time of surgery, the appendix was found to be thickened and inflamed. A retroiliac abscess was also found, filled with thick creamy contents that resembled cottage cheese. The appendix was removed and the abscess drained. The patient was given ampicillin, gentamicin and metronidazole for six days and then amoxicillin–clavulanate for another five days. His recovery was uneventful. When seen in follow-up six weeks postoperatively, the patient was doing well and had no specific symptoms. The pathology report of the appendix, however, described florid necrotizing granulomatous inflammation with extension into the mucosa, and tuberculosis (TB) was suspected (Figure 1).

The patient was referred to the infectious diseases clinic for further assessment. He described a mild cough 48 hours before the surgery, which recurred five to six weeks postoperatively. His review of systems was otherwise unremarkable, and he had no history of chronic fever or night sweats. Before emigrating from the Philippines four years earlier he had lived with his grandmother, who had received treatment for pulmonary TB seven years earlier.

Given our strong suspicion of TB, we performed a Mantoux test, which was strongly positive at 17 mm. Chest radiography showed ill-defined parenchymal opacities and multiple bilateral upper-lobe nodules scattered throughout both lung fields (Figure 2). An induced sputum sample tested positive for acid fast bacilli.

We diagnosed multifocal tuberculosis. The patient was referred to the regional TB clinic and started on four-drug therapy with isoniazid, rifampin, ethambutol and pyrazinamide. He was confirmed to be HIV-negative. Mycobacterium tuberculosis was isolated from his sputum cultures. Susceptibility tests showed that the isolate was resistant to isoniazid, and his treatment was tailored appropriately. Results of repeat sputum smear and culture were negative within two months, and repeat chest radiography was normal within five months. He tolerated his treatment well and successfully completed nine months of therapy with complete resolution of symptoms.

Discussion

The most recent TB report from the Public Health Agency of Canada showed a TB incidence of 4.6 per 100 000 population in Canada. Of the reported TB cases, 66% occurred in foreign-born individuals, 21% in Canadian-born Aboriginal people and 12% in Canadian-born non-Aboriginal people. The World Health Organization has identified 22 developing countries with a high burden of TB that account for about 80% of all new TB cases arising each year. In the case described, the patient emigrated from a TB-endemic region and had a known family contact who had received treatment for TB in the past. Immigrants to Canada...
Tuberculosis of the appendix

Abdominal TB complicates untreated pulmonary TB in 6%–38% of cases.5 Any organ of the gastrointestinal tract can have TB involvement, but the ileocecal region with its rich lymphoid tissue is the most common area involved. Three pathways have been described to cause occurrence of TB in the appendix: a direct lymphohematogenous route, an intestinal route and direct spread from peripheral organs.6 First described in 1837, appendicular TB is quite rare, with about 70 case reports published in English.7

Foreign-born individuals from TB-endemic regions and Aboriginal Canadians have a higher risk of TB infection, as do close contacts of an individual with known TB disease. Tuberculosis should be considered in all patients with abdominal pain, fever and non–liver related ascites, if they fit the epidemiological profile noted above.3 Intraoperative clues may also be present. Tissues that are necrotic and have a “cottage-cheese” appearance similar to peritoneal carcinosis may indicate the presence of gastrointestinal and disseminated TB.3 Suspicion of TB appendicitis may also be raised by the patient’s postoperative course. If, despite use of appropriate intravenous antibiot-
ics and absence of an inflammatory focus, the patient continues to have prolonged peritoneal drain discharge and fever, TB involvement should be considered. The patient described in this case report had an uneventful postoperative course.

Granulomatous appendicitis is a rare entity, appearing in only 0.1%–2% of all appendectomy specimens. A four-year review of 3381 appendectomy specimens showed only 13 cases (0.38%) of granulomatous appendicitis; of those, only one was believed to have been caused by TB. Other infectious causes for a granulomatous appendix include infection with *Yersinia enterocolitica* and parasitic infections (such as trichinosis and enterobiasis), whereas non-infectious causes include Crohn disease, sarcoidosis, diverticulitis and foreign body.

The treatment for appendicular TB is both surgical and medical. In conjunction with appendectomy, these patients must be given multiple antituberculous antibiotics as per standard guidelines. It is crucial that patients with appendicular or extrapulmonary TB be assessed for pulmonary involvement with chest radiography and sputum collection. This can help determine the risk of transmission to others, as well as guide therapeutic decisions and aid with diagnosis, as occurred in this case.

### References


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