CASE REPORT

A 29-year-old male presented in September 1998 with a sudden-onset lower abdominal pain accompanied by episodes of nausea and vomiting. Past history revealed ileus one year earlier. These symptoms worsened gradually, and he was admitted to our hospital for further evaluation. On physical examination, the abdomen was flat and soft with tenderness in the lower abdomen. Laboratory data demonstrated mild anemia (hemoglobin 11.5g/dl) but other values were within normal limits.

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Plain X-ray of the abdomen showed intestinal obstruction with dilated small bowel loops and air-fluid levels suggesting small bowel obstruction (Fig. 1). He
was treated by naso-jejunal tube. X-ray examination of the small intestine with gastrografin showed a "beak-like" filling defect in the terminal ileum (Fig. 2). Ultrasonography failed to show a "target-like" appearance. Abdominal CT showed cecal wall thickening (Fig. 3). Barium enema showed filling defect in the cecum (Fig. 4). Colonoscopy revealed edematous and reddish mucosa but no tumor was found in the cecum (Fig. 5), and biopsy showed no evidence of neoplasia. These finding suggested that the patient had an intussusception.

At laparotomy, the intussusception was spontaneously resolved but there was focal thickening of the serosa at the distal 20 cm of the terminal ileum. Therefore, partial ileocolectomy was performed to prevent a causing of recurrent ileus. Grossly, the mucosa appeared intact. Microscopically, the lesion extended through the muscularis propria into the serosa (Fig. 6A). The lesion was composed of fibrous stroma and chronic inflammatory cells including lymphocytes, plasma cells (Fig. 6B) with calcification (Fig. 6C) providing a diagnosis of inflammatory pseudotumor. Postoperative course was satisfactory, and the patient remains in good health two years after the operation.

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Fig. 2. X-ray examination of the small intestine with gastrografin showing the so called "beak-like" filling defect in the terminal ileum.

Fig. 3. Abdominal CT showing cecal wall thickening.

Fig. 4. Barium enema showing a filling defect in the cecum.

Fig. 5. Colonoscopy showed an edematous and reddish mucosa with no tumor in the cecum.
Microscopically, the lesion extended through the muscularis propria into the serosa (A) (×10). The pseudotumor was composed of fibrous stroma and a mixture of chronic inflammatory cells including lymphocytes, plasma cells (B) (×200), and contained areas of calcification (C) (×200).

Discussion

Intussusception in adults is almost invariably caused by some preexisting lesions, including benign tumors, such as leiomyoma, hamartoma, and neurofibroma, or malignant neoplasms. Furthermore, trauma, post-operative adhesion, Meckel’s diverticulum, and lymphoid hyperplasia are other potential predisposing factors in the production of intussusception. Inflammatory pseudotumor is not a true neoplasia. It occurs rarely in the small intestine, and a solitary or polypoid lesion may present clinically as intussusception and bowel obstruction. In the small intestine inflammatory pseudotumors are commonly located in the mucosal layer of the ileum. In our case, the pseudotumor was located in the muscularis propria and serosa, that are rare sites for this tumor. The true nature of pseudotumor in the present case is obscure. There are several pathoetiologic factors as for inflammatory pseudotumor reported in the literature. Although the microscopic findings were similar to the post-operative change, he has no history of abdominal operation. Ramsden et al. reported a 76-year-old man...
with recurrent intussusception associated with vascular proliferation. The present case did not have such a vascular change. One of the histologic feature in our case is calcification within the tumor. Entero-colic calcification is probably the most common radiographic manifestation of schistosomal infestation of the gastrointestinal tract. Kimura et al. reported a patient with a complete invagination of the cecum that contained two large laminated calcified fecaliths. Most of the eggs of the schistosomioma are deposited in the submucosa, whereas serosal calcification is rare. In the present case, there were no granulomas, fecalis nor pathogens. Subsequently, etiology of the pseudotumor was not determined.

Mesenteric panniculitis is an extremely rare disease in which the adipose tissue is replaced by fibrosis, necrosis, and calcification. However, our case did not show any of the features of mesenteric panniculitis.

In conclusion, we have described a rare case of intussusception caused by subserosal inflammatory pseudotumor of the terminal ileum, although the etiology of the lesion remains obscure.

References