Rectal diverticulitis mimicking rectal carcinoma with intestinal obstruction: Case report

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ABSTRACT

Although diverticular disease of the colon is common, the occurrence of rectal diverticula is extremely rare with only sporadic reports in the literature since 1911. Symptomatic rectal diverticula are seen even less frequently, and surgical intervention is needed for only complicated cases. Here we report the case of a 63-year-old woman presenting with rectal diverticulitis mimicking rectal carcinoma with intestinal obstruction.

Keywords: Rectal diverticulitis, rectal carcinoma, intestinal obstruction

INTRODUCTION

Diverticulosis is common in the Western world, manifesting as a sac of an abnormally protruding bowel wall and usually occurring in the distal colon; 90% of patients have sigmoid colon involvement (1) and only 15% have right-sided diverticula (1). Diverticulosis is an acquired disease with herniations of the mucosa and part of the submucosa through the muscularis propria due to a lack of dietary fiber, but the pathogenesis of colonic diverticula remains poorly understood.

The occurrence of rectal diverticula is extremely rare, with only a few reports published in the literature since 1911 (2). The cause of rectal diverticula is unknown; however, most are asymptomatic and do not require treatment. Surgical intervention is generally only necessary for complicated cases. In this report, we describe a rectal diverticulum that mimicked rectal carcinoma with intestinal obstruction. Low anterior resection and temporary colostomy were performed via emergent laparotomy because of complete intestinal obstruction.

CASE PRESENTATION

A 63-year-old woman was admitted into the hospital with abdominal pain, nausea, and vomiting. She had experienced chronic constipation for 10 years, but had stopped producing gas and gaita discharge 4 days prior to admission. The patient’s medical history included an appendectomy 30 years previously, coronary bypass surgery 10 years earlier, and an explorative laparotomy for a suspected left ovarian mass that resulted in a left salpingo-oophorectomy 3 years earlier. Upon physical examination, scars from a median incision and the appendectomy incision were seen. She had abdominal distension with tenderness, but did not have rebound tenderness, and her bowel sounds were increased and obstructive. During a digital rectal examination, her ampulla recti was empty. A perianal fistula suggestive of inflammatory bowel disease was not apparent, and her laboratory results were within normal ranges. A computed tomography (CT) scan was suggestive of inflammatory bowel disease was not apparent, and her laboratory results were within normal ranges. A computed tomography (CT) scan was suggestive of rectal cancer with a diffuse, irregular wall thickening that was 9-cm long and located in the superior rectum, approximately 10 cm above the anal verge (Figure 1); a rectal contrast transition above the mass was not observed (Figure 2). The proximal colon segments were dilated (maximum, 8 cm) with fecal retention due to a mechanical obstruction. Multiple pericolic lymph nodes (<10 mm in size) were detected around the mass and at the obturator chains.

With these findings, an emergent explorative laparotomy was performed for mechanical ileus and severe
bowel distension due to a rectal obstructing tumoral lesion. During surgery, a 5-cm infiltrative mass was detected adherent to the bladder and uterus, as well as neighboring severe fibrosis. A small bladder perforation occurred during the isolation of the bladder from the mass, and was sutured primarily. Because of uterine infiltration, a gynecologist participated into the operation and hysterectomy; a right salpingo-oophorectomy was also performed. After this, low anterior resection and terminal colostomy were performed. Frozen section examination revealed a 2-cm tumor-free zone to the distal surgical border.

A pathologic examination revealed a 4-cm, off white, infiltrative lesion around the rectum. Upon microscopic examination, mucosal ulceration, diffuse active chronic inflammation with abscess formation reaching to the serosal surface and removing the submucosa, and multiple diverticuli were detected; there was no evidence of rectal cancer. The microscopic examination was compatible with diverticulitis with micro-perforations (Figure 3). Ten lymph nodes were detected; all were reactional. The patient was discharged on postoperative day 5 without complications.

Written informed consent obtained from the patient for the publication of this case report.

DISCUSSION
Diverticular disease usually involves the sigmoid colon (65%), followed by the ascending colon and cecum. Involvement of the entire colon occurs in up to 10% of cases; however, rectal involvement is extremely rare, with an estimated incidence of <0.1% of the cases (3,4). Two theories have been described to explain the low incidence of rectal diverticula. The first theory suggests that the tenia coli muscle fibers, which surround the rectum, protect it from intraluminal pressures. The second theory suggests that the rectum is protected due to reduced internal pressure from accumulated feces and the reduced peristaltic activity associated with the rectum compared with that associated with the sigmoid colon. Although the cause of rectal diverticula remains unknown, possible predisposing factors have been described, including weakness in the circumferential muscles surrounding the rectum, primary muscle atrophy, absence of supporting structures such as the coccyx, a relaxed rectovaginal septum, constipation or recurrent impaction causing rectal distension, and rectal trauma or infection that leads to a weakened rectal wall (5,6).
Most patients with rectal diverticula are asymptomatic and diagnosed incidentally, and do not require treatment. However, rectal diverticula may become inflamed with impacted feces resulting in abscess formation and perforation. Other complications associated with rectal diverticula include rectal strictures, rectovesical fistulas, rectal prolapses, and misdiagnosis of rectal carcinoma (2). Surgical intervention is necessary only in complicated patients, and operative treatment includes draining the abscess, a diverting colostomy, resection of the diverticular mass, or rectal resection.

Radiologically, differentiating diverticulitis from colon cancer is difficult. Although they are completely different pathologic entities, they show similar segmental colonic wall thickenings, with varying degrees of pericolonic infiltration on CT. Pericolic infiltration, lymph node visibility and size, inflamed and non-inflamed diverticula, degree and pattern of enhancement, intestinal obstruction, involved wall thickness and length, fascial thickening, and free air and abscess formation have been determined to be significant parameters for differentiating diverticulitis from colon cancer (7). Diverticula tend to be longer and thinner, with more pericolonic inflammatory changes, and are minimally enhanced lesions that show a target or homogenous enhancement pattern (7). Usually, the lymph nodes associated with colon cancer are visible and >10 mm in diameter (7). Sometimes the identification of diverticulosis is challenging because of severe colonic structure destruction or the presence of fluid-filled intramuscular diverticula, and the inflammatory process appears as an inflammatory mass (7). Inflamed and non-inflamed diverticula are the most reliable signs for the diagnosis of diverticulitis, whereas coincidental diverticula have been reported in 14-29% of colon cancer cases (7). Colonic obstruction is another significant factor for differentiating diverticulitis from colon cancer, with a reported incidence of 3.8-23% in colon cancer patients (8). In diverticulitis, the chronic inflammatory process results in the intramuscular fibrosis that is responsible for the lumen rigidity, and colonic obstruction occurs in 10-20% of cases (7).

In conclusion, rectal diverticula are extremely rare, and usually do not require treatment; surgical intervention is only necessary in complicated cases. The final diagnosis, without postsurgical pathologic confirmation, is very difficult. Although CT enables accurate diagnosis of diverticulitis, colon cancer remains difficult to exclude. The presence of diverticula, absence of lymph nodes, and minimal or target pattern enhancement are the major CT findings used for discriminating between diverticulitis and colon cancer. Rectal diverticulitis should also be considered in the differential diagnosis of patients admitted with rectal masses and mechanical ileus.

**Ethics Committee Approval:** Ethics committee approval was received for this study

**Informed Consent:** Written informed consent obtained from the patient.

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