IMAGE | ENDOSCOPY



Inverted Meckel's Diverticulum with Intussusception and Ulceration Diagnosed after Rectal Double-Balloon Enteroscopy

Suliman Al-Shankiti, MD¹, Brendan Halloran, MD¹, Dante D'Urbano, MD², and Sergio Zepeda-Gómez, MD¹

¹Division of Gastroenterology, University of Alberta, Edmonton, Alberta, Canada ²Division of Anatomical Pathology, University of Alberta, Edmonton, Alberta, Canada

CASE REPORT

A 24-year-old man was admitted for abdominal pain and obscure overt gastrointestinal bleeding. The patient had several episodes of hematochezia, and his hemoglobin level dropped to 5.7 g/dL. Initial esophagogastroduodenoscopy and colonoscopy after transfusion of 3 units of blood and fluid resuscitation were negative. The patient reported a lifelong history of intermittent episodes of diffuse abdominal pain, frequently accompanied by nausea and vomiting, which was associated with mild iron-deficiency anemia. Past investigations included upper endoscopy, colonoscopy, magnetic resonance imaging of the abdomen, and a Meckel's scan, all of which were negative.

A computed tomography enterography revealed a tubular lesion at the distal ileum without additional findings. We performed a rectal double-balloon enteroscopy (DBE) to reach Figure 1. Endoscopic view of an inverted diverticulum with large ulcerathis area. At the distal ileum, approximately 50 cm from the ileocecal valve, we found a polypoid lesion of 4-5 cm in length with a broad-based stalk. There was an area of ulceration (about 1 cm in diameter) at the tip of the lesion without active bleeding. A spot tattoo was injected 2 cm proximal and distal to the base of the polypoid lesion (Figure 1 and Video 1). The patient underwent subsequent segmental resection of this



tion located at the distal ileum.

Video 1. Endoscopic view of an inverted diverticulum with large ulceration located at the distal ileum. Watch the video: http://s3.gi.org/media/ links/16-0136Video.mp4.

area through surgery. Histological analysis confirmed the diagnosis of gastric heterotopia with associated ulceration and intussusception (Figure 2). The patient experienced a full recovery and has remained stable, without further episodes of bleeding or abdominal pain at 12 months of follow-up.

Endoscopic resection of a Meckel's diverticulum has been reported previously. One case had a successful outcome after placement of hemoclips at the base of the resection site, and the other case was complicated by perforation.^{1,2} There have been reports in the literature of inverted Meckel's diverticulum diagnosed with balloon-assisted enteroscopy, but only a minority of cases were associated with intussusception.³⁻⁶ The diagnosis of an inverted Meckel's diverticulum can be challenging. High clinical suspicion along with appropriate imaging tests can give useful information. Direct endoscopic evaluation with balloon-assisted enteroscopy can be helpful in confirming the exact nature and location of the lesion.

Correspondence: Sergio Zepeda-Gómez, 1-20 A, Zeidler Ledcor Centre, 130 University Campus, Division of Gastroenterology, University of Alberta, Edmonton, Canada, T6G2X8 (zepedago@ualberta.ca).



🚯 🕲 Copyright: © 2016 Al-Shankiti et al. This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License. To view a copy of this license, visit http://creativecommons.org/licenses/by-nc-nd/4.0.

ACG Case Rep J 2016;3(4):e171. doi:10.14309/crj.2016.144. Published online: December 7, 2016.

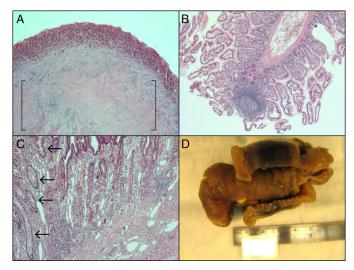


Figure 2. (A) Low-power view of the ulcerated tip with prominent reactive muscularis mucosa hypertrophy (brackets). (B) Normal ileal mucosa adjacent to pyloric metaplasia at the base of the intussusceptum. Notice the lymphoid follicle, frequent in ileal mucosa. Villous architecture is preserved. (C) Pyloric metaplasia is identified throughout the intussusceptum. This corresponds to gastric heterotopia. Notice the normal ileal mucosal crypts on the left (arrows). (D) Intussusception with intussuscipens cut open.

DISCLOSURES

Author contributions: S. Zepeda-Gómez wrote the manuscript and is the article guarantor. S. Al-Shankiti performed literature research. B. Halloran reviewed and edited the manuscript. D. D'Urbano performed histopathological analysis and reviewed the manuscript. Financial disclosure: None to report.

Informed consent was obtained for this case report.

Received February 23, 2016; Accepted May 25, 2016

REFERENCES

- Fukushima M, Suga Y, Kawanami C. Successful endoscopic resection of inverted Meckel's diverticulum by double-balloon enteroscopy. Clin Gastroenterol Hepatol. 2013;11:e35.
- Huang TY, Liu YC, Lee HS, et al. Inverted Meckel's diverticulum mimicking an ulcerated pedunculated polyp: Detection by single balloon enteroscopy. *Endoscopy*. 2011;43:E244-5.
- Fukushima M, Kawanami C, Inoue S, et al. A case series of Meckel's diverticulum: Usefulness of double-balloon enteroscopy for diagnosis. BMC Gastroenterol. 2014;14:155.
- Hotta K, Oyama T, Tomori A, et al. Meckel's diverticulum with ulceration diagnosed by double-balloon enteroscopy. *Digestive Endoscopy*. 2007;19:52-4.
- Takeda H, Sato T, Orii T, et al. Heterotopic gastric mucosa in Meckel's diverticulum incidentally found by double-balloon enteroscopy. *Digestive Endoscopy*. 2008;20:159-61.
- Cakmak GK, Emre AU, Tascilar O, et al. Lipoma within inverted Meckel's diverticulum as a cause of recurrent partial intestinal obstruction and hemorrhage: A case report and review of literature. World J Gastroenterol 2007;13:1141-3.