Check for updates

OPEN ACCESS

EDITED BY Anna Maria Musolino, Bambino Gesù Children's Hospital (IRCCS), Italy

REVIEWED BY Rahul Gupta, Synergy Institute of Medical Sciences, India José Estevão-Costa, University of Porto, Portugal

*CORRESPONDENCE Lihao Tang 🖾 nedved1987@163.com

[†]These authors have contributed equally to this work

RECEIVED 08 May 2024 ACCEPTED 12 August 2024 PUBLISHED 16 September 2024

CITATION

Cui Q, Xu X, Li C and Tang L (2024) Wandering spleen combined with pedicle torsion and splenic infarction: a rare case report and literature review. Front. Pediatr. 12:1429490.

doi: 10.3389/fped.2024.1429490

COPYRIGHT

© 2024 Cui, Xu, Li and Tang. 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.

Wandering spleen combined with pedicle torsion and splenic infarction: a rare case report and literature review

Qian Cui¹, Xiaoyan Xu^{2†}, Chang Li^{3†} and Lihao Tang^{2*}

¹Department of Radiology, Shaoxing Central Hospital, Shaoxing, China, ²Imaging Center, Tengzhou Central People's Hospital, Tengzhou, China, ³Department of Pathology, Tengzhou Central People's Hospital, Tengzhou, China

Wandering spleen (WS) concurrent with splenic pedicle torsion and infarction has been described rarely. We reported our experience in diagnosing and treating such a condition in a 16-year-old girl with acute abdominal pain. A plain CT scan showed the wandering of the spleen from the left upper quadrant. Contrast-enhanced CT indicated dilatation and distortion in the splenic vein, a counterclockwise "whirl sign" in the splenic pedicle, pancreatic tail torsion, and splenic infarction. The patient was diagnosed with WS combined with splenic pedicle torsion and splenic infarction and underwent splenectomy for treatment. She showed a satisfactory outcome during the follow-up. To enhance our understanding of it, we performed a comprehensive literature research to summarize the clinical manifestations, treatment options, and outcomes among adolescent patients.

KEYWORDS

wandering spleen, torsion, infarction, contrast-enhanced computed tomography, acute

1 Introduction

Wandering spleen (WS), also known as ectopic spleen, is a rare clinical condition in which the spleen migrates to an abnormal position within the abdomen or pelvis (1). It mainly affects children aged between 3 months and 10 years old and women of childbearing age (2). A major complication of WS is the torsion of the splenic pedicle, which can subsequently cause splenic infarction and rupture (3). In clinical practice, WS combined with acute splenic torsion is considered a life-threatening emergency (4). However, the diagnosis of this condition remains challenging due to its asymptomatic progression.

Currently, the diagnosis of WS mainly depends on clinical and imaging features, especially contrast-enhanced CT. This imaging technique can reveal the location, size, vascular condition of the spleen, and presence of splenic pedicle torsion. To date, less attention has been paid to the WS and the splenic torsion due to the rarity of cases. In this study, we present a case of a 16-year-old girl who presented with acute abdominal pain and was eventually diagnosed with WS, pedicle torsion, and splenic infarction.

2 Case presentation

A 16-year-old girl presented to our hospital's emergency department with progressive aggravation of abdominal pain for at least 24 h. Physical examination revealed a migrating

mass with tenderness in the left lower quadrant. A plain CT scan showed the spleen absent from the left upper quadrant (Figure 1, Plate 1A). In other areas of the left abdomen, a soft tissue density shadow in an elliptic shape was observed. The visualization of hyperdense splenic vessels suggested thrombosis of the splenic vein (Figure 1, Plate 1B). A contrast-enhanced CT scan showed a counterclockwise "whirl sign" in the splenic pedicle, along with pancreatic tail torsion and splenic infarction (Figure 1, Plate 1C). Furthermore, the visualized splenic veins showed dilatation and distortion (Figure 1, Plate 1D). Vascular reconstruction showed an elongated splenic pedicle vessel (Figure 1, Plate 2). The CT scan revealed an enlarged spleen with no enhancement in the arterial and venous phase, suggesting splenic ischemia and infarction (Figure 1, Plate 3).



FIGURE 1

Plate 1, contrast-enhanced computed tomography (CECT) images. (A) Absence of the spleen in the left upper abdomen after plain CT scan. (B) CT scan indicated an elliptic soft tissue density shadow (yellow arrow) in the left upper quadrant. The hyperdense shadows in splenic veins suggested splenic vein thrombosis (red arrow). (C) The axial section showed a "whirl sign" (red arrow) of the splenic pedicle vessels counterclockwise, together with pancreatic tail torsion (green arrow). The white arrow indicated the absence of obvious enhancement in the spleen, showing a possibility of splenic infarction. (D) The visualized splenic veins showed dilatation and tortuosity (yellow arrow). Plate 2, computed tomography (CT) image showing elongated splenic pedicle vessels (yellow arrow). Plate 3, coronal and sagittal CT images. (A) Contrast-enhanced CT scan indicated splenomegaly. There was pelvic fluid induced by splenic infarction. (B) There was a filling of contrast agent in the upper segment of the vessels (yellow arrow), while there was no filling in the lower segment (red arrow). This indicated a block of the splenic artery.

The patient was finally diagnosed with WS combined with splenic pedicle torsion and splenic infarction. Laparoscopy was then performed in a supine position after anesthesia. A 10 mm trocar was inserted through an arc-shaped incision below the umbilicus. Carbon dioxide was injected to establish pneumoperitoneum, with pressure set to 14 mmHg. The laparoscope showed a black and infarcted spleen without ligamentous fixation that was merely suspended by the vascular pedicles in the left lower abdomen. In addition, the splenic pedicle was twisted 720° counterclockwise, along with pancreatic tail torsion.

For the treatment, splenectomy (Figure 2) was performed since the splenic ischemia showed no improvement even after splenic repositioning. Three trocars (i.e., 10 mm, 5 mm, and 5 mm) were placed in the right lower abdomen, right umbilicus, and left upper abdomen, respectively. The spleen was repositioned to the splenic fossa, where it showed no adhesion to the surrounding structure and omentum. The splenic pedicle vessels were doubleligated at the pancreatic tail, and the splenic artery and vein branches were separated and cut using Hem-o-lock clips. The spleen was placed in a retrieval bag and was cut into pieces, and then the pieces were removed through the umbilical incision. The abdominal cavity was flushed, and the splenic blood vessels were ligated. A drainage tube was placed in the left upper abdomen and exited through the left trocar hole. The surgical time was 200 min, and the intraoperative bleeding volume was ~30 ml. Postoperative histopathological analysis confirmed ischemic necrosis. During the 10-month follow-up, the patient showed normal conditions with no recurrence of abdominal pain.

3 Discussion

WS with torsion of the splenic pedicle is an extremely rare condition with an incidence rate of <0.2% (5). There is no sex difference under the age of 10 years; however, after the first

decade, females have a higher prevalence of WS than that in males, with a ratio of 7:1 (6). Nevertheless, the diagnosis of WS combined with splenic pedicle torsion in adolescents remains challenging due to the rarity of cases. In this study, we report a 16-year-old girl with WS accompanied by splenic pedicle torsion and splenic infarction. Meanwhile, a comprehensive literature review was undertaken to fully enhance our understanding of this condition.

Anatomically, the spleen is fixed by six primary suspensory ligaments (i.e., gastrosplenic, splenorenal, splenophrenic, splenocolic, splenopancreatic, and presplenic fold ligaments) and two ligaments (pancreaticocolic and phrenicocolic ligaments) in an indirect association. The gastrosplenic, splenorenal, and phrenicocolic ligaments are usually loose or absent in the etiology of WS (2). Congenitally, WS is caused by the absence or deficits of ligaments resulting from the failure of the dorsal mesentery to fuse with the posterior abdominal wall during embryogenesis, which can be accompanied by developmental abnormalities such as congenital diaphragmatic hernia or intestinal malrotation (7). Additionally, the laxity of the supporting ligaments has been identified as an acquired cause of WS. This condition was mainly presented in women of childbearing age, secondary to multiple pregnancies and hormonal changes (8). Other acquired causes included malaria-, infectious mononucleosis- or lymphoma-induced chronic splenomegaly, abdominal trauma, and surgery (9).

Congenital WS is more common in adolescents than in adults. Cases are generally asymptomatic or show mild symptoms in childhood as the WS does not twist in the early years. The risk of WS increases with age and exercise. In adulthood, the prevalence of WS increases, with many patients developing acquired WS. In this study, we present a case of a 16-year-old girl with acute abdominal pain due to WS and splenic pedicle torsion. Therefore, we focused on the literature search on this condition in adolescents rather than adults. Herein, we identified 32 (10–41) previous case reports and case series published from



Representative intraoperative images. (A) The twisted splenic pedicle. (B) The splenic tissues of the spleen.

2003 to 2023 involving 37 WS cases (female, 25; male, 12) aged 12– 18 years, as presented in Table 1. There was no difference in the sex ratio under the age of 10 years, but females showed a higher prevalence of WS than that in males (6).

The diagnosis of WS is usually a challenge in clinical practice. Its symptoms vary according to the size and location of the spleen. For example, in our literature review, some cases were completely asymptomatic in the early stage (11). Along with disease progression, migration of the spleen predisposes to the torsion of the lengthy and loosening of the splenic pedicle. In the presence of mild torsion, the thin-walled splenic vein and its tributaries were first affected, resulting in congestive splenomegaly, abdominal discomfort or intermittent abdominal pain (13), and enlarged palpable mass with tenderness. Regional portal hypertension and gastric varices bleeding may occur in a longterm incomplete torsion (22). Moreover, the chronic splenomegaly can induce complete torsion. On this basis, the splenic artery blood flow will be blocked, and the spleen will be infarcted. The clinical symptoms usually include acute abdominal pain, hypotension, and even shock, accompanied by peritoneal irritation (31). The additional signs include fever, nausea, vomiting, loss of appetite, and a palpable mass in the mid-lower abdomen or pelvic cavity. Our case showed acute abdominal pain when presenting to our department. Indeed, attention should be paid to the differential diagnosis of WS from other diseases with acute abdominal pain as the initial symptom.

Laboratory tests for WS patients are mostly normal and nonspecific, and occasionally, some patients may present with leukocytosis (10, 17, 28, 29, 33), thrombocytopenia (14, 17, 24, 31, 41), and neutrophilia (18, 29). Additionally, two patients (i.e., Patients 13 and 17) (32, 38) exhibited mild pancytopenia. One patient (i.e., Patient 22) (36) showed a twofold increase in serum amylase and lipase counts. Another patient (i.e., Patient 31) (33) showed an increased C-reactive protein level. Our case showed normal laboratory indicators, which implied that these laboratory indices may somehow contribute to the diagnosis of WS, but with low efficiency.

WS is usually confirmed by imaging methods such as plain x-ray (42), Doppler ultrasonography (USG) (43), CT (44), and magnetic resonance imaging (MRI) (39). In addition to these methods, one study (17) applied contrast-enhanced ultrasound (CEUS) for the diagnosis of WS. The images were characterized by the absence of the spleen in the left upper quadrant and the presence of a soft tissue mass elsewhere in the abdomen or pelvis. Nevertheless, CT is still the modality of choice for the final diagnosis of WS (45) as it shows multiple advantages such as high sensitivity (46) for the identification of splenic pedicle torsion. In our literature review, 30/37 (81%) adolescents were diagnosed with WS by CT. For the incomplete torsion, CT showed congestive enlargement of the spleen, together with dilated and twisted splenic veins. For complete torsion, splenic artery blood flow blockage was observed after the CT scan, and some or all non-enhancement areas appeared in the spleen. This indicated partial or total splenic infarction. Splenic pedicle torsion is characterized by splenomegaly and abnormal orientation of the splenic hilum, and a "whirl sign" can be used to predict parenchymal organ torsion (44). The hyperdense splenic pedicle is indicative of the thrombosed splenic vein (47). Capsule sign and peritoneal effusion can indirectly reflect splenic infarction. The VR reconstruction would visualize the position of the torsion and the elongated splenic pedicle. Our case presented with WS, combined with complete torsion of the splenic pedicle and splenic infarction. Contrast-enhanced CT showed the absence of mass enhancement in the left lower quadrant, and it was necessary to differentiate mesenteric cysts, lymphangiocysts, and adnexal tumors. Additionally, pancreatic volvulus was also reported in Patients 21 (26), 22 (36), 27 (19), and 32 (23). There may be rare symptoms such as pancreatic tail necrosis and pancreatitis under the simultaneous torsion of the WS and the pancreatic tail. Our case showed torsion of the pancreatic tail. The splenic artery was a terminal artery without communication and was also the terminal arterial circulation. Therefore, the incidence of splenic infarction was higher than that of other organs. Theoretically, any disease that can form an embolism can lead to splenic infarction. In addition to splenic pedicle torsion caused by WS, the other common causes of splenic infarction in children included infection, blood system diseases, and autoimmune diseases, such as infective endocarditis, malaria, sickle cell anemia, infectious mononucleosis, chronic myeloid leukemia, lymphoma, antiphospholipid syndrome, and arteritis (48). In the future, attention should be paid to the exclusion of in situ splenic infarction before considering the possibility of WS with splenic pedicle torsion.

Splenopexy and splenectomy have been used for the treatment of WS. In our literature review, 27 (73%) patients underwent splenectomy. Among these patients, 22 (81%) cases presented with different degrees of splenic torsion (Table 2). Splenopexy is the preferred treatment option for WS when there is no splenic torsion or when there is normal circulation in the distorted splenic vessels or normal blood supply after repositioning the splenic torsion. It is worth noting that a long-term follow-up is necessary given the likelihood of postoperative recurrence of splenic torsion. Splenectomy is required for treating WS accompanied by complications such as splenomegaly, splenic infarction, rupture, and thrombosis (6, 49). However, there is a high possibility of septicemia and severe infection after splenectomy. In our literature review, Patient 1 (10) received polyvalent pneumococcal vaccine and penicillin after splenectomy. Patient 12 (31) received conjugate vaccines against pneumococcus and Haemophilus influenza type B before laparoscopic splenectomy, as well as postoperative penicillin V. Patient 14 (38) received aspirin for about 2.5 months after splenectomy. Three patients (i.e., Patients 25, 26, and 39) (28, 35, 39) received triple vaccination against pneumococcus, meningococcus, and Haemophilus influenza after splenectomy, while one patient (i.e., Patient 32) (23) received the same vaccine before splenectomy. Patient 30 (24) received lifelong amoxicillin. Therefore, it is recommended to use antibiotics and capsular bacterial vaccines (e.g., Haemophilus influenzae, meningococcal, and pneumococcal vaccines).

TABLE 1 Thirty-seven adolescent patients with wandering spleen, from 2003 to 2023.

First author, year (Ref.)	Case No.	Gender/ age (year)	Past medical history	Clinical and imaging findings	Diagnostic imaging approaches	Wandering position	Pedicle torsion	Treatment	Outcome
Alimoglu (10)	01	M/17	Hydrocephalus Ventriculoperitoneal shunt Chronic constipation Mental retardation	Abdominal pain and distention Low-grade fever Leukocytosis of 14 000/mm ³ Massive colonic gas Intestinal adhesions Dilated small intestine Dolichocolon	Abdominal x-ray US	Right lower abdomen	Yes, 360°	Splenectomy Postoperative polyvalent pneumococcal vaccine and penicillin	Sigmoid colon resection was performed 3 months after splenectomy Uneventful recovery
Tucker (14)	02	F/17	4-year history of abdominal pain	Recurrent episodes of cramping pain in various regions of the abdomen Splenomegaly, hypersplenism with thrombocytopenia	СТ	Mid-abdomen	Yes	Splenectomy	Uneventful recovery
Kleiner (12)	03	F/12	Mental retardation Myotonic dystrophy	Non-specific abdominal pain Normal size spleen Unremarkable palpation	US Radionuclide scan Laparoscopy CT	Left lower abdomen	Yes, clockwise 180°	Laparoscopic splenopexy	Uneventful recovery
Falchetti (11)	04	F/14	No	Mobile, slightly tender mass in the left lower abdomen Non-enhanced splenic cyst	US CT CECT	Lower pole of the left kidney	No	Laparoscopic splenopexy	Uneventful recovery
Upadhyaya (15)	05	F/16	No, but ride dirt bikes extensively	Symptoms of dysmenorrhea Large, mobile, well- vascularized mass 22 cm spleen extending from the level of the mid-pole of the left kidney to the pelvis, with a cyst measuring 4.6×2 cm in its inferior pole	US CT	Pelvis	No	Laparoscopic splenopexy Cystectomy	Uneventful recovery
Soleimani (13)	06	M/18	No	Intermittent abdominal pain + mass	US CT	Not available	No	Splenectomy	Uneventful recovery
	07	M/12	No	Tender abdominal mass	US	Not available	No	Splenectomy	Uneventful recovery
	08	F/14	No	Intermittent abdominal pain + mass	US	Not available	No	Splenectomy	Uneventful recovery
	09	F/12	No	Abdominal pain + mass	US CT	Not available	No	Splenopexy with absorbable mesh	Uneventful recovery
Feroci (25)	10	M/15	No	Diffuse abdominal tenderness and guarding Large hypogastric abdominal mass White cell count of $19.4 \times 10^{9}/$ L, hemoglobin 11.0 g/dl, and platelets $410 \times 10^{9}/$ L Splenic infarction and vein thrombosis	US CECT	Left lumbar and iliac region	Yes, 720°	Splenectomy	Uneventful recovery

10.3389/fped.2024.1429490

(Continued)

First author, year (Ref.)	Case No.	Gender/ age (year)	Past medical history	Clinical and imaging findings	Diagnostic imaging approaches	Wandering position	Pedicle torsion	Treatment	Outcome
Cho (21)	11	F/14	Vague recurrent periumbilical pain	Homogenous hypoechoic mobile mass Stomach distended into the left upper abdomen	US CT CECT	Left lower abdomen	No	Splenopexy	Uneventful recovery
Lien (31)	12	M/14	No	Acute abdominal pain, intensified by lying down, and relieved by standing up or walking around Depressed platelet count of $106 \times 10^3/\mu l$ Severe gastric spasm Splenic infarction and vein thrombosis	US Gastroendoscopy CECT Multi-detector row CT (MDCT) angiography	Anterior aspect of the middle to lower abdomen	Yes, 540°	Laparoscopic splenectomy Preoperative conjugate vaccines for pneumococcus and <i>H. influenza</i> type B Postoperative penicillin V	Uneventful recovery
Zarroug (38)	13	F/16	Prematurity (30 weeks)	Vague abdominal pain Malaise and vomiting for several weeks Mobile lower abdominal mass Mild pancytopenia	US CT	Pelvis	Yes, greater than 360°	Laparoscopic splenectomy	Uneventful recovery
	14	F/12	No	Non-tender abdomen with a 10 cm fixed mass in the infraumbilical region	US MRI	Pelvis	Yes	Splenectomy Postoperative aspirin for ~2.5 months	Uneventful recovery
Yılmaz (37)	15	F/15	No	Sharp and persistent pain in abdomen Irritability and loss of appetite Enlarged spleen with partial infarction	US CT	Midline just below the umbilicus	Yes, counterclockwise 540°	Splenectomy	Uneventful recovery
Holst (27)	16	F/16	No	Acute right lower quadrant abdominal pain No appreciable mass and peritoneal signs	CT Laparoscopy	Right lower pelvis	Yes, 720°	Laparoscopic splenectomy	Uneventful recovery
Lombardi (32)	17	M/12	No	Tense abdomen with large palpable mass in pelvis Reduced hematocrit values Enlarged spleen with a large lesion arising from its inferior pole Separated bowel loops	Radiographs Doppler US CECT	Mesogastrium, left iliac fossa	Yes, 360°	Splenopexy	Not available
	18	M/15	No	Asymptomatic Large congested spleen with a sign of venous thrombosis	X-ray Color Doppler US CECT	From the medial margin of the left kidney to the pelvis	Yes	Splenectomy	Not available
Katsura (30)	19	F/13	No	Slightly tender, mobile mass in the left lower abdomen	СТ	Left lower abdomen	Yes, counterclockwise 360°	Laparoscopic splenectomy Splenic autotransplantation Pneumococcal vaccination	Uneventful recovery

(Continued)

10.3389/fped.2024.1429490

First author, year (Ref.)	Case No.	Gender/ age (vear)	Past medical history	Clinical and imaging findings	Diagnostic imaging approaches	Wandering position	Pedicle torsion	Treatment	Outcome
Rellum (34)	20	F/14	No	3 weeks of recurrent abdominal pain, malaise, and fatigue without fever Palpable spleen in the normal left upper abdomen	Color Doppler US MRI	Pelvis	Yes, clockwise 360°	Laparoscopic splenopexy	Uneventful recovery
Gorsi (26)	21	F/16	No	Severe abdominal pain Nausea and recurrent episodes of vomiting Distended fluid-filled stomach Splenic infarction, gastric volvulus (mesenteroaxial), and pancreatic volvulus	US CECT	Right side below the liver	Yes	Splenectomy Gastropexy Sham gastrojejunostomy	Uneventful recovery
Torri (36)	22	F/13	No	7 days of pain in the left abdomen and hypochondrium Fever and nausea Large palpable mesogastric mass, without tenderness and guarding A twofold increase of seric amylase and lipase count Splenomegaly and pelvic effusion Distal pancreatic torsion	US CECT	Mesogastrium	Yes	Laparoscopic splenopexy	Uneventful recovery
Jiang (29)	23	F/17	No	Intermittent lower abdominal pain Neutrophilia and leukocytosis Global splenic ischemia Cystic formations at hilar level	CECT	Pelvis	Yes	Splenectomy	Uneventful recovery
Aguirre Pascual (17)	24	M/18	No	Left flank bulge, nausea, and slight left lower quadrant pain Decreased platelet count Increased white blood cell count	Doppler US CEUS CT MRI	Left lower abdomen	Yes	Laparoscopic splenectomy	Uneventful recovery
Hui Lian (28)	25	M/16	No	Lower abdominal pain Palpable mass over the suprapubic region Mildly tender with dullness on percussion Leukocytosis	US	Pelvis	Yes, clockwise 720°	Splenectomy Triple vaccination against pneumococcus, meningococcus, and <i>Haemophilus influenza</i> 2 weeks after surgery	Uneventful recovery
Rizzuto (35)	26	F/17	Minor beta thalassemia	Leucopenia The triad of classic physical examination findings Coexistence of wandering spleen ad accessory spleen	US CECT	Right lumbar region to pelvis	No	Laparoscopic splenectomy Postoperative low molecular weight heparin (LMWH) for 30 days and triple vaccination against pneumococcus, meningococcus, and <i>Haemophilus influenza</i>	Uneventful recovery

(Continued)

10.3389/fped.2024.1429490

First author, year (Ref.)	Case No.	Gender/ age (year)	Past medical history	Clinical and imaging findings	Diagnostic imaging approaches	Wandering position	Pedicle torsion	Treatment	Outcome
Asafu Adjaye Frimpong (19)	27	F/14	Milder episodes of abdominal pains since childhood	5-day history of nausea, vomiting, loss of appetite, fever, headache, constipation, and central abdominal pain Tenderness around umbilicus Hypokalemia of 3.2 mmol/L and significant ketonuria Organoaxial gastric volvulus, with volvulus of the pancreas, wandering spleen, cholestasis Acute small bowel obstruction secondary to an ileo-ileal intussusception after the first laparotomy	CECT	Anteromedial of the abdomen	No	Laparotomy (twice)	Not available
Assaf (20)	28	F/13	No	Generalized, acute abdominal pain Fever, anorexia, and vomiting Spleen was infarcted without any ligamentous attachments	US CT	Hypogastric region	Yes, 540°	Splenectomy	Uneventful recovery
Abaszadeh (16)	29	M/15	No	Abdominal pain, right lower quadrant pain, pain under his bladder, pelvic pain, rebound tenderness, fever, nausea, vomiting, and lack of appetite Cystic lesion Appendix in the upper right quadrant	СТ	Pelvis	Yes	Splenectomy Postoperative vaccination	Uneventful recovery
Dangen (24)	30	F/17	No	Firm abdominal mass Thrombocytopaenia Very large congenital diaphragmatic hernia Hypoplastic left lung and left pulmonary Left pneumothorax	US CT X-ray	Mid-abdomen	Yes	Splenectomy Postoperative vaccination and lifelong amoxicillin	Uneventful recovery
Perez-Rosillo (33)	31	F/18	Left diaphragmatic hernia repair two years earlier	High levels of C-reactive protein and leukocytosis Low-enhancing, comma- shaped mass in the hypogastric region	US CT	Hypogastric region	Yes	Splenectomy	Uneventful recovery

80

(Continued)

First author, year (Ref.)	Case No.	Gender/ age (vear)	Past medical history	Clinical and imaging findings	Diagnostic imaging approaches	Wandering position	Pedicle torsion	Treatment	Outcome
Colombo (23)	32	F/18	Wandering spleen	Frequent episodes of lower abdominal pain Multi-infarcted pelvic splenomegaly Normocromic normocytic anemia and piastrinopenia <i>Helicobacter pylori</i> gastritis Partially twisted pancreatic tail	CT Laparoscopy	Pelvis	Yes	Laparoscopic splenectomy Triple vaccination against <i>Haemophilus influenzae</i> , pneumococcus, and meningococcus 3 weeks before surgery	Uneventful recovery
Chue (22)	33	F/18	No	2-month history of intermittent left abdominal pain Palpable splenomegaly Gastric varices with features of portal hypertension	CT Oesophagogastroduodenoscopy	Pelvis	No	Laparoscopic splenectomy	Uneventful recovery
Ahmed (18)	34	M/12	Appetite loss, constipation, and mucoid diarrhea	Crampy lower abdominal pain Tachycardia (pulse rate of 134 beats per min) and respiratory rate of 22 breaths per minute Firm, tender mass across the left lower quadrant White blood cell count 10,300/ micl with a neutrophilia of 78.5%, with moderate anemia (Hg 8.6 g/dl, Hct 28%), MCV 86 fl and platelet 775 × 103/ micl Secondarily inflamed appendix	US	Lower abdomen	Yes, 720°	Splenectomy Appendectomy Vaccination against pneumococcus and meningococcus Antibiotic prophylaxis	Uneventful recovery
Bairwa (39)	35	F/14	No	Abdominal mass for ~7 months Mild tachypneic Acute abdomen, abdominal distension, tenderness, rebound tenderness, diffuse guarding, and rigidity Increased platelet count Splenic ischemia and infarction	US CECT	Mid-abdomen	Yes, 720°	Splenectomy Triple vaccination against <i>Haemophilus influenzae</i> , pneumococcus, and meningococcus	Uneventful recovery
Petroianu (41)	36	M/14	Long-term history of frequent episodes of progressively worsening lower abdominal pain associated with dysuria, tenesmus, and constipation	Visible tender pelvic mass A reduced platelet count of $114 \times 103/\mu$ l, a white blood cell count of $6.2 \times 103/\mu$ l, and a hemoglobin level of 11.2 g/dl	CT CECT	Pelvis	Yes	Splenopexy	Uneventful recovery
Lugo-Fagundo, 2022 (40)	37	F/16	No	Severe periumbilical abdominal pain Wandering spleen with torsion and infarction	CT Laparoscopy	Right lower abdomen	Yes, 720°	Splenectomy	Uneventful recovery

CT, computed tomography; CECT, contrast-enhanced CT; US, ultrasound; CEUS, contrast-enhanced ultrasound; MRI, magnetic resonance imaging.

TABLE 2 Summary of surgical procedures for wandering spleen with and without torsion in the literature review.

Surgery	WS without torsion (<i>n</i> = 10)	WS with torsion (<i>n</i> = 27)	Total (n = 37)
Splenectomy	5 (50.0%)	22 (81.5%)	27 (73.0%)
Splenopexy	5 (50.0%)	5 (18.5%)	10 (27.0%)

4 Conclusion

We reported a rare case of WS combined with splenic pedicle torsion and infarction with acute abdominal pain as the initial symptom. WS with splenic pedicle torsion should be considered in the differential diagnosis in children who present with recurrent abdominal pain and a palpable abdominal mass. CECT is the first choice for early diagnosis of WS, which may improve the possibility of spleen preservation and prevent the occurrence of complications.

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 the Ethical Committee of Shaoxing Central Hospital (No. 2023-018). The studies were conducted in accordance with the local legislation and institutional requirements. Written informed consent for participation in this study was provided by the participants' legal guardians/next of kin. Written informed consent was obtained from the participant/patient(s) for the publication of this case report.

References

1. Reisner DC, Burgan CM. Wandering spleen: an overview. Curr Probl Diagn Radiol. (2018) 47(1):68-70. doi: 10.1067/j.cpradiol.2017.02.007

2. Puranik AK, Mehra R, Chauhan S, Pandey R. Wandering spleen: a surgical enigma. *Gastroenterol Rep (Oxf)*. (2017) 5(3):241-3. doi: 10.1093/gastro/gov034

3. Ely AB, Zissin R, Copel L, Vasserman M, Hertz M, Gottlieb P, et al. The wandering spleen: CT findings and possible pitfalls in diagnosis. *Clin Radiol.* (2006) 61(11):954–8. doi: 10.1016/j.crad.2006.06.007

4. Viana C, Cristino H, Veiga C, Leão P. Splenic torsion, a challenging diagnosis: case report and review of literature. *Int J Surg Case Rep.* (2018) 44:212-6. doi: 10. 1016/j.ijscr.2018.02.032

5. Memari M, Nikzad M, Nikzad H, Taherian A. Wandering spleen in an adult man associated with the horseshoe kidney. *Arch Trauma Res.* (2013) 2(3):129–32. doi: 10. 5812/atr.9332

6. Nastiti NA, Niam MS, Khoo PJ. Emergency laparoscopic splenectomy for torsion of wandering spleen in a geriatric patient: a case report. *Int J Surg Case Rep.* (2019) 61:91–5. doi: 10.1016/j.ijscr.2019.07.021

7. Brown CV, Virgilio GR, Vazquez WD. Wandering spleen and its complications in children: a case series and review of the literature. *J Pediatr Surg.* (2003) 38 (11):1676–9. doi: 10.1016/s0022-3468(03)00582-7

8. Romero JR, Barksdale EM Jr. Wandering spleen: a rare cause of abdominal pain. *Pediatr Emerg Care.* (2003) 19(6):412–4. doi: 10.1097/01.pec.0000101582.65509.cc

Author contributions

QC: Data curation, Formal Analysis, Investigation, Resources, Software, Writing – original draft. XX: Data curation, Formal Analysis, Visualization, Writing – original draft. CL: Data curation, Formal Analysis, Methodology, Writing – original draft. LT: Conceptualization, Project administration, Supervision, Writing – review & editing.

Funding

The authors declare that no financial support was received for the research, authorship, and/or publication of this article.

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.

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.

9. Mehta A, Vana PG, Glynn L. Splenic torsion after congenital diaphragmatic hernia repair: case report and review of the literature. *J Pediatr Surg.* (2013) 48(3): e29–31. doi: 10.1016/j.jpedsurg.2013.01.003

10. Alimoglu O, Sahin M, Akdag M. Torsion of a wandering spleen presenting with acute abdomen: a case report. *Acta Chir Belg.* (2004) 104(2):221–3. doi: 10.1080/00015458.2004.11679541

11. Falchetti D, Torri F, Dughi S, Porto C, Manciana A, Boroni G, et al. Splenic cyst in a wandering spleen: laparoscopic treatment with preservation of splenic function. *J Pediatr Surg.* (2007) 42(8):1457–9. doi: 10.1016/j.jpedsurg.2007.03.063

12. Kleiner O, Newman N, Cohen Z. Pediatric wandering spleen successfully treated by laparoscopic splenopexy. *J Laparoendosc Adv Surg Tech A*. (2006) 16(3):328–30. doi: 10.1089/lap.2006.16.328

13. Soleimani M, Mehrabi A, Kashfi A, Fonouni H, Büchler MW, Kraus TW. Surgical treatment of patients with wandering spleen: report of six cases with a review of the literature. *Surg Today.* (2007) 37(3):261–9. doi: 10.1007/s00595-006-3389-0

14. Tucker ON, Smith J, Fenlon HM, McEntee GP. Recurrent torsion of a wandering spleen. Am J Surg. (2004) 188(1):96–7. doi: 10.1016/j.amjsurg.2003.11.042

15. Upadhyaya P, St Peter SD, Holcomb GW 3rd. Laparoscopic splenopexy and cystectomy for an enlarged wandering spleen and splenic cyst. *J Pediatr Surg.* (2007) 42(5):E23–7. doi: 10.1016/j.jpedsurg.2007.03.001

16. Abaszadeh F, Taebi M, Nikzad Jamnani H. Torsion of wandering spleen attached to the omentum: a rare case report from Iran. *Int J Gen Med.* (2020) 13:333–6. doi: 10.2147/ijgm.S248259

17. Pascual EA, Fontanilla T, Pérez Í, Muñoz B, Carmona MS, Minaya J. Wandering spleen torsion-use of contrast-enhanced ultrasound. *BJR Case Rep.* (2017) 3 (1):20150342. doi: 10.1259/bjrcr.20150342

18. Ahmed M, Nasir M, Negash A, Haile K. Wandering spleen with splenic torsion: unusual cause of acute abdomen. *Int Med Case Rep J.* (2022) 15:625–30. doi: 10.2147/ imcrj.S388271

19. Frimpong GAA, Aboagye E, Ayisi-Boateng NK, Antwi K, Bawuah KA, Coleman NE, et al. Concurrent occurrence of a wandering spleen, organoaxial gastric volvulus, pancreatic volvulus, and cholestasis - a rare cause of an acute abdomen. *Radiol Case Rep.* (2019) 14(8):946–51. doi: 10.1016/j.radcr.2019.05.018

20. Assaf R, Shebli B, Alzahran A, Rahmeh AR, Mansour A, Hamza R, et al. Acute abdomen due to an infarction of wandering spleen: case report. J Surg Case Rep. (2020) 2020(2):rjz378. doi: 10.1093/jscr/rjz378

21. Cho SA, Choh NA, Dar I, Jehangir M, Yousuf R. Wandering spleen presenting as recurrent abdominal pain in a young female. *Indian J Pediatr.* (2008) 75 (11):1181–2. doi: 10.1007/s12098-008-0181-8

22. Chue KM, Tan JKH, Pang NQ, Kow AWC. Laparoscopic splenectomy for a wandering spleen with resultant splenomegaly and gastric varices. *ANZ J Surg.* (2020) 90(10):2124–5. doi: 10.1111/ans.15737

23. Colombo F, D'Amore P, Crespi M, Sampietro G, Foschi D. Torsion of wandering spleen involving the pancreatic tail. *Ann Med Surg (Lond)*. (2020) 50:10–3. doi: 10. 1016/j.amsu.2019.12.001

24. Dangen J, Lau S, Abbas S. Treatment of a congenital diaphragmatic hernia with associated wandering spleen: case report of a 17-year-old girl. *Int J Surg Case Rep.* (2020) 77:32–5. doi: 10.1016/j.ijscr.2020.10.049

25. Feroci F, Miranda E, Moraldi L, Moretti R. The torsion of a wandering pelvic spleen: a case report. *Cases J.* (2008) 1(1):149. doi: 10.1186/1757-1626-1-149

26. Gorsi U, Bhatia A, Gupta R, Bharathi S, Khandelwal N. Pancreatic Volvulus with wandering spleen and gastric Volvulus: an unusual triad for acute abdomen in a surgical emergency. *Saudi J Gastroenterol.* (2014) 20(3):195–8. doi: 10.4103/1319-3767.133026

27. Holst JM. Acute abdominal pain in the pediatric emergency department: a case of right lower quadrant wandering spleen that acutely torsed. *J Emerg Med.* (2013) 44 (6):e395–6. doi: 10.1016/j.jemermed.2012.11.063

28. Lian HH, Hayati F, Ali AA, Azizan N, Ani MFC, Suhaili MA, et al. Wandering spleen: a unique cause of acute abdomen. *Folia Morphol (Warsz)*. (2018) 77(2):400–2. doi: 10.5603/FM.a2017.0097

29. Jiang M, Chen P, Ruan X, Ye X, Huang Q. Acute torsion of wandering spleen in a 17-year-old girl. *Int J Clin Exp Med.* (2015) 8(7):11621–3.

30. Katsura S, Kawamura D, Harada E, Enoki T, Hamano K. Single-incision laparoscopic splenectomy and splenic autotransplantation for an enlarged wandering spleen with torsion. *Eur J Pediatr Surg Rep.* (2014) 2(1):23–5. doi: 10.1055/s-0033-1357262

31. Lien CH, Lee HC, Yeung CY, Chan WT, Wang NL. Acute torsion of wandering spleen: report of one case. *Pediatr Neonatol.* (2009) 50(4):177–80. doi: 10.1016/s1875-9572(09)60059-0

32. Lombardi R, Menchini L, Corneli T, Magistrelli A, Accinni A, Monti L, et al. Wandering spleen in children: a report of 3 cases and a brief literature review

underlining the importance of diagnostic imaging. Pediatr Radiol. (2014) 44 (3):279-88. doi: 10.1007/s00247-013-2851-6

33. Perez-Rosillo MA, Gomez-Huertas M, Salmeron-Ruiz A, Lainez-Ramos-Bossini AJ. Acute abdomen secondary to torsion and infarction of a wandering spleen. *Gastroenterol Hepatol.* (2021) 44(8):585–6. doi: 10.1016/j.gastrohep.2020.05.013

34. Rellum R, Risseeuw G, Blaauw I, Lequin M. Splenorenal collaterals as hallmark for a twisted wandering spleen in a 14-year-old girl with abdominal pain: a case report. *Eur J Pediatr Surg Rep.* (2014) 2(1):26–8. doi: 10.1055/s-0034-1370776

35. Rizzuto A, Di Saverio S. Laparoscopic splenectomy for a simultaneous wandering spleen along with an ectopic accessory spleen. Case report and review of the literature. *Int J Surg Case Rep.* (2018) 43:36–40. doi: 10.1016/j.ijscr.2018.01.017

36. Torri F, Parolini F, Vanzetti E, Milianti S, Cheli M, Alberti D. Urgent laparoscopic mesh splenopexy for torsion of wandering spleen and distal pancreas: a case report. *Asian J Endosc Surg.* (2015) 8(3):350–3. doi: 10.1111/ases.12188

37. Yılmaz Ö, Bayrak V, Daştan E, Kotan Ç. Torsion of wandering spleen as a rare reason for acute abdomen: a presentation of two cases. *Ulus Cerrahi Derg.* (2013) 29 (4):200–2. doi: 10.5152/ucd.2013.47

38. Zarroug AE, Hashim Y, El-Youssef M, Zeidan MM, Moir CR. Wandering spleen as a cause of mesenteric and portal varices: a new etiology? *J Pediatr Surg.* (2013) 48 (3):e1–4. doi: 10.1016/j.jpedsurg.2012.12.042

39. Bairwa BL, Gupta S, Singh AK, Gupta P. Wandering spleen with torsion: a rare cause of acute abdomen in a 14-year-old girl. *Arch Clin Cases.* (2022) 9(2):56–61. doi: 10.22551/2022.35.0902.10204

40. Lugo-Fagundo E, Fishman EK. Wandering spleen with torsion and infarction: a case report. Radiol Case Rep. (2022) 17(9):3377-9. doi: 10.1016/j.radcr.2022.06.073

41. Petroianu A, Sabino KR. Wandering splenomegaly reduction after splenopexy. *Int J Surg Case Rep.* (2021) 85:106273. doi: 10.1016/j.ijscr.2021.106273

42. Gordon DH, Burrell MI, Levin DC, Mueller CF, Becker JA. Wandering spleen-the radiological and clinical Spectrum. *Radiology*. (1977) 125(1):39–46. doi: 10.1148/125.1.39

43. Nemcek AA Jr, Miller FH, Fitzgerald SW. Acute torsion of a wandering spleen: diagnosis by CT and duplex Doppler and color flow sonography. *AJR Am J Roentgenol.* (1991) 157(2):307–9. doi: 10.2214/ajr.157.2.1853811

44. Blázquez MJP, Vargas DR, Ferrer MG, González JT, Serrano BV. Torsion of wandering spleen: radiological findings. *Emerg Radiol.* (2020) 27(5):555–60. doi: 10. 1007/s10140-020-01786-1

45. Bough GM, Gargan KE, Cleeve SJ, Farrell S. Diagnosis, management and outcome of splenic torsion; a systematic review of published studies. *Surgeon*. (2022) 20(5):e296–305. doi: 10.1016/j.surge.2021.08.006

46. Nel D, Kloppers C, Panieri E. Laparoscopic splenopexy for wandering spleen with absorbable mesh fixation. *S Afr J Surg.* (2021) 59(1):30a-b. doi: 10.17159/2078-5151/2021/v59n1a353

47. Chu J, Li Z, Luo B, Yang J. Wandering spleen with torsion and complete infarction. *Acta Radiol.* (2011) 52(8):911–3. doi: 10.1258/ar.2011.100504

48. Estevão-Costa J, Correia-Pinto J, Fragoso A, Soares-Oliveira M, Carvalho JL. Neonatal splenic necrosis not related to wandering spleen. *Eur J Pediatr Surg.* (2003) 13(5):344–6. doi: 10.1055/s-2003-43579

49. Lourdusamy V, Patel D, Docobo R, Tantry S, Lourdusamy D, Farukh S. The importance of recognizing wandering spleen as a cause of recurrent acute pancreatitis. *Case Rep Gastrointest Med.* (2018) 2018:7573835. doi: 10.1155/2018/7573835