

REFERENCES

1. Farrell RJ, Kelly CP. Celiac sprue. N Engl J Med 2002; 346: 180-8.
2. Bardella MT, Trovato C, Quatrini M, et al. Mesenteric lymph node cavitation: a rare hallmark of celiac disease. Scand J Gastroenterol 1999; 34(12): 1257-9.
3. Howat AJ, McPhie JL, Smith DA, et al. Cavitation of mesenteric lymph nodes: a rare complication of celiac disease, associated with a poor outcome. Histopathology 1995; 27: 349-54.
4. Howatt AJ, McPhie JL, Smith DA, et al. Cavitation of mesenteric lymph nodes: a rare complication of coeliac disease, associated with a poor outcome. Histopathology 1995; 27: 349-54.
5. DeBoer WA, Maas M, Tytgat GN. Disappearance of mesenteric lymphadenopathy with gluten-free diet in celiac sprue. J Clin Gastroenterol 1993; 16: 317-9.
6. Tomei E, Diacinti D, Marini M, et al. Abdominal CT findings may suggest coeliac disease. Dig Liver Dis 2005; 37: 402-6. Epub 2005 Mar 17.
7. Tomei E, Semelka RC, Braga L, et al. Adult celiac disease: what is the role of MRI? J Magn Reson Imaging 2006; 24: 625-9.

Müge USTAOĞLU¹, Ahmet BEKTAŞ¹,
Tülay BAKIR¹, Murat DANACI²

Department of ¹Gastroenterology and ²Radiology, Ondokuz
Mayıs University School of Medicine, Samsun

Cecal duplication cyst presenting as perforation in an adult patient

Erişkin bir hastada perforasyon sonucu saptanan çekal duplikasyon kisti

To the Editor,

Alimentary tract duplications are uncommon congenital abnormalities that may occur anywhere in the digestive tract from the lingual root to the anus (1). More than 80% of the cases can be detected prenatally or in the first two years of life, but this rare entity may remain asymptomatic for years, even until adulthood, unless complications occur (2). Many complications related to colonic duplications have been reported in adults, such as obstruction, bleeding, intussusception, or melena (3-6). Peritonitis related to perforation of the duplication is a rare condition in adults (1,3). To our knowledge, this report describes the first case of an unusual cause of acute abdomen in an adult patient related to cecal duplication cyst perforation (CDCP).

A 27-year-old female patient was admitted to our emergency service with a two-day history of right lower quadrant abdominal pain. The vital signs were stable. Upon physical examination, right lower quadrant abdominal tenderness and rebound were detected. No disorder was determined in the

laboratory parameters. Abdominal ultrasonography and computerized tomography revealed a cystic mass measuring 20x15 cm located in the right lower abdominal quadrant with pericecal fluid, which could be compatible with a mesenteric cyst rupture. After obtaining the patient's consent, laparotomy was performed. On exploration, a 20x15 cm perforated cecal duplication cyst was observed (Figure 1). The cyst was totally excised without colonic resection. The patient recovered uneventfully. Histopathological examination revealed a colonic duplication cyst with no evidence of malignancy or heterotopic mucosa.

Approximately 75% of duplications have been reported to be located within the abdominal cavity. The ileum is the most frequently involved, accounting for over 60% of cases, while colonic duplications are comparatively rare, representing only 6.8% of all duplications and often located in the cecum, as in our patient (7).

Symptomatic colonic duplication is a rarity in adults. The clinical picture varies according to the location and size of the lesion, as well as the type

Address for correspondence: Alper SÖZÜTEK

Department of General Surgery, Division of Gastroenterological
Surgery, Mersin University Medical Faculty, Mersin, Turkey
Phone: + 90 324 337 43 00
E-mail: dralpers@hotmail.com

Manuscript received: 27.06.2011 Accepted: 10.10.2011

doi: 10.4318/tjg.2012.0419

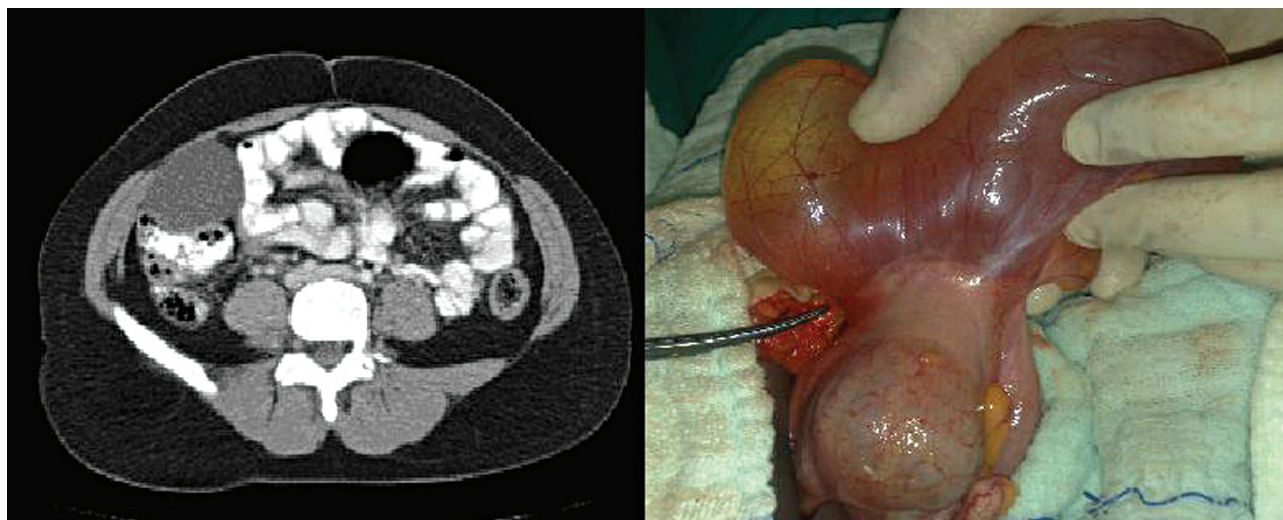


Figure 1. The contrast-enhanced abdominal CT findings of the cystic mass and intraoperative view of the duplication cyst perforation as indicated by the scissor tip.

of mucosal lining (6). The most common symptoms are mild abdominal pain with or without intestinal obstruction signs that may be related to the direct compression of the adjacent bowel or distension of the duplication.

The recommended treatment is surgery either for the treatment of the complications or to avoid further complications, including the possibility of malignant degeneration of the duplication. However, duplications can be considered as benign lesions (7). Hence, the surgical procedure should not be more radical than necessary but should involve

complete resection of the duplication along with the adjacent part of the bowel (6). En bloc resection of the cyst and adjacent viscera is sufficient, as observed in our patient (2,5,6). The prognosis of duplications is good because of the localized and benign character of the lesions. The overall outcome is generally favorable.

In conclusion, CDCP should be considered as an unusual cause of acute abdomen in adult patients. Duplication cysts should be addressed by surgery to avoid further complications.

REFERENCES

1. Sakamoto K, Hasegawa S, Yamazaki Y, et al. Ileal duplication presenting as perforation: report of a case. *Surg Today* 2000; 30: 445-7.
2. Mourra N, Chafai N, Bessoud B, et al. Colorectal duplication in adults: report of seven cases and review of the literature. *J Clin Pathol* 2010; 63: 1080-3.
3. Ryckman FC, Glenn JD, Moazam F. Spontaneous perforation of a colonic duplication. *Dis Colon Rectum* 1983; 26: 287-9.
4. Reiser-Erkan C, Erkan M, Ulbrich E, et al. Cystic colonic duplication causing intussusception in a 25-year-old man: report of a case and review of the literature. *BMC Surg* 2010; 10: 19.
5. Fotiadis C, Genetzakis M, Papandreou I, et al. Colonic duplication in adults: report of two cases presenting with rectal bleeding. *World J Gastroenterol* 2005; 11: 5072-4.
6. Simsek A, Zeybek N, Yagci G, et al. Enteric and rectal duplications and duplication cysts in the adult. *ANZ J Surg* 2005; 75: 174-6.
7. Puligandla PS, Nguyen LT, St-Vil D, et al. Gastrointestinal duplications. *J Pediatr Surg* 2003; 38: 740-4.

Alper SÖZÜTEK, Tahsin ÇOLAK, Ahmet DAĞ,
Ozan KARAK

*Department of Gastroenterological Surgery, Mersin University
School of Medicine, Mersin*