Stercoral perforation of the colon proximal to an end colostomy

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Introduction

Despite Grinvalsky's finding of an overall incidence of 4.6% of stercoral ulceration at post-mortem, stercoral perforation of the colon is a rarely reported condition. We report the first two cases of stercoral perforation occurring as a post-operative complication proximal to an end colostomy.

Case reports

Case 1

A 34 year old female presented with a 6 hour history of severe lower abdominal pain. On examination she had a temperature of 38°C and a tachycardia of 120. There was tenderness and guarding in the lower abdomen, but rectal examination was normal. Vaginal examination revealed a tender mass in the right fornix and a presumptive diagnosis of a pyosalpinx was made. Appropriate antibiotic therapy was instituted; however, after failing to improve over a 12 hour period, laparoscopy was performed. This revealed a perforation of the sigmoid colon with faecal peritonitis. Subsequent laparotomy confirmed this to be a stercoral perforation of the lower sigmoid colon, with multiple perforations along a 6 cm segment of colon. This segment was excised, the rectal stump oversewn, peritoneal lavage performed, and an end colostomy created in the left iliac fossa after manually emptying the loaded proximal colon of faecal content. The patient was discharged 7 days later after an uneventful recovery.

However, 24 hours later she returned with signs of generalized peritonitis. At laparotomy a recurrent stercoral perforation was present 6 cm proximal to the end colostomy. This segment was excised and the colostomy refashioned. Her subsequent course was uneventful and the colostomy was closed 3 months later. Pathological examination of both specimens revealed multiple stercoral perforations with ischaemic necrosis of the surrounding bowel wall. There was no intrinsic colonic pathology or vasculitis in the excised mesentery.

Case 2

A 53 year old male presented with a 9 month history of rectal bleeding, passage of mucus, tenesmus and gross constipation. On examination an extensive circumferential, stenosing, immobile carcinoma was present in the lower third of the rectum. Biopsy confirmed adenocarcinoma of the rectum. At laparotomy a massive, fixed tumour was found completely occupying the pelvis, and the proximal colon was loaded with scybala. A defecting left iliac fossa colostomy was created and a mucous fistula fashioned in the lower end of the wound to avoid obstruction in the defunctioned distal bowel. Five days postoperatively the colostomy was functioning satisfactorily. On the tenth day, however, the patient experienced colicky abdominal pain and distension, although the colostomy continued to function. Eighteen hours later generalized peritonitis was present, and plain abdominal X-rays revealed colonic distension, but...
no free gas. At laparotomy, the findings were of a diffuse faeculent peritonitis due to a stercoral perforation of the colon immediately proximal to the exit site of the colostomy. A limited sigmoid colectomy was performed, the colon manually milked of its contents and the colostomy refashioned. Four months later his condition was satisfactory. Histological examination confirmed multiple stercoral ulcers, with a full thickness perforating ulcer.

Discussion

Stercoral perforation has been reported proximal to an end transverse colostomy. However, the two cases described are the first reported cases to our knowledge occurring as a postoperative complication, proximal to an end sigmoid colostomy.

The diagnosis, as in the majority of previous reports, was not made prior to laparotomy. The first patient did not admit to constipation and radiological features of faecal loading were absent. However, there was a long history of marijuana abuse, which is a known constipative agent, and may have contributed to stercoral ulceration. Faecal impaction proximal to the stenosing tumour was recognized in the second case. However, despite careful manual emptying of the proximal colon at the initial laparotomy, this manoeuvre was not adequate to prevent stercoral perforation.

Both cases demonstrate the typical pathology of stercoral perforation, with multiple, round or ovoid, ischaemic ulcers, depressed below mucosal level, and a marked inflammatory reaction. These features distinguish the condition from idiopathic perforation where the primary pathology is a linear tear of the colonic wall, without associated inflammatory or ischaemic changes.

The principles of treatment as outlined by Guyton et al. were applicable to our cases, and involved preoperative resuscitation, elimination of all faecal soiling, resection of involved bowel with exteriorization and antibiotic therapy. Furthermore, the first case demonstrates the multifocal nature of this disease, which affects a segment rather than a focal point of the colon. This is the first reported case of recurrent stercoral perforation, and emphasizes the necessity fully to resect the diseased segment, rather than exteriorization of the perforation alone.

Intra-operative orthograde colonic lavage is used to protect colonic anastomoses, and is effective at cleaning the colon of faecal content. The two cases described have demonstrated a failure of standard techniques to deal with the problem of the loaded proximal colon, even when an end colostomy is established. It is possible that stercoral perforation might have been avoided in these two cases had intra-operative orthograde colonic lavage been undertaken at the original operations.

We therefore suggest that intra-operative orthograde colonic lavage be performed to protect a terminal colostomy where the proximal colon is loaded with scybala, and is therefore at risk of stercoral perforation.

References

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