Y Shaped Colonic Duplication Mimicking Intestinal Volvulus: A Case Report and Review of Literature

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ABSTRACT

Enteric duplications are rare congenital anomalies found anywhere from mouth to anus. Colonic duplications constitute about 13% of all enteric duplications. In this report a 6-year-old boy with chronic abdominal pain for a duration of last 2 years requiring intermittent hospital admissions was diagnosed as colonic duplication mimicking intestinal volvulus. Clinical findings are nonspecific and definitive diagnosis can only be made during surgical intervention and surgical treatment is advocated for all duplications. The topic is discussed under the light of relevant literature with a brief literature review.

Keywords: Colonic duplication, intestinal volvulus, children

INTRODUCTION

Enteric duplications (“ED”) are rare congenital anomalies in the gastrointestinal tract from mouth to anus (1). Sites of involvement include ileum (33%), esophagus (20%), colon (13%), jejunum (10%), stomach (7%), and duodenum in 5% of cases (2-4). More than 80% of patients present before the age of 2 years and findings in presentations vary from case to case. These include nonspecific findings like; abdominal pain and mass, acute abdomen or intestinal obstruction like volvulus, or intussusception and rectal bleeding (5-7). The aim of this study is to present a case with Y shaped colonic duplication presenting like intestinal volvulus and to discuss the topic with regard to relevant literature and to give a brief literature review. CASE A 3-year-old boy with a complaint of abdominal pain and vomiting was admitted to our clinic. He had a chronic abdominal pain for a duration of last 2 years requiring intermittent hospital admissions. The physical examination revealed that the boy was dehydrated and tachycardic. Resuscitation with IV fluid and electrolyte was commenced promptly. He had a moderate abdominal distention and laboratory tests were within normal range. Standing abdominal X-ray showed large air collection at the middle abdomen with multiple gas filled, grossly dilated bowel loops (Figure 1). Abdominal ultrasonography (US) reported a diffuse collection of fluid in the abdominal cavity at the region of hepatorenal, pelvic, and superior to bladder. Urgent computerized tomography (CT) scan of the abdomen revealed findings compatible with an intestinal volvulus located periumbilically at the right of mid abdominal line (Figure 2). Emergent laparotomy was performed and a tubular bowel seg-
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Figure 1. Standing abdominal X-ray showing large air collection in the abdomen with multiple gas filled, grossly dilated bowel loops.

Figure 2. Abdominal CT scan showing intestinal volvulus in the abdominal cavity.

ment originating from ascending colon, measuring 21.5x5x8 cm with a dilated blind end floating freely in the abdominal cavity together with hemorrhagic fluid collection were found. Duplicated colonic segment was found to be congested and twisted 360° anticlockwise around mesentery with caecum and appendix found in the right lower quadrant (Figure 3). After detorsion of the volvulus, in addition to an incidental appendectomy, resection of colonic duplicated segment was performed with the aid of linear 6 cm stapler device (Figures 4 and 5). A second suture layer with 4/0 polyglicolic acid was performed for reinforcement of stapler suture line at the resection site in the native colon. Histopathologic examination of the excised specimen revealed a colonic duplication containing all layers of large intestine without evidence of ectopic or abnormal tissue. For a possibility of concomitant urinary, cardiac and vertebral anomalies, the patient was evaluated accordingly and had no accompanying anomalies. The child did well post operation and was commenced on oral feeds on the 4th postoperative day and discharged on 7th postoperative day. He is disease free and gaining weight with no symptoms.

DISCUSSION

Alimentary tract duplications in children are rare congenital anomalies commonly seen under the age of 2 years as an acute abdomen or bowel obstruction \(^{(8,9)}\). The incidence of gastrointestinal duplications is 1 in 4500 autopsies \(^{(10)}\). The first report of ED was made by Calder in 1733 and the term “Duplications of the Alimentary Tract” was coined by Ladd in 1937 \(^{(6,11)}\). In a meta analysis comprising 580 cases, it was found that 80% of lesions occurred in the abdomen and 20% in the chest \(^{(12)}\). There are numerous terms for defining these masses including: enterogenous cysts,
giant diverticula, ileal or jejuna duplex, and unusual meckel diverticula (7). The etiology of EDs is still unclear and it is believed that it occurs between 4th and 8th weeks of gestation (13). There are several proposed theories to explain the pathophysiology of EDs suggesting that the origin of ED can be multifactorial. These are the theories of split notochord, luminal recanalisation, partial twinning, persistent embryonic diverticula, and intrauterine vascular accident (2,3,14-16). EDs have 3 characteristics in common; epithelial lining containing alimentary mucosa, smooth muscle envelope, and close attachment with gastrointestinal tract showing common wall (13). Structurally, they can be cystic in 80% and tubular in 20% of cases. Cystic duplications are not related to adjacent intestinal lumen whereas tubular lesions may be related to adjacent colonic lumen adjacent as in our case (17). It has been reported that ectopic tissue is present in 25-30% of duplicated specimens and most common types of ectopic tissues are gastric followed by pancreatic tissue (6). Presentation of colonic duplication is variable and asymptomatic in 10% of patients and can be discovered accidentally at surgery (18). Vague abdominal pain and distention, vomiting, constipation or failure to thrive may be observed. As an emergency setting, the children may present with an acute intestinal obstruction due to intussusception or volvulus as in the presented case. If there is ectopic gastric tissue in the epithelial lining of duplicated colon, rectal bleeding may be observed. Extra gastrointestinal anomalies including genital, urinary or cardiovascular systems have been reported in 80% of patients with colonic duplications (19,20). Our patient did not reveal any finding related to these systems. Imaging findings may be helpful in diagnosing colonic duplications in children. Plain abdominal X-ray is usually nonspecific and shows features of intestinal obstruction and air filled intestinal loops.
Ultrasonography ("US") is the imaging modality of choice in the diagnosis of ED but is operator dependent. Classical findings of uncomplicated cystic EDs are the presence of a cyst adjacent to the gut with double-wall or muscular sign (gut signature sign) but US may be non-helpful in diagnosing tubular duplications [13]. Sonographic finding in the presented case is nonspecific and includes massive abdominal fluid collection in the abdominal cavity. Due to ionizing radiation computerized tomography (CT) is not typically performed to evaluate the EDs but may depict location and extension of duplication and anatomical relationship with surrounding structures as well as complications like volvulus [13]. CT finding in our case was an intestinal volvulus necessitating urgent surgical intervention. The treatment in colonic duplications is surgical excision of the duplicated intestinal segment. The aims of surgery are to relieve the symptoms, to eliminate the risks of complications like volvulus, intussusception or bleeding from an ectopic gastric mucosa. Resection of duplicated colonic segment can also decrease the risk of adenocarcinoma because the occurrence of adenocarcinoma in the duplicated colon is higher than duplications located at any other locations [21,22]. Other surgical treatment options especially in extensive tubular colonic duplications include cyst marsupialisation, partial cystectomy, and mucosal stripping. In conclusion, colonic duplications especially Y shaped lesions in children may be a challenge for clinicians with regard to not the surgical treatment but the clinical diagnosis because these cases usually can not be diagnosed usually without surgical intervention. Significant morbidity and even mortality may be observed if these patients are left untreated. A high index of suspicion is necessary to recognize this anomaly and clinicians should keep this entity in their minds in children with nonspecific complaints of gastrointestinal tract including abdominal pain, vomiting or intestinal obstruction and these children should be provided treatment and care promptly for an uneventful recovery.

Conflict of Interest: None.
Informed Consent: Obtained from the patient’s relatives.

REFERENCES

https://doi.org/10.1097/00007611-193704000-00002
https://doi.org/10.1148/radiographics.13.5.8210590
https://doi.org/10.4067/S0718-40262009000200011
https://doi.org/10.1002/jcu.22007
https://doi.org/10.5334/jbr-btr.92
https://doi.org/10.1177/000313480807400315
https://doi.org/10.1007/s00383-004-1248-x
https://doi.org/10.1016/j.ejrad.2006.03.012
https://doi.org/10.1007/s002619900305
https://doi.org/10.1136/jcp.2010.083238