CASE REPORT

Long-tract ileocolic intussusception due to mucinous adenocarcinoma of the ileocecal valve: a case report and literature review

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Abstract

Intussusception in adults lacks specific symptoms and is often diagnosed emergently when they present with obstruction. Though intussusception certainly varies in size and location, the increased likelihood of ischemia or obstruction make large intussusceptions very rare in the literature. A patient admitted to our facility for small bowel obstruction was found to have extensive intussusception from the right lower quadrant to the splenic flexure, where a lead point was identified. Histopathology revealed multiple satellite lesions surrounding the lead point tumor, which was found to be invasive mucinous adenocarcinoma of the ileocecal valve. While malignancy is found in 60% of lead point identifiable adult intussusceptions, a malignancy is always found in the case of an exceptionally large intussusception.

INTRODUCTION

Intussusception has classically been described as a childhood condition and a rarity in adults [1]. The overall annual incidence of intussusception in adults in the general population has been reported to be as low as 2 cases/1 000 000 persons; adults account for ~5% of confirmed intussusceptions. As high as 90% of cases of adult intussusception are due to a pathological process, with neoplasms accounting for ~60% of identifiable leadpoints [2]. One 6-year prospective study encompassing screening of 380 999 computed tomography (CT) scan reports that were read as intussusception, 0.04% of cases were in adults [3]. In adults, surgery is typically performed owing to the likelihood that a neoplastic or otherwise chronic pathology is the cause of the intussusception [1]. Indications to take any case to the operating room include large caliber of telescoped bowel, long length of telescoped portion, identifiable lead point and evidence of obstruction [2–4]. The former indications owe to the increased likelihood of malignancy requiring surgical exploration and incision, and bowel wall and mesenteric ischemia in the setting of large intussusceptions [4].

CASE REPORT

A 61-year-old gentleman presented with a 3-day history of right-greater-than-left lower quadrant abdominal pain, distention, anorexia and one episode of emesis. The patient also reported only consuming liquids for 5–7 days prior to presentation as solid foods caused significant discomfort. He endorsed intermittent cramping in the past year with a 20-lb unintentional weight loss. Three weeks prior, colonoscopic evaluation of his cramping was aborted at 20 cm due to an impassable sigmoid stricture. Several hyperplastic polyps and one tubular adenoma were removed at that time.

At the time of presentation, he was afebrile and hemodynamically stable. His abdomen was distended and tympanic, with tenderness and fullness appreciated in the right lower
quadrant. Initial laboratory analysis suggested mild dehydration without additional abnormalities. A CT obtained at that time (Fig. 1) demonstrated small bowel obstruction due to an ileocolic intussusception, from the ileocecal valve to the splenic flexure, and collapsed distal colon. The patient was rehydrated and a decision was made to perform exploratory laparotomy.

SURGERY AND PATHOLOGY

The telescoped bowel contained terminal ileum, cecum, appendix and the entire ascending colon (Fig. 2). A 4.5 cm × 5 cm mass was identified at the lead point, the ileocecal valve. Resection included the right hemi-colon, cecum, appendix and 15 cm of terminal ileum. A diverting ileostomy was created as well as a mucus fistula for post-operative antegrade colono-scopic evaluation of the remaining colon.

Additional tubulovillous adenomas, tubular adenomas and hyperplastic polyps were identified throughout the excised portion of bowel. Microsatellite instability (MSI) testing was negative. Final grading and staging of the lead point tumor (Fig. 3) was ‘Invasive Mucinous Adenocarcinoma of the ileocecal valve,’ T2N0M0.

DISCUSSION

This patient’s presentation is exceptionally unique for both the size and cause of intussusception. To our knowledge, very few case reports have ever described such a long-tract intussusception. The few cases we found were always in the setting of malignancy, the patients presented with a similar chronicologic progression with escalation of symptoms including liquid-only diets shortly before presentation. Additionally, the very large intussusceptions were always found in the setting of malignancy [10–12]. It seems that a favorable (mobile) anatomy with an indolent process is a common thread amongst these rare cases.

Further compounding the uniqueness of this case is the lead point tumor’s location and histology, along with the presence of additional terminal ileum and colonic polyps. The annual incidence in the United States of small bowel adenocarcinoma from 1973–2005 is 6.8 cases per 1,000,000 person years, as calculated by Surveillance Epidemiology and Results (SEER) program. Additionally, ileocecal adenocarcinoma of any type is sparsely described outside of a few case reports [8, 9, 13–15]. Mucinous adenocarcinoma tends to be a locally aggressive tumor, and metastasizes by redistribution phenomenon rather than vascular or lymphatic invasion. Pseudomyxoma peritonei, or ‘jelly belly,’ is a rare complication [16].

Presently, we have no unifying diagnosis to correlate the multiple lesions found throughout the telescoped bowel and distal 20 cm of colon and rectum. However, it is currently estimated that up to 40% of patients with extensive polyps and a positive family history are ruled out for HNPCC by genetic testing for mismatch repair and MSI, as was the case for this patient [17]. Nevertheless, the unique presentation of this patient allowed for expeditious removal of a classically aggressive tumor, in what appears to be the first reported case of a long-tract intussusception secondary to a mucinous adenocarcinoma of the ileocecal valve.

CONFLICT OF INTEREST STATEMENT

None declared.

REFERENCES

1. Davis CF, McCabe AJ, Raine PAM. The ins and outs of intussusception: history and management over the past fifty years. J Pediatr Surg 2003;38:60–4.
2. Sarr MG, Nagorney DM, McIlrath DC. Postoperative intussusception in the adult: a previously unrecognized entity? Arch Surg 1981;116:144–8.
3. Rea JD, Lockhart ME, Yarbrough DE, Leeth RR, Bledsoe SE, Clements RH. Approach to management of intussusception in adults: a new paradigm in the computed tomography era. Am Surg 2007;73:1098–1105.
4. Aydin N, Roth A, Misra S. Surgical versus conservative management of adult intussusception: Case series and review. Int J Surg Case Rep 2016 Jan 22.
5. Bosman FT, Carneiro F, Hruban RH, Theise ND. *International Agency for Research on Cancer, WHO Classification of Tumours*. Volume 3, 4th edn, 2010; WHO Regional Office for Europe, Copenhagen, Denmark.

6. Odze RD, Goldblum JR. *Odze and Goldblum Surgical Pathology of the GI Tract, Liver, Biliary Tract and Pancreas*, 3e, 2015. Elsevier Saunders, Philadelphia, PA, USA.

7. Neugut AI, Marvin MR, Chabot JA. Adenocarcinoma of the small bowel. In: Holzheuemer RG, Mannick JA, eds. *Surgical Treatment: Evidence-Based and Problem-Oriented*. Munich: Zuckscherwerdt, 2001. Available from: https://www.ncbi.nlm.nih.gov/books/NBK6880/.

8. Yoruk G, Aksoz K, Buyrac Z, Unsal B, Nazli O, Ekinci N. Adenocarcinoma of the ileocecal valve: report of a case. *Turk J Gastroenterol* 2004;15:268–9.

9. Chand P, Patel AA, Cervellione KL, Sulh M. A rare case of mucinous adenocarcinoma of the colon presenting as ileo-ileal intussusception in an adult. *Case Rep Med* 2012;2012:1–4. doi:10.1155/2012/940947.

10. Kraniotis P, Pastromas G, Tsota I, Patsoura M, Petsas T. Giant ileocolic intussusception in an adult induced by a double ileal lipoma: a case report with pathologic correlation. *Radiol Case Rep* 2016;11:148–51. doi:10.1016/j.radcr.2016.04.014.

11. Yol S, Bostanci EB, Ozogul Y, Akoglu M. Extensive adult colo-colonic intussusception from ascending colon to sigmoid colon: report of a case. *Turkish J Gastroenterol* 2004;15:201–3.

12. Marinis A, Yiallourou A, Samanides L, Dafnios N, Anastasopoulos G, Vassiliou I, et al. Intussusception of the bowel in adults: a review. *World J Gastroenterol* 2009;15:407–11. doi:10.3748/wjg.15.407.

13. Horton KM, Jones B, Bayless TM, Lazenby AJ, Fishman EK. Mucinous adenocarcinoma at the ileocolic valve mimicking crohn’s disease. *Digest Dis Sci Digestive Diseases a Sci* 1994;39:2276–81. doi:10.1007/bf02090384.

14. Winder T, Lenz H-J. Mucinous adenocarcinomas with intra-abdominal dissemination: a review of current therapy. *Oncologist* 2010;15:836–44.

15. Maksimovic S. Survival rates of patients with mucinous adenocarcinoma of the colorectum. *Med Arh* 2007;61:26–9.

16. Landry CS. Analysis of 900 appendiceal carcinoid tumors for a proposed predictive staging system. *Arch Surg Arch Surg Surg* 2008;143:664 doi:10.1001/archsurg.143.7.664.

17. Lindor NM, Rabe K, Petersen GM, Casey G, Baron J, Gallinger S, et al. Lower cancer incidence in Amsterdam-I criteria families without mismatch repair deficiency: familial colorectal cancer type X. *JAMA* 2005;293:1979–85.