Cecal Intussusception in an Adult with Cronkhite-Canada Syndrome Relieved by Colonoscopy

Emi Ishikawa¹, Masatoshi Kudo¹, Yasunori Minami¹, Kazuomi Ueshima¹, Satoshi Kitai¹ and Kazuki Ueda²

Abstract

Cronkhite-Canada syndrome (CCS) is a rare, noninherited gastrointestinal polyposis syndrome associated with characteristic ectodermal abnormalities. Here, we report a case of Cronkhite-Canada syndrome with cecal intussusception relieved by colonoscopy. A 52-year-old man who was diagnosed as CCS pathologically two years previously presented abdominal pain and sub fever-up. Physical examination revealed the palpable mass sized approximate 10 cm in diameter in the upper abdominal site, in addition to the symptoms of alopecia, absent fingernails and toenails. However, abdominal wall rigidity and rebound tenderness were never expressed. Abdominal plain CT showed concentric circles from the ascending to the middle of the transverse colon, and a tumor in the lumen at the middle of the transverse colon. Colonoscopic reduction was performed first because we diagnosed it as intussusception due to CCS polyps without peritoneal irritation, and his symptoms were improved dramatically after careful reduction. Therefore, he was able to undergo the laparoscopic ascending colectomy as scheduled.

Key words: Cronkhite-Canada syndrome, intussusception, colonoscopic reduction

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Introduction

The patients with sigmoid volvulus or intussusception require an emergent surgery or colonoscopy to avoid a necrosis or perforation of digestive organs. The cause of intussusception in adulthood is often a tumor in the colon or small intestine (1). Therefore, some investigators have pointed out the risks of perforation and cancer cells seeding into the abdominal cavity in patients with advanced cancers caused by colonoscopic reduction (2-4). However, colonoscopic reduction may be able to reduce the necessity of an emergent surgery in intussusceptive patients with low malignant potentials and no peritoneal irritation sign.

Cronkhite-Canada syndrome (CCS) is a rare acquired gastrointestinal polyposis syndrome of unknown etiopathogenesis, accompanied by alopecia, cutaneous hyper pigmentation, onychodystrophy, diarrhea, and dysgeusia (5). We report a case of Cronkhite-Canada syndrome presenting cecal intussusception relieved by colonoscopy.

Case Report

After several days of abdominal pain and low-grade fever, a 52-year-old man presented at our hospital. This patient had been seen regularly at our hospital during the past two years as an outpatient for follow-up of CCS. He had taken cox-2 inhibitor orally, but did not have a history of steroid use because of no phenomenon of protein-losing enteropathy and malnutrition. The present physical examination revealed a palpable mass approximately 10 cm in diameter in the upper abdomen, as well as symptoms of alopecia and missing finger and toenails. However, no abdominal wall rigidity or rebound tenderness was observed. Laboratory data indicated normal electrolytes and kidney and liver function. His he-

¹Division of Gastroenterology and Hepatology, Department of Internal Medicine, Kinki University School of Medicine, Osaka-Sayama and ²Department of Surgery, Kinki University School of Medicine, Osaka-Sayama

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Correspondence to Dr. Masatoshi Kudo, minkun@med.kindai.ac.jp

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moglobin was 12.9 g/dL, white blood cell count was 4,100/mm³, CRP 0.063 mg/dL. Serum proteins were 6.4 g/dL and albumin was 3.7 g/dL. Serum CEA and CA19-9 were 1.1 ng/mL and 10 IU/mL, respectively. Plain abdominal CT showed a pattern of concentric circles from the ascending colon to the middle of the transverse colon, where a tumor was lodged in the lumen (Fig. 1). There were no signs of severe ischemia or necrosis at the base and the head of the intussusception.

Based on our diagnosis of intussusception due to CCS polyps, but with no peritoneal irritation, we performed colonoscopic reduction with the support of a surgeon. Colonoscopy revealed a focus of redness in the middle of the transverse colon that centered on a large polyp with an irregular surface (Fig. 2a). Fluorography with meglumine sodium amidotrizoate during colonoscopy showed an obstruction that did not fill the proximal transverse colon (Fig. 2b). We took care that the scope was not pushed against consistent rigidity, and used the handling of air insufflation efficiently. Therefore, with careful handling of the colonoscope, we confirmed that the intussusceptions were relieved (Fig. 3a). Images also revealed a large and irregular polyp in the cecum (Fig. 3b), and small polyps in the colon.

The patient improved dramatically after intussusception reduction, however he underwent laparoscopic ascending colectomy as scheduled due to relapses. The main polyp in the cecum was grounded with a thick stalk that had an irregular, granulated surface. Dilative cystic ductal structures and a severe infiltration of inflammatory cells were confirmed pathologically for this polyp, but no colon cancer cells were detected. Thereafter, the patient showed no postoperative symptoms and remained in good condition for one year after surgical operation.

**Discussion**

CCS is characterized by the presence of diffuse gastrointestinal (GI) polyposis, dystrophic changes in the fingernails, alopecia, cutaneous hyperpigmentation, diarrhea, weight loss, abdominal pain, and other GI complications such as...
Intussusception itself in adults occurs relatively rarely; however, a specific lead point is identified in more than 90% of cases (14, 15). A correct and timely diagnosis is not only necessary to avoid the complications of bowel infarction and perforation secondary to high-grade obstruction but also to resect the underlying lesion that serves as a lead point. Therefore, knowledge of the imaging spectrum and the clinical features of intussusception are important because imaging plays a crucial role in the diagnosis and management of these patients. Typical intussusception is well diagnosed on CT, which shows a pathognomonic bowel-within-bowel configuration (15, 16), appears as a sausage-shaped mass when CT images is obtained parallel to its longitudinal axis of digestive tract but as a target-like mass when CT images is perpendicular to the cross sections of digestive tract (15, 16). Sonography can facilitate the diagnostic decision when the characteristic sign of a target like lesion or bull’s eye lesion is shown, similar to the CT findings (15). We could diagnose the intussusception in this patient immediately by these typical findings of imaging.

Intussusceptions associated with CCS have been reported as a case report in at least four patients in Japanese language up to 2007 (17-19) in spite of no report in the English language literature. According to these literatures, intussusceptions had occurred at the ascending colon/cecum. This patient underwent laparoscopy as a therapeutic procedure, in which the intussusception due to CCS polyps that was relieved by colonoscopic reduction. Therefore, surveillance laparoscopy should be performed regularly.

Several reports have discussed intussusceptions associated with Peutz-Jeghers syndrome, a kind of disease of multiple polyposis (21-23). Peutz-Jeghers syndrome is characterized by hamartomas throughout the gastrointestinal tract, mucocutaneous melanotic spots and increased predisposition to malignancy. The polyp stalk of Peutz-Jeghers syndrome tends to lengthen with the growing of polyps. The bigger polyp with long stalk as the lead point of an intussusception may be outlined distal to the tapered lumen of the intussusceptum. On the other hand, polyps of CCS tend to be relatively small without a long stalk. However, in these patients, the larger size of the cecum polyp might be occur in spite of the thick stalk of the polyp. Not only polypysis but also inflammatory bowel disease could cause giant pseudopolyps. Colonolic intussusception of a giant pseudopolyp was reported in a patient with inflammatory bowel disease (24, 25).

In conclusion, we described an adult with cecal intussusception due to CCS polyps that was relieved by colonoscopy. Colonoscopic reduction may reduce the necessity of an emergent surgery in intussusceptive patients with low malignant potential and no peritoneal irritation sign.

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