Inflammatory bowel disease and superior mesenteric artery thromboembolism

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While patients with inflammatory bowel disease are known to be at increased risk of venous thromboembolism, the risk of arterial thrombosis is less well recognized. Here, we describe the case of a middle-aged female with a recent diagnosis of Crohn’s disease who presented to her local emergency department with acute abdominal pain. Subsequent investigations revealed a thrombus in the superior mesenteric artery resulting in multi-organ infarction requiring major intra-abdominal surgery and extensive resection of segments of small and large bowel. (Intest Res 2020;18:130-133)

Key Words: Inflammatory bowel disease; Ischemia; Thromboembolism

INTRODUCTION

Inflammatory bowel disease (IBD), particularly CD, is characterized by a state of chronic inflammation that can involve any segment of the GI tract. Furthermore, this inflammatory state can influence multiple organ systems external to the bowel. These include but are not limited to the musculoskeletal, ocular and vascular systems. One well documented systemic complication is the propensity for the development of venous thromboembolism. Less well documented or understood is the increase in risk of arterial thromboembolism. Here, we describe a case of a patient, recently diagnosed with CD who subsequently presented with acute severe abdominal pain secondary to a superior mesenteric artery thromboembolism.

CASE REPORT

A 48-year-old female patient was diagnosed with ileocolonic CD after experiencing months of abdominal discomfort, loose stools and bloating. Her initial colonoscopy demonstrated an inflamed intestinal mucosa with a cobblestone appearance and longitudinal ulcerations throughout the colon extending into the terminal ileum with rectal sparing. Based on these findings she was commenced on oral prednisolone. Her symptoms improved initially however upon weaning the prednisolone, she developed recurrence of symptoms. She was subsequently commenced on 6-mercaptopurine 25 mg daily and allopurinol 100 mg daily. Despite therapeutic thiopurine metabolites and 3 months of therapy, she continued to have abdominal pain and loose stools, opening her bowels 5 to 6 times a day, there was no rectal bleeding.

There was no family history of IBD, no prior use of NSAID, oral contraceptive pill or illicit drugs and no recent travel overseas. She was a current smoker with no other risk factors for cardiovascular disease and had a normal BMI. She did not display any extraintestinal manifestations of CD on history or physical examination.

She underwent a second colonoscopy which showed mild patchy colitis throughout the colon with rectal sparing (Fig. 1). Biopsy revealed patchy mild active chronic colitis. There were rare small clusters of histiocytes without well-formed granulo-
mata (Fig. 2).

Following this, she had an MRI of the small bowel to assess the extent of disease. This demonstrated a 25 cm segment of distal and terminal ileum with bowel wall thickening and enhancement. There was no proximal dilatation. A decision was made to commence adalimumab.

Two weeks later and just prior to the commencement of anti-TNF therapy, she presented to the local emergency department with severe abdominal pain, vomiting, and watery diarrhea. On examination, she was hemodynamically stable, afebrile and her abdomen was soft with a mildly tender epigastrium but no signs of peritonism. Blood tests revealed anemia with a hemoglobin 113 g/L, a mild leukocytosis 14.2 × 10⁹/L, CRP 6 mg/L, ALT 23 IU/L, GGT 31 IU/L, ALP 81 IU/L, albumin 36 g/L, lactate 1 mmol/L, and lipase of 394 IU/L.

An initial CT scan of the abdomen and pelvis did not show any evidence of colitis or bowel obstruction. However, the spleen was heterogenous and showed features consistent with splenic infarction (Fig. 3).

On day 2 of her admission, she developed worsening abdominal pain and rectal bleeding with no relief from opioid analgesia. A thrombophilia and vasculitis screen did not identify a cause and she was treated empirically with therapeutic enoxaparin. By day 4, she had on-going pain refractory to opioid analgesia so a CT angiogram was performed which revealed a nearly occlusive thrombus within the superior mesenteric artery and diffusely thickened loops of small bowel suggestive of ischemia, plus focal areas of nonenhancement within the liver suggestive of hepatic infarction (Fig. 4). There were no features to suggest aortitis.

Lactate was 2 mmol/L with a pH of 7.44. White cell count had risen to 18.8 × 10⁹/L and hemoglobin had dropped to 95 g/L, CRP peaked at 13 mg/L, ALT 179 IU/L, GGT 58 IU/L, ALP 88 IU/L and albumin 16 g/L. Extensive investigation did not show any evidence of an underlying thrombophilia including protein C or S deficiency, antithrombin III, factor V Leiden, antiphospholipid syndrome and JAK2 gene testing. Serial electrocardiography demonstrated sinus rhythm consistently (no atrial fibrillation or other arrhythmias detected) and transesophageal echocardiogram was unremarkable with no evidence of valvular, atrial or ventricular thrombus and no atrial septal defect or cardiac shunt. There were no clinical features

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Fig. 1. Colonoscopic finding. Colonoscopy demonstrating patchy colitis throughout the colon.

Fig. 2. H&E stained slides from colonic biopsies (× 200). The section shows mildly active colitis with increased mixed inflammatory cells in the lamina propria and an occasional cluster of histiocytes beneath the surface epithelium.

Fig. 3. Abdominal CT finding. CT of the abdomen demonstrates area of infarction within the spleen (arrow).
of deep venous thrombosis.

She was subsequently commenced on a heparin infusion and underwent a laparotomy, which revealed a dilated, ischemic gallbladder, ischemic caecum, and distal small bowel. An embolectomy of the superior mesenteric artery was performed and a resection of small and large bowel starting 140 cm from the pylorus to the hepatic flexure leaving 140 cm of viable small bowel remaining. She was commenced on intravenous antibiotics and 2 days later had a repeat laparotomy showing healthy small and large bowel and a small bowel to transverse colon anastomosis was performed.

Histopathology from the resected specimens was consistent with ischemic colitis (Fig. 5). Her postoperative course was uncomplicated apart from ongoing diarrhea likely secondary to short gut syndrome. She required a short period of total parenteral nutrition. Her prednisolone was gradually weaned and long-term warfarin therapy was commenced.

DISCUSSION

Thromboembolic occlusion of the superior mesenteric artery is recognized to be a common cause of acute mesenteric ischemia. Despite this there have been few reported cases of superior mesenteric artery thromboembolism related to IBD reported in the literature and most of these relate to chronic ischemia and none of the reported cases resulted in multi-organ infarction. There is one meta-analysis which looked at thromboembolic events and cardiovascular mortality in IBD. This was a systematic review of observational studies. It found that the overall risk of venous thromboembolism in IBD patients was increased almost double that of the general population (RR, 1.96; CI 95%, 1.67–2.30). Comparatively a significant increased risk of mesenteric ischemia was observed in IBD patients compared to the general population (RR, 3.46; 95% CI, 1.78–6.71). Despite this, there are still conflicting data on the risk of arterial thromboembolism in IBD. While there is an increased risk in other chronic inflammatory diseases such as rheumatoid arthritis, studies have failed to demonstrate any increase in risk of arterial thromboembolism in the IBD cohort.

The causes of acute mesenteric ischemia are usually secondary to an underlying cardiac abnormality either structural or electrophysiological. The other common cause is from any process that increases the general risk of arterial thrombosis including smoking, hypertension, hypercholesterolemia, peripheral vascular atherosclerosis, advanced age or chronic inflammatory disease affecting the vascular system such as vasculitis.

The segment of bowel affected by CD is also particularly susceptible to ischemia given the greater metabolic demand of the inflamed segment. It has previously been noted that mesenteric blood flow is increased in CD. Hence, any alteration in the supply of oxygenated blood to affected bowel could result in acute ischemia. The mechanism leading to ischemia may result from either unstable thrombus rupture secondary to increased arterial wall stress or chronic ischemia secondary to the build-up of plaque in those with multiple cardiovascular risk factors.
This case highlights the difficulty in diagnosis of ischemia in patients with IBD presenting with a suspected disease flare. Delay in diagnosis poses a substantial risk of morbidity and mortality. A high index of suspicion for ischemia is warranted in patients who present with persistent abdominal pain disproportionate to the severity of intestinal inflammation and/or unresponsive to corticosteroids and analgesia.

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CONFLICT OF INTEREST

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AUTHOR CONTRIBUTION

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