Perforation of the jejunum secondary to AIDS-related gastrointestinal Kaposi’s sarcoma

ERIC M YOSHIDA MD FRCPC, NORMAN HL CHAN MD FRCPC, CLIFFORD CHAN-YAN MD FRCPC, ROBERT M BAIRD MD FRCSC

Kaposi’s sarcoma (KS) is a multicentric malignant vascular neoplasm of unknown cause. The classic form of this disease is an indolent cutaneous neoplasm that afflicts mainly elderly Mediterranean and Jewish men. The endemic form of this disease manifests as a locally aggressive neoplasm found in Africans (1). This disease is now well known to clinicians as a common manifestation of the acquired immunodeficiency syndrome (AIDS). As a manifestation of AIDS, KS may be limited to the skin or may present as aggressive disseminated disease with visceral involvement. Involvement of the gastrointestinal tract by KS occurs in up to 40% of patients with AIDS (2). Gastrointestinal KS is rarely symptomatic, but can occasionally present as gastrointestinal bleed (3). Reported instances of intestinal perforation due to KS have been rare. We report a human immunodeficiency virus (HIV)-positive man with cutaneous KS who had a perforation of the jejunum due to KS. The current literature pertaining to this subject is reviewed.

CASE PRESENTATION

A 36-year-old HIV seropositive homosexual man, with a three-year history of cutaneous KS and primary pulmonary hypertension, presented with recent onset of cramp-like mid-abdominal pain and frequent loose stools. He did not have fever, chills, nausea or vomiting. Background history revealed that his CD4 count two
months previously was 140/mm³. Cutaneous KS was the only AIDS defining illness. There was no history of opportunistic infections. His pulmonary hypertension had been investigated three years previously and was felt to be primarily AIDS-related. Medications at admission included zidovudine, didanosine, cotrimoxazole, erythromycin, digoxin and diltiazem. There was no history of nonsteroidal anti-inflammatory drug use.

Physical examination revealed an acutely distressed patient. Auscultation of the chest was unremarkable. Examination of the cardiovascular system revealed jugulovenous distension 10 cm above the sternal angle, a right parasternal heave, displaced apical impulse, a right-sided third heart sound and a II/VI pansystolic murmur at the left sternal border. Examination of the abdomen revealed diffuse tenderness and decreased bowel sounds. Dermatological survey revealed lesions consistent with KS on the forehead and upper body.

Initial laboratory data revealed the following: leukocyte count, 4.5x10⁶/L; hemoglobin, 131 g/L; platelet count, 171x10⁶/L; normal serum electrolytes; serum creatinine, 90 /c10⁹ mol/L (normal 40 to 120); aspartate aminotransferase, 52 U/L (normal greater than 40); alkaline phosphatase, 321 U/L (normal 35 to 105); and serum lactate, 6 mmol/L (normal 0.5 to 1.8). Abdominal radiographs revealed the presence of free air.

Intraoperatively, five slightly raised pigmented lesions consistent with KS were seen in the jejunum and ileum. One of the lesions in the jejunum was noted to have a perforation. Resection of the affected bowel was undertaken followed by primary reanastomosis.

Gross pathological examination of the jejunum revealed a 1 mm perforation in an ulcerating KS lesion. Histopathological examination of the solid area surrounding the perforation revealed a malignant spindle cell neoplasm with numerous vascular clefts, features typical of KS (Figures 1, 2). Special stains, including a rhodamine-auromine fluorescent stain for mycobacteria, periodic acid-Schiff stain for fungus, methenamine silver stain for Pneumocystis carinii and in situ hybridization for cytomegalovirus (CMV), were all negative. The final pathological diagnosis was perforation of the jejunum secondary to KS. The patient died of respiratory complications in the intensive care unit following surgery.

DISCUSSION

AIDS-related KS can involve every aspect of the gastrointestinal tract from the oropharynx to the rectum. Hepatobiliary, pancreatic and splenic involvement have also been described (3). KS of the gut is typically asymptomatic. Endoscopically, gastrointestinal KS appears as purple nodules, polyloid masses or hemorrhagic macules. These lesions are frequently submucosal and may be entirely missed on endoscopic biopsy (2). Other manifestations of gastrointestinal KS include diarrhea and intussusception (4), appendicitis (5) and obstruction due to bulky disease (3), as well as symptoms referable to nonintestinal intra-abdominal involvement, such as cholangitis secondary to biliary involvement (6).

Whereas intestinal perforation is a well described complication of gastrointestinal CMV infection (7,8), intestinal perforation due to KS alone has rarely been reported. The first case was reported by Mitchell and Feder (9) in 1949. They described an elderly Jewish man who had obstructive symptoms and developed an acute abdomen while straining at stool. Postmortem examination revealed bulky KS encircling the jejunum with perforation. In the AIDS era there have been several reports of intestinal perforation associated with gastrointestinal KS. Cases of KS presenting with duodenal perforation (10) and ruptured appendicitis (5) have also been reported. However, the etiology of the perforation in most of these cases is not attributed to KS alone. Perforation of the terminal ileum with KS in close proximity to the perforation and with coexisting CMV vasculitis was reported by Burke et al (11). Ileal perforation with mycobacteria and candida within a KS mucosal ulceration was also reported by Bieluch and colleagues (12). These authors believed that superinfection of the KS lesion was a contributing factor leading to perforation. Intestinal mycobacterial infection has
been reported to result in bowel perforation (13). Intestinal lymphoma is yet another important cause of AIDS-related small bowel perforation (8,14).

Our patient with AIDS-related KS had a jejunal perforation secondary to KS but did not have bulky disease, making this case significantly different from the patient reported by Mitchell and Feder (9). In our patient this perforation was a direct result of central ulceration within a small KS lesion. There were no other identifiable causes, either infectious or neoplastic, to cause this small bowel perforation.

Another interesting AIDS-related illness in our patient was primary pulmonary hypertension. Pulmonary hypertension has been recognized to be associated with HIV, although its occurrence is rare (15). Its association with our patient’s intestinal perforation was coincidental, but was a major factor in his death.

CONCLUSIONS
This case demonstrates that intestinal perforation can be secondary to KS. Unlike the original case reported over 40 years ago in the pre-AIDS era, gastrointestinal KS does not have to manifest as bulky disease with resulting obstruction in order to cause perforation. KS itself should be considered in the differential diagnosis of any HIV-seropositive patient presenting with a bowel perforation. This may have therapeutic and clinical relevance.

REFERENCES