



Congenital trans-mesenteric internal hernia: a rare cause of acute intestinal obstruction in adult

Nor A. Yasin, MD, Mohamed R. Ahmed, MD, Abdulkadir N. Mohammed, Abdihamid M. Ali

Introduction and Importance: Internal hernia is responsible for 0.6–5.8% of all small intestinal obstructions. Only 8% of internal hernias are of the congenital trans mesenteric variant. Urgent surgical intervention should be considered in individuals who exhibit intestinal obstruction before the development of irreversible bowel ischemia and necrosis.

Case Presentation: The authors report a 38-year-old male patient who presented to the emergency department with abdominal pain, distension, and vomiting for the last three days. After an explorative laparotomy, it was confirmed that there was a trans mesenteric hernia defect with strangulated distal ileal loops. End-to-end ileo-ileal anastomosis was done.

Clinical Discussion: Early recognition and subsequent surgical treatment permit proper management and prevent complications. There should be a differential diagnosis. In this case, there is no prior history of abdominal surgery, and the patient presents with recurrent abdominal pain and intestinal obstruction.

Conclusion: Early diagnosis and emergency laparotomy can save the intestine before gangrene, lowering morbidity and mortality, correcting the mesenteric defect to prevent recurrences, and enhancing clinical outcomes because many studies have shown that some cases are missed before radiological investigation. Laparotomy is still the method of choice for acute cases of incarceration with bowel obstruction, strangulation, and ischemia. The entire mesentery needs to be evaluated, and all mesenteric defects need to be sutured to prevent recurrence.

Keywords: internal hernia, intestinal obstruction, morbidity, mortality

Introduction and importance

Internal hernia is defined as the protrusion of abdominal viscera into a normal or aberrant hole within the abdominal and pelvic cavity, most commonly the small bowel^[1]. Despite having an overall incidence of less than 1%, it is responsible for 0.6–5.8% of all small intestinal obstructions, and it is interesting to note that only 8% of internal hernias are of the congenital transmesenteric variant^[2].

Aberrant intestinal rotation that takes place during embryonic development as well as pre-existing anatomical structures can both result in congenital internal hernias (CIHs). Although there are no distinctive symptoms or signs that clearly separate internal hernia from other types of intestinal obstruction, the presentation might be either acute or chronic^[3]. Since early diagnosis is often

HIGHLIGHTS

- Internal hernia is responsible for 0.6–5.8% of all small intestinal obstructions. Only 8% of internal hernias are of the congenital transmesenteric variant.
- Despite having an overall incidence of less than 1%, it is responsible for 0.6–5.8% of all small intestinal obstructions and It's interesting to note that only 8% of internal hernias are of the congenital transmesenteric variant.
- Congenital internal hernias (CIHs) can be caused by aberrant intestinal rotation that occurs during embryonic development as well as pre-existing anatomical structures.
- Early recognition and subsequent surgical treatment permit proper management and prevent complications.
- There are no distinctive symptoms or signs that clearly separate internal hernia from other types of intestinal obstruction, the presentation might be either acute or chronic.
- Urgent surgical intervention should be considered in individuals who exhibit the key indicators of acute intestinal obstruction, such as high fever, persistent hypotension, and abdominal rigidity.

Department of General Surgery, Mogadishu Somali Turkey Recep Tayyip Erdoğan Training and Research Hospital, Mogadishu, Somalia

Sponsorships or competing interests that may be relevant to content are disclosed at the end of this article.

*Corresponding author. Address: Department of General Surgery, Yaqshiid District, Mogadishu, Somalia. Tel.: +252 615 732 173. E-mail: dalnurshe68@gmail.com (N.A. Yasin).

Copyright © 2024 The Author(s). Published by Wolters Kluwer Health, Inc. This is an open access article distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as the author is credited and the new creations are licensed under the identical terms.

Annals of Medicine & Surgery (2024) 86:6121–6124

Received 9 November 2023; Accepted 22 February 2024

Published online 3 April 2024

<https://dx.doi.org/10.1097/MS9.0000000000001931>

hard and can show up quickly and in an emergency situation, people who have high fevers, persistent low blood pressure, and rigid abdominal muscles should think about having surgery right away. These are the main signs of an acute intestinal obstruction.

This work has been reported in line with the Surgical CAsE REport (SCARE) 2020 criteria^[4].



Figure 1. Erect abdominal radiograph reveals multiple irregular small bowel loops without classic small bowel obstruction.

Case presentation

A 38-year-old man with three days of colicky abdominal pain, distension, and vomiting visited the emergency department. His medical history was unknown, including abdominal surgery and trauma. He was fit and healthy. A previous abdominal pain incident resolved spontaneously. This attack started with umbilical pain and became generalized to the abdomen. His general examination revealed fever, dehydration, normal tension, and tachycardia. Distension, rigidity, abdominal peritonitis, and a slow bowel sound were seen on abdominal inspection. Investigation-wise, leukocytosis (WBC $21 \times 1000 \text{ mm}^3$) and increased CRP (153 mg/l) were present, although LDH was normal (210 u/l). An erect abdomen radiograph showed several air-fluid levels (Fig. 1), and an abdominal ultrasound showed pelvic-free fluid and intestinal dilatation. A decompressive NGT tube was introduced to adjust intravenous fluid and electrolyte levels, and explorative laparotomy was planned for acute intestinal obstruction. Exploration revealed a trans mesenteric hernia defect with strangulated distal ileal loops (Figs 2, 3, 4). Retrieving and untwisting strangled intestinal loops from the hernia sac required expanding the sac edge, and bowel viability evaluation revealed impaired blood supply.

Due to delayed presentation and gangrenous bowel, two-stage segmental ischemic resection, enterostomy, and end-to-end ileo-ileal anastomosis were performed. Uninteresting postoperative. Semisolid food and toleration were introduced on day two of the stoma's postoperative work. The pathological analysis of the resected intestine revealed an ischemic and necrotic state of the small bowel.

Clinical discussion

Internal hernias account for 0.5 to 3% of all cases of intestinal obstruction. A trans mesenteric hernia is a type of internal hernia caused by a congenital mesentery defect. Rokitsky recorded the first trans mesenteric internal hernia in an autopsy in 1836^[4]. About 0.5% of all autopsies have mesenteric defects. Intestinal obstructions related to internal hernias, which are viscus protrusions through intra-abdominal apertures without traversing fascial planes, are rare^[5]. Predominantly preduodenal (53%), trans mesenteric, and trans mesocolic account for 8%. Although rare, trans mesenteric hernias have increased their incidence after surgical procedures like Roux-en-Y bypass and liver transplant surgeries^[6]. 5–10% of internal hernias are CTMH. The mesentery window allows intestinal loops to herniate without a sac in CTMH, causing complications^[7]. Due to their particular pathology, only 1% of the population has them, although they cause roughly 6% of small bowel obstruction^[8]. This case was extremely unusual because of a congenital sac and a trans mesenteric defect that caused the proximal bowel to herniate. The defect was trapped in the congenital sac, which subsequently became strangled within it.

Due to the lack of clinical indicators, preoperative diagnosis is difficult. An inaccurate diagnosis might cause delayed exploration, which could result in irreversible bowel ischemia and a subsequent increase in morbidity and mortality^[9]. An isolated mesenteric defect can be a rare type of intestinal atresia, and in the majority of the reported cases, the diagnosis of an internal hernia was not made prior to surgery in most cases^[10]. Clinical symptoms range from moderate digestive issues to significant abdominal pain, depending on hernia size^[11]. Although similar to acute small bowel obstruction, a palpable abdominal mass with

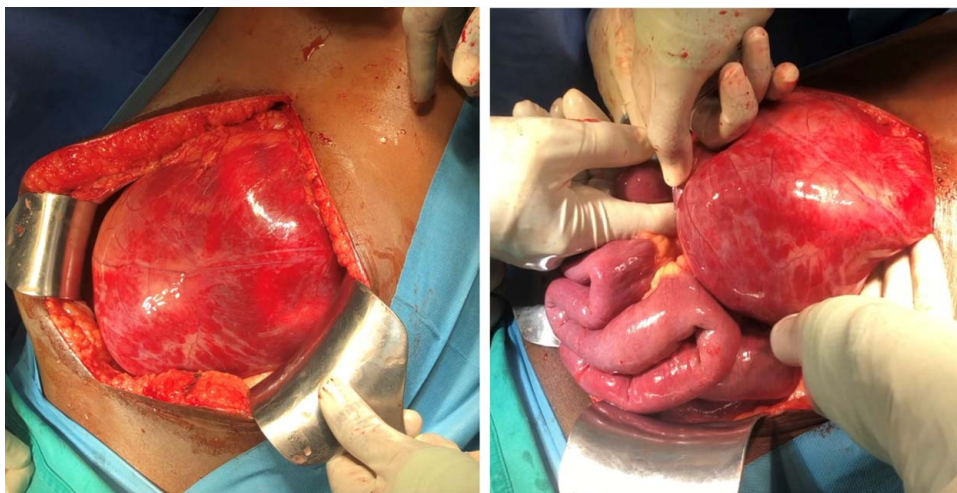


Figure 2. Intraoperative image illustrating a huge congenital internal hernia sac was seen occupying infracolic compartment containing multiple loops of ileum.

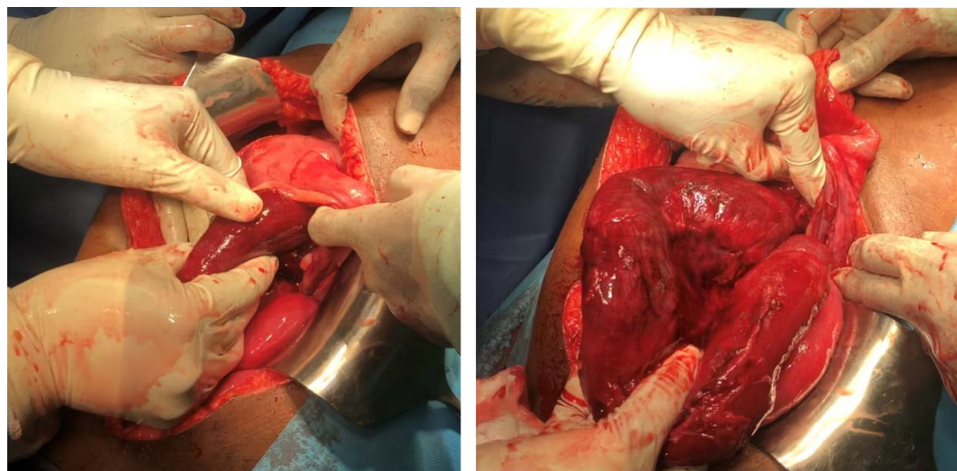


Figure 3. An intraoperative image demonstrated a ischemic small intestinal through mesenteric defect.

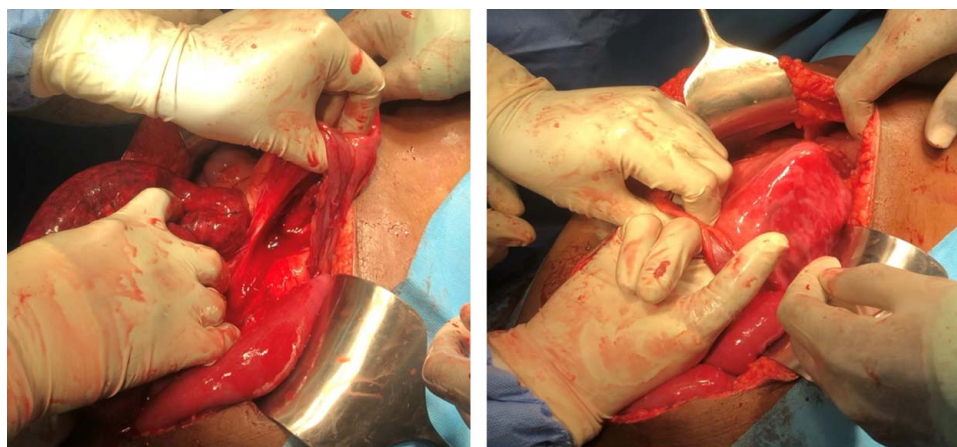


Figure 4. Mesenteric defect through which the hernia occurred. Trans-mesenteric type of congenital internal hernia.

localized discomfort is noted. They include nausea, vomiting, pain in the stomach, distension, and constipation. When a patient has bowel obstructions without having had abdominal surgery, an inflammatory disease, or trauma, a congenital internal hernia should be taken into consideration^[12]. The positive prognostic markers for intestinal ischemia included abdominal rebound pain, leukocytosis ($> 18\,000/\text{mm}^3$), or a high level of manual band form ($> 6\%$); however, a history of chronic intermittent abdominal pain was a negative indicator^[11].

In our case, due to peritonitis, distention, and late presentation over 48 h, we decided on an explorative laparotomy and used clinical and abdominal radiograph and ultrasounds to make our decision.

Treatment of this hernia requires reduction of herniated structures and closure of broad ligament defects to prevent recurrence. Surgical management of congenital internal hernias comprises reducing the hernia's closure, resection of the strangulated ischemic bowel^[11]. Performing a primary anastomosis or an enterostomy depending on local and systemic factors related to patients, and fixing hernia defects with a nonabsorbable suture^[13]. In our cases, we had two problems with congenital

masseteric defects. With the congenital sac, we reduced the bowel and the bowel was ischemic. We did resection of the strangulated part of the bowel without anastomosis, resulting in bowel edema, and we closed the defect with a silk nonabsorbable suture. We also resected the congenital sac that trapped the bowel by ligature with an electric vascular and tissue sealer device. We washed the abdomen and the patient followed aggressive in the postoperative period.

Early identification and exploratory laparotomy are the gold standard to prevent fatal complications in congenital trans mesenteric intra-abdominal hernias because bowel strangulation has a high morbidity and mortality rate. Internal hernias have a mortality rate that reportedly ranges from 20 to 75%^[14].

Conclusion

A strong index of suspicion is needed in all cases of small intestinal obstruction and nonspecific clinical and radiographic characteristics to diagnose internal hernias, a rare but fatal problem. Congenital trans-mesenteric hernias seldom cause small intestine obstruction in adulthood without surgery, trauma, or

peritoneal inflammation. Internal hernia, especially transmesenteric, can cause intestinal obstruction. Early diagnosis and emergency laparotomy can save the intestine before gangrene, lowering morbidity and mortality, correcting the mesenteric defect to prevent recurrences, and enhancing clinical outcomes because many studies have shown that some cases are missed before radiological investigation. Laparotomy is still the method of choice for acute cases of incarceration with bowel obstruction, strangulation, and ischemia. The entire mesentery needs to be evaluated, and all mesenteric defects need to be sutured to prevent recurrence.

Ethics approval and consent to participate

Ethical approval was not needed for writing a case report in our settings.

Consent

Written informed consent was obtained from the patient for publication of this case report and accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal on request.

Consent for publication: for the publication of this case report and any accompanying pictures, the patient provided written informed consent. A copy of the written consent is available for review by the Editor-in-Chief of this journal on request.

Source of funding

No funding for this research.

Author contribution

N.A.Y., A.M.A., A.N.M., and E.M.: wrote the manuscript and corrected the manuscript for its scientific basis; N.A.Y., A.M.A., A.N.M., and E.M.: collected the data for the study. All authors have read and approved the final manuscript.

Conflicts of interest disclosure

The authors declare that they have no conflict of interest in writing this case report.

Research registration unique identifying number (UIN)

Not applicable.

Guarantor

Nor Abdi Yasin.

Availability of data and materials

The author declare that all data in this article are available within the article.

Provenance and peer review

Not commissioned, externally peer-reviewed.

Acknowledgements

Not applicable.

References

- [1] Kelahan L, Menias CO, Chow L. A review of internal hernias related to congenital peritoneal fossae and apertures. *Abdom Radiol* 2021;46: 1825–36.
- [2] Doishita S, Takeshita T, Uchima Y, *et al.* Internal hernias in the era of multidetector CT: correlation of imaging and surgical findings. *Radiographics* 2016;36:88–106.
- [3] Agha RA, Franchi T, Sohrabi C, *et al* for the SCARE Group. The SCARE 2020 guideline: updating consensus Surgical CAse REport (SCARE) Guidelines. *Int J Surg* 2020;84:226–30.
- [4] Noura F, Dhaou B, Charieg A, *et al.* Small bowel obstruction caused by congenital transmesenteric defect. *Afr J Paediatr Surg* 2011;8:75.
- [5] Fan HP, Yang AD, Chang YJ, *et al.* Clinical spectrum of internal hernia: a surgical emergency. *Surg Today* 2008;38:899–904.
- [6] Newsom BD, Kukora JS. Congenital and acquired internal hernias: unusual causes of small bowel obstruction. *Am J Surg* 1986;152:279–85.
- [7] Akyildiz H, Artis T, Sozuer E, *et al.* Internal hernia: complex diagnostic and therapeutic problem. *Int J Surg* 2009;7:334–7.
- [8] Banker A, Martin L, Merkle E, *et al.* Congenital transmesenteric hernia—importance of a timely intervention. *Surg J* 2020;6:e98–100.
- [9] Al Buainain L, Kaundinya KB, Hammed FN. Transomental hernia—an enigmatic case report causing bowel obstruction in a virgin abdomen. *Int J Surg Case Rep* 2020;66:88–90.
- [10] Czeiger D, Vaynshtein J, Kukeev I, *et al.* Emergency management of internal hernia. *Textbook of Emergency General Surgery: Traumatic and Non-traumatic Surgical Emergencies*. Springer International Publishing; 2023:1155–1161.
- [11] Lingegowda AP, Amit B, Pillai PR, *et al.* Congenital transmesenteric hernia—A rare presentation. *J Indian Assoc Pediatr Surg* 2023;28:72.
- [12] Dave A, McMahon J, Zahid A. Congenital peritoneal encapsulation: a review and novel classification system. *World J Gastroenterol* 2019;25: 2294.
- [13] Islam S, Shah AN, Dial S, *et al.* Congenital dual internal hernias causing small bowel obstruction in a man with no prior surgical history: a report of a very rare case. *Am J Case Rep* 2021;22:e932132–1.
- [14] Chowdhury A, Tata RC, Shah A, *et al.* Congenital peritoneal encapsulation: a literature review of a rare pathology. *Cureus* 2022;14:e31765.