Wandering spleen is a rare entity with a constant danger of splenic torsion leading to splenomegaly and infarction. It occurs consequent to an embryonal disturbance in the development of the ligaments connecting the spleen with the surrounding tissues. It may be associated with other congenital or hereditary disorders. There is always a constant threat of torsion, infarction, rupture, infection or trauma. The diagnosis of this condition is very difficult due to the lack of specific clinical manifestations, and a high index of suspicion is required. Ultrasonography (US), computerized tomography (CT), magnetic resonance angiography, arteriography, scintigraphy and Doppler scanning are very useful tools for confirming the diagnosis.

**Case Report**

**Patient one.** A 28-year-old woman presented with recurrent attacks of lower abdominal pain for the last 3-4 months which on presentation were severe. The pain was stabbing in nature and was not associated with vomiting, fever or urogenital symptoms. On examination there was a tender, freely mobile, mid-abdominal mass measuring 8 x 10 cm. Her complete blood count was normal. Abdominal ultrasound revealed an empty splenic area and a homogenous, hypoechoic mass in the center of the abdomen highly suggestive of an ectopic spleen. A Doppler sonogram showed no blood flow in the hilum of the spleen consistent with torsion. At laparotomy, the mass was found to be a spleen located in the center of the abdomen and displacing the bowels. It had a long thrombosed pedicle which was pulling the tail of the pancreas inferiorly. Splenectomy was carried out. The spleen was found viable. The histopathological examination confirmed a normal spleen with thrombosed vascular pedicle. The patient had an uneventful postoperative recovery.

**Patient 2.** A 40-year-old woman presented with severe pain in the left hypochondrium of few days duration. There was no fever, vomiting or urogenital symptoms. On examination there was tenderness in the left hypochondrium and left renal angle. Abdominal ultrasound showed an empty splenic region and a 12 x 6 cm homogenous mass was replacing the absent left kidney. At laparotomy, the left kidney was absent and the mass was proved to be a spleen with thrombosed pedicle. Splenectomy was carried out. Histopathology revealed multiple infarcted areas. Post-operative period was uneventful.

**Patient 3.** A 65-year-old female patient was complaining of recurrent lower abdominal pain of 2 months duration. The pain was intermittent, stabbing
and localized. There were no associated symptoms, history of abdominal trauma or urinary complaints. On examination, there was a pelvi-abdominal mass (10 x 15 cm). Her complete blood count, urea, creatinine, and electrolytes and blood clotting profile were within normal limits. Abdominal ultrasound revealed the spleen was not in its normal site. In the centre of the lower abdomen there is a splenic-shaped structure measuring 7 x 13 cm, showing multiple large irregular echolucent areas representing necrosis. The picture is highly suggestive of ectopic spleen with multiple infarctions. Color Doppler ultrasound showed no flow in the splenic artery or vein. The findings are those of wandering spleen with areas of infarctions. Liver scan (Tc-99m Tin colloid) showed homogenous uptake in the liver. There was no uptake in the spleen. Computerized tomography angiogram of the abdomen and pelvis demonstrated absence of the spleen from its normal location. There was a large pelvi-abdominal mass (9 x 12 x 14 cm). The mass was predominantly of fluid density and showed a thick enhancing capsule. At the left superior aspect of the mass there was a whorled appearance containing the splenic vessels, pancreatic tail and some peri-pancreatic fat (Figure 1). The mass was indenting the urinary bladder and loops of small bowel were adherent to it. This mass represents a wandering spleen with chronic torsion and infarction (Figure 2). Laparotomy was carried out. There was a large spleen in the lower abdomen surrounded by peri-splenic hematoma with thrombosed splenic vessels. The splenic pedicle was long and pulled down together with the pancreas. Loops of small bowel were adherent to the mass. Splenectomy was carried out. Histopathology revealed an infarcted spleen with peri-splenic hematoma. The patient had uneventful recovery and was discharged on the 9th post-operative day.

Discussion. Wandering spleen remains an elusive clinical diagnosis. It is a distinctive rare clinical entity. It is characterized by a normal spleen with extreme mobility associated with an elongated pedicle. The condition results from congenital maldevelopment or acquired laxity of the spleen’s suspensory ligaments. The ectopic spleen is usually located in the pelvis, the pelvi-abdominal region or rarely in the retroperitoneum. It usually occurs at 20-40 years of age and most cases are seen in women. The clinical presentations include acute abdomen, chronic abdominal pain, thrombocytopenia or it may be asymptomatic. The acute and chronic abdominal pain are related to torsion, infarction or gangrene. The torsion is predisposed by the elongated splenic pedicle. The most common symptom is an acute surgical abdomen related to acute torsion of the ectopic spleen. The wandering spleen may be associated with Beckwith-Wiedemann syndrome, osteopetrosis, Gaucher disease, gastric volvulus, congenital diaphragmatic hernia, diaphragmatic eventration or epidermoid cyst. The major concern in ectopic spleen is the development of complications. Many case reports have documented the occurrence of various complications in the wandering spleen. These include trauma, torsion, infarction, infection, subcapsular hematoma, obstructive uropathy, gastric outlet obstruction, duodenal obstruction, intestinal obstruction, herniation through a defect in the transverse colon or left-sided portal hypertension. The pre-operative diagnosis of this rare condition is often difficult. Various diagnostic modalities are available to identify the spleen. These include US, CT, magnetic resonance imaging (MRI), scintigraphy, arteriography and Doppler ultrasound. Ultrasound, CT, MRI, arteriography and scintigraphy are extremely valuable. Ultrasound plays a special role and is the elective diagnostic method.
homogenous hypoechoic mass and empty splenic area, while CT shows a homogenous, unenhanced mass.\textsuperscript{5,6} The Doppler ultrasound has proven to be a useful tool indicating the presence or absence of blood flow in the splenic pedicle.\textsuperscript{5,6}

The treatment of the wandering spleen is controversial.\textsuperscript{14} There are 2 lines in the treatment of this condition, conservative or surgical. A conservative approach with Doppler ultrasound follow-up can be offered to patients without symptoms.\textsuperscript{22,40} Surgical intervention is indicated for symptomatic patients and those who develop complications. Splenectomy should be considered for complicated cases only such as infarction, gangrene, subsacular hematoma or absent splenic blood flow.\textsuperscript{12,18,41} Every effort should be practiced to preserve the spleen and splenopexy is the treatment of choice in symptomatic patients and in children.\textsuperscript{14,17,22,25,31} However, some surgeons may perform it for non-complaining patients.\textsuperscript{11} Laparoscopic splenopexy is a state of the art. It is safe, feasible, minimally invasive, without impairing splenic function and is applicable even for adult splenomegaly.

In conclusion, an ectopic spleen is a rare clinical entity and torsion remains an important differential diagnosis in patients presenting with acute abdomen. Early diagnosis is necessary to reduce the risk of complications. Awareness of the condition and the judicious use of appropriate imaging modalities can lead to the correct diagnosis. Splenopexy is the treatment of choice. However, in complicated cases splenectomy is a must.

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References


