

OPEN ACCESS

EDITED BY Zhuang Aobo, Xiamen University, China

REVIEWED BY
Chengfei Xu,
Xiamen University, China
Mahdi Parvizi Mashhadi,
Mashhad University of Medical Sciences, Iran
Gloria Caterine Pérez Mingan,
Universidad de Cartagena, Colombia
Dhouha Bacha,
Université de Tunis El Manar, Tunisia

*CORRESPONDENCE
Haihua Yu
Myhhsd200308@163.com

[†]These authors have contributed equally to this work and share first authorship

RECEIVED 02 September 2025 ACCEPTED 22 September 2025 PUBLISHED 08 October 2025

CITATION

Shu Z, Li Z, Lin S and Yu H (2025) Recurrent intestinal intussusception due to ileal angiolipoma: an uncommon, rare, and potentially life-threatening entity: a case report. *Front. Oncol.* 15:1680934. doi: 10.3389/fonc.2025.1680934

COPYRIGHT

© 2025 Shu, Li, Lin and Yu. This is an openaccess article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

Recurrent intestinal intussusception due to ileal angiolipoma: an uncommon, rare, and potentially life-threatening entity: a case report

Zheng Shu^{1,2†}, Zhe Li^{3†}, Shengnan Lin^{2,4} and Haihua Yu^{1,3*}

¹Department of General Surgery, The First Affiliated Hospital of Shandong First Medical University & Shandong Provincial Qianfoshan Hospital, Jinan, China, ²Shandong First Medical University & Shandong Academy of Medical Sciences, Jinan, China, ³Department of Emergency Medicine, The First Affiliated Hospital of Shandong First Medical University & Shandong Provincial Qianfoshan Hospital, Jinan, China, ⁴Department of Endocrinology and Metabology, The First Affiliated Hospital of Shandong First Medical University & Shandong Provincial Qianfoshan Hospital, Shandong Institute of Nephrology, Jinan, China

Angiolipoma, a rare benign intestinal tumor, is primarily diagnosed through abdominal imaging and pathological examination. Intestinal angiolipomas frequently cause intussusception, necessitating prompt surgical resection. This paper reports a 62-year-old female patient admitted to the First Affiliated Hospital of Shandong First Medical University with "unprovoked paroxysmal abdominal pain for 3 months." Preoperative diagnosis indicated ascending colonic intussusception secondary to ileal angiolipoma. Pathological examination following surgical bowel resection confirmed the lesion as an (ileal) angiolipoma. Given the rarity of this pathological entity in the intestinal tract, we present this case.

KEYWORDS

angiolipoma, intussusception, rare pathological type, cancer, case

Introduction

Angiolipoma is a distinct benign adipose tissue tumor (1), composed of mature adipocytes and abnormally proliferating blood vessels (predominantly small-caliber). It most commonly occurs in subcutaneous tissues (2), particularly the forearms (near the elbow region), trunk (back, shoulders, abdomen), and lower extremities (thighs, calves). Involvement of the breast, paravertebral region, or gastrointestinal tract is exceedingly rare (3–5). Over the past decade, only six articles have reported cases of angiolipomas occurring in the gastrointestinal tract (4, 6–10). The low incidence rate poses challenges for the management of this case. This case report describes a 62-year-old female patient who presented with CT findings suggestive of intussusception involving the ascending colon and ileocecal region. Following endoscopic reduction, the intussusception recurred rapidly, necessitating partial small bowel resection with anastomosis. Serial abdominal computed

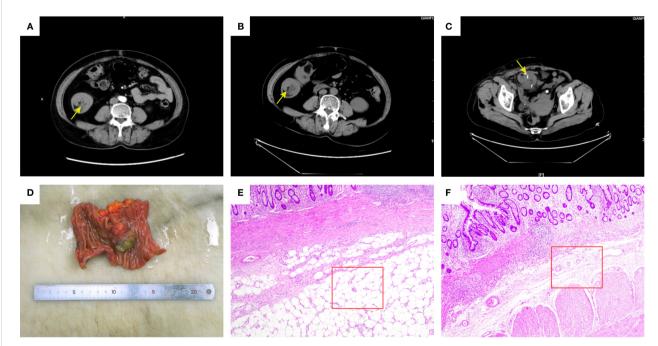
tomography (CT) imaging demonstrated temporal evolution of the lesion. Histopathological examination of the resected bowel specimen confirmed the rare diagnosis of an ileal angiolipoma. This paper will offer new insights into the diagnosis and treatment of ascending colon intussusception caused by ileal hemangioma.

Case report

A 62-year-old woman presented to The First Affiliated Hospital of Shandong First Medical University in February 2025 with a 3month history of paroxysmal abdominal pain, exacerbated over the preceding 4 days. Her medical history included renal cyst unroofing decortication in 2018, with no family history of cancer. Physical examination revealed mild abdominal distension without visible bowel patterns or peristalsis, abdominal wall tension and resistance, tenderness with rebound tenderness; liver and spleen were nonpalpable, Murphy's sign was negative, no costovertebral tenderness was elicited, bowel sounds were slightly hyperactive (6-8/min), and shifting dullness was absent. Pre-admission gastroscopy showed no significant abnormalities. Admission lower abdominal CT demonstrated an irregular structure in the ascending colon with adipose tissue invagination into the lumen, irregular wall thickening, and no enlarged pericolonic lymph nodes (Figure 1A). Following clinical assessment, endoscopic reduction of the intussusception was performed. However, a repeat CT one day post-reduction revealed tortuous bowel loops in the ascending colon and ileocecal region, a double-tube sign in the ascending colon with mesenteric fat herniation, associated wall thickening, and pericolonic fat stranding, suggesting recurrent ascending colonic intussusception (Figure 1B). Given the confirmed diagnosis and prompt recurrence after endoscopic reduction, laparoscopic partial small bowel resection was undertaken. Intraoperatively, after reducing the intussuscepted bowel, a firm mass was identified approximately 15 cm proximal to the ileocecal valve, deemed the cause of the refractory intussusception, and resected for pathology. Postoperative CT showed findings consistent with laparoscopic small bowel resection, including linear high-density opacities and patchy soft tissue densities in the subcutaneous surgical site, with mild wall thickening around the anastomosis (Figure 1C). Pathological examination of the small bowel specimen revealed a submucosal angiolipoma (3.7 x 3.0 x 2.7 cm) with surface ulceration, located in the submucosa compressing the underlying muscularis propria downward, with uninvolved resection margins (Figures 1D-F). Immunohistochemistry was positive for CD34 (vascular component) and negative for HMB-45 and Melan-A.

Discussion

Lipomas represent the most common benign mesenchymal tumors. They are broadly categorized into simple lipomas, composed solely of mature adipocytes, and fibrolipomas, which contain an admixture of



Abdominal CT imaging findings and pathological analysis of resected bowel segments. (A) Pre-endoscopic reduction CT: Axial view showing intussusception of the ascending colon (yellow arrow); (B) Post-endoscopic reduction CT: Recurrent intussusception (yellow arrow) with mesenteric fat incarceration; (C) Post-laparoscopic resection CT: Surgical site showing anastomotic bowel (yellow arrow) with mild wall thickening; (D) Macroscopic specimen shows a well-defined submucosal mass protruding into the lumen with surface ulceration; (E) Microscopic observation (HE stain, 40x): Submucosal angiomyolipoma composed of mature adipose tissue and proliferative vessels (red box); (F) Microscopic observation (HE stain, 40x): Normal submucosal intestinal tissue (red box).

adipose and fibrous tissue (11). Within the intestines, angiolipomas predominantly arise in the submucosa or subserosa, with a higher incidence in the small bowel compared to the colon (12). To evaluate the clinical relevance of lipomas causing intussusception, a literature search was conducted using PubMed (pubmed.ncbi.nlm.nih.gov), the free public database developed by the US National Library of Medicine (NLM). Employing the search strategy ("lipoma" OR "angiolipoma") AND "intussusception" for publications between January 2015 and January 2025, and applying predefined inclusion/ exclusion criteria supplemented by manual review of reference lists, 635 articles related to "lipoma" were identified. Of these, 22 specifically addressed "angiolipoma", and only 6 reported cases occurring in the gastrointestinal tract (4, 6-10). Most gastrointestinal angiolipomas are small (< 2 cm) and asymptomatic. However, when tumors exceed 2 cm in size or undergo hemorrhagic rupture, they can precipitate gastrointestinal bleeding, intestinal obstruction, or intussusception (13). Intussusception occurs precisely because the angiolipomatous mass acts as a "lead point," propelled by peristalsis into the distal bowel lumen (14).

The pathophysiological mechanism by which ileal lipomas cause ascending colon intussusception is now largely understood (15-17). This pathological process begins with the ileal lipoma acting as a local inflammatory stimulus and fixation point, initially triggering asynchronous contraction and hyperperistalsis in the proximal ileum. This abnormal peristalsis, using the tumor as a fulcrum, propels the terminal ileum into the ascending colon. During this process, the physiological anchoring effect of colonic peristalsis in the ascending colon provides essential resistance, directly facilitating the occurrence of ileo-colonic intussusception. The invagination immediately causes mechanical intestinal obstruction, leading to proximal intestinal dilation, fluid and gas accumulation, and a rapid increase in abdominal cavity volume, ultimately resulting in intra-abdominal hypertension. Intraabdominal hypertension is not merely a consequence but actively exacerbates intestinal wall edema and compression of mesenteric vessels. This creates a vicious cycle where the compression of the intussusception sheath mutually intensifies, collectively leading to severe pathological outcomes such as intestinal ischemia and necrosis.

The etiology of intussusception in adults differs fundamentally from that in children. Over 90% of adult cases have an identifiable pathological cause, broadly categorized as neoplastic or nonneoplastic (18). Non-neoplastic causes account for only 30–40% of cases, including anatomic abnormalities (e.g., Meckel's diverticulum, intestinal adhesions) and inflammatory or infectious processes (19–21). In contrast, neoplastic factors constitute 60–70% of etiologies, encompassing both malignant tumors (colorectal carcinoma, lymphoma, or metastatic cancer) and benign neoplasms (lipomas, gastrointestinal stromal tumors, or adenomatous polyps) (21–24). Current management primarily involves surgical intervention. Comprehensive preoperative evaluation typically identifies the underlying cause, including the

precise location of intussusception and presence of a pathological "lead point (25). The initial surgical step involves manual reduction of the intussuscepted bowel segment, followed by assessment for ischemic bowel necrosis and underlying tumor pathology. For benign tumors, surgical resection is generally curative. Reported recurrence rates following resection of benign lead points are less than 5% (12). Gastrointestinal angiolipomas are exceptionally rare. Dr. Juan Sun et al. aggregated all reported cases over the past two decades, identifying only 30 documented instances. Their analysis revealed that these tumors predominantly affect middle-aged and elderly males, with the ileum and colon being the most common sites. Notably, small bowel involvement exceeds colonic occurrence (12). Abdominal pain presents non-specifically; however, angiolipomas in the upper gastrointestinal tract typically manifest with hemorrhage, whereas those in the lower tract primarily cause pain and obstruction. Remarkably, during follow-up extending to 10 years post-resection, no recurrences were documented among the 30 patients. The diagnostic challenge lies in the non-specificity of abdominal symptoms, underscoring the critical role of abdominal CT. In the present case, the patient's longstanding gastric history initially directed evaluation toward upper GI pathology, leading to gastroscopy alone. Crucially, abdominal CT was not performed at initial symptom onset. Consequently, surgical intervention was delayed until bowel necrosis developed. While resection was ultimately performed, earlier imaging could have prevented progression to transmural compromise and mitigated risks of hemorrhagic shock or intra-abdominal infection.

In summary, ileal angiolipoma represents a rare yet treatable etiology of adult intussusception. In the present case, preoperative CT identification of a fat-density mass with vascular enhancement proved crucial for differentiating this benign lesion from malignancies such as gastrointestinal stromal tumors (GISTs) or lymphoma. Complete surgical resection remains the gold standard for preventing recurrence and complications. Clinicians should include angiolipoma in the differential diagnosis of adults presenting with occult GI pain or unexplained intussusception, particularly when imaging demonstrates fat-containing lesions. The accumulation of additional cases remains necessary to elucidate its pathogenesis and optimize management strategies.

Data availability statement

The original contributions presented in the study are included in the article/supplementary material. Further inquiries can be directed to the corresponding author.

Ethics statement

The studies involving humans were approved by Ethics Committee of the First Affiliated Hospital of Shandong First Medical University. The studies were conducted in accordance

with the local legislation and institutional requirements. The human samples used in this study were acquired from primarily isolated as part of your previous study for which ethical approval was obtained. Written informed consent for participation was not required from the participants or the participants' legal guardians/next of kin in accordance with the national legislation and institutional requirements. Written informed consent was obtained from the individual(s) for the publication of any potentially identifiable images or data included in this article. Written informed consent has been obtained from the participant/patient permitting the publication of this case report.

Author contributions

ZS: Writing – original draft, Writing – review & editing. ZL: Formal Analysis, Project administration, Visualization, Writing – original draft. SL: Project administration, Software, Supervision, Writing – review & editing. HY: Methodology, Resources, Supervision, Writing – review & editing.

Funding

The author(s) declare that no financial support was received for the research, and/or publication of this article.

References

- 1. Dupuis H, Lemaitre M, Jannin A, Douillard C, Espiard S, Vantyghem MC. Lipomatoses. *Ann Endocrinol (Paris)*. (2024) 85:231–47. doi: 10.1016/j.ando.2024.05.003
- 2. Kransdorf MJ, Larsen BT, Goulding KA, Cumsky JL, Hwang S, Long JR. Angiolipoma: a review of 778 lesions in 344 patients. *Skeletal Radiol.* (2023) 52:541–52. doi: 10.1007/s00256-022-04075-9
- 3. Noel JC, Van Geertruyden J, Engohan-Aloghe C. Angiolipoma of the breast in a male: a case report and review of the literature. *Int J Surg Pathol.* (2011) 19(6):813–6. doi: 10.1177/1066896909350467
- 4. Vara-Luiz F, Nunes G, Oliveira C, Mendes I, Dahlstedt-Ferreira C, Currais P, et al. Duodenal angiolipoma: a rare tumor causing recurrent upper gastrointestinal bleeding. *Rev Esp Enferm Dig.* (2023) 115(11):674–5. doi: 10.17235/reed.2023.9744/2023
- 5. Iampreechakul P, Tangviriyapaiboon T, Liengudom A, Lertbutsayanukul P, Thammachantha S, Siriwimonmas S. Sacral extradural angiolipoma associated with tight filum terminale and spina bifida coexisting with spinal arteriovenous fistula. *World Neurosurg.* (2020) 140:37–45. doi: 10.1016/j.wneu.2020.04.239
- 6. Ni X, Yin J, Chen L, Zhu Z. Endoscopic resection of a rare duodenal polypoid mass: duodenal angiolipoma. *Rev Esp Enferm Dig.* (2023) 115:332–3. doi: 10.17235/reed 2022 2068/2022
- 7. Lou XH, Chen WG, Ning LG, Chen HT, Xu GQ. Multiple gastric angiolipomas: A case report. World J Clin Cases. (2019) 7:778–84. doi: 10.12998/wjcc.v7.i6.778
- 8. Alghamdi SA, Alshammery SM, Alamari MS, Alshehri MA, Alshahrani AS. An unusual case of gastric angiolipoma presenting with upper gastrointestinal bleeding. Oxf Med Case Rep. (2022) 2022:omac119. doi: 10.1093/omcr/omac119
- 9. Choi JH, Park SB, Lee JB, Lee TJ, Park HJ, Lee ES. A huge gastric angiolipoma presenting with acute upper gastrointestinal hemorrhage: A case report. *Curr Med Imaging*. (2023) 19:945–9. doi: 10.2174/1573405618666220822095201
- 10. Kimura A, Morinaga N, Tajima K, Ogata K, Kato H, Sohda M, et al. Colon angiolipoma with intussusception resected by laparoscopy-assisted surgery: a case report. *J Surg Case Rep.* (2021) 2021(8):rjab329. doi: 10.1093/jscr/rjab329
- 11. Charifa A, Azmat CE, Badri T. Lipoma pathology. In: *StatPearls*. StatPearls Publishing, Treasure Island (FL (2022).

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Generative AI statement

The author(s) declare that no Generative AI was used in the creation of this manuscript.

Any alternative text (alt text) provided alongside figures in this article has been generated by Frontiers with the support of artificial intelligence and reasonable efforts have been made to ensure accuracy, including review by the authors wherever possible. If you identify any issues, please contact us.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

- 12. Sun J, Kang W, Zeng Z, Zhang Z, Wang X. Rare localization of angiolipoma in the gastrointestinal tract: a case series. *J Int Med Res.* (2020) 48:300060520938589. doi: 10.1177/0300060520938589
- 13. Alharbi K, Bilal A. Colonic angiolipoma: A rare cause of chronic anemia and rectal bleeding. *Cureus*. (2024) 16:e56678. doi: 10.7759/cureus.56678
- 14. Nissen A, Yi F. Rare cause of colonic intussusception in an adult. *BMJ Case Rep.* (2017) 2017:bcr2017221976. doi: 10.1136/bcr-2017-221976
- 15. Marinis A, Yiallourou A, Samanides L, Dafnios N, Anastasopoulos G, Theodosopoulos T. Intussusception of the bowel in adults: a review. World J Gastroenterol. (2009) 15(4):407–11. doi: 10.3748/wjg.15.407
- 16. Tarchouli M, Ait Ali A. Adult intussusception: an uncommon condition and challenging management. Visc Med. (2021) 37:120–7. doi: 10.1159/000507380
- 17. Heersche S, Hirt J, Butti F, Hübner M, Hahnloser D, Joliat GR, et al. Intestinal intussusception in adults: A systematic review. $World\ J\ Surg.\ (2025)\ 10.1002/wjs.70055.$ doi: 10.1002/wjs.70055
- 18. Neymark M, Abramov R, Dronov M, Gilshtein H. Management of adult intussusception-A case series experience from a tertiary center. *World J Surg.* (2021) 45:3584–91. doi: 10.1007/s00268-021-06277-z
- 19. Xia F, Sun Z, Wu JH, Zou Y. Intussusception after reconstruction following gastrectomy for gastric cancer. World J Surg Oncol. (2021) 19:340. doi: 10.1186/s12957-021-02456-3
- 20. Huang Q, Peng X, Hu S, Tan X, Li X. Diabetic ketoacidosis complicated with intussusception: a case report. *World J Emerg Med.* (2024) 15:419–21. doi: 10.5847/wjem.j.1920-8642.2024.084
- 21. Chong EH, Kim DJ, Kim S, Kim G, Kim WR. Inverted Meckel's diverticulum: Two case reports and a review of the literature. World J Gastrointest Surg. (2018) 10:70-4. doi: 10.4240/wjgs.v10.i6.70
- 22. Yang Y, Jia D, Jiang C. Multiple intestinal hemangioma concurrent with low-grade appendiceal mucinous neoplasm presenting as intussusception-a case report and literature review. *World J Surg Oncol.* (2022) 20:44. doi: 10.1186/s12957-022-02519-z
- 23. Zhu LB, Li PF, Xiao WH, Zhang PB, Li JQ, Sun MF. A distal ileum Malignant peripheral nerve sheath tumor causing intussusception in a patient in China: a case report. *World J Surg Oncol.* (2017) 15:29. doi: 10.1186/s12957-017-1101-z

24. Panzera F, Di Venere B, Rizzi M, Biscaglia A, Praticò CA, Nasti G, et al. Bowel intussusception in adult: Prevalence, diagnostic tools and therapy. *World J Methodol.* (2021) 11(3):81–7. doi: 10.5662/wjm.v11.i3.81

25. Patsikas MN, Papazoglou LG, Paraskevas GK. Current views in the diagnosis and treatment of intestinal intussusception. *Top Companion Anim Med.* (2019) 37:100360. doi: 10.1016/j.tcam.2019.100360