

When the unexpected happens: Isolated fallopian tube torsion with large hematosalpinx mistaken for ovarian cyst torsion in a pediatric patient: A case report

Naina Kumar¹, Annapurna Srirambhatla², Shailaja Prabhala³, Mishu Mangla⁴

From ¹Additional Professor, ⁴Associate Professor, Department of Obstetrics and Gynecology, ²Additional Professor, Department of Radiology, ³Additional Professor and Head, Department of Pathology, All India Institute of Medical Sciences, Bibinagar, Hyderabad, Telangana, India

ABSTRACT

Isolated fallopian tube torsion (IFTT) in adolescent girls is an infrequent but significant cause of acute abdominal pain in gynecology, often misdiagnosed as ovarian torsion. A 13-year-old unmarried virgin girl presented to the gynecological outpatient department with acute abdominal pain lasting 5 days, accompanied by nausea, vomiting, and fever for 2 days. Pelvic ultrasound revealed a normally sized uterus and a normal right ovary, while the left ovary was slightly enlarged (4×5 cm) and displaced medially, demonstrating a “whirlpool sign” indicative of vascular pedicle twisting, suggestive of left ovarian torsion. A large exophytic lesion measuring 8×6 cm was also noted in the left adnexa, raising concerns for a left exophytic or paraovarian cyst. Given her clinical presentation, the patient underwent an emergency laparotomy. Intraoperative findings revealed minimal hemoperitoneum (50–80 mL), a normally sized uterus deviated to the left, and a left fallopian tube tightly twisted 3 times, with large hematosalpinx (8×6 cm), distended with clots. Due to complete damage to the left fallopian tube, left salpingectomy was performed, with histopathological examination confirming hematosalpinx. Hence although IFTT is uncommon, it should be included in the differential diagnosis for acute abdominal pain in young girls and women of reproductive age. Early recognition and prompt surgical intervention are crucial to prevent complications and preserve tubal function.

Key words: Acute abdomen, Hematosalpinx, Isolated fallopian tube torsion, Ovary, Salpingectomy


Isolated fallopian tube torsion (IFTT) in adolescent girls is an exceptionally rare cause of acute abdominal pain, often misdiagnosed as ovarian torsion [1]. IFTT was first identified by Bland-Sutton in 1890 and is recognized as a rare yet significant cause of acute abdominal pain in gynecology [1,2]. To date, there are very few reported cases of hematosalpinx resulting from fallopian tube torsion in the absence of outflow tract obstruction [2,3]. Hematosalpinx, characterized by the accumulation of blood within the fallopian tube, is typically associated with conditions such as ectopic pregnancy, pelvic inflammatory disease, tubal cancer, pelvic trauma, and endometriosis. In addition, non-tubal obstructive factors, such as tumors, adhesions, an imperforate hymen, and uterine abnormalities can contribute to its development [3,4]. The presence of hematosalpinx alongside tubal torsion in a young girl adds a layer of complexity to diagnosis and management. With an average incidence of isolated

tubal torsion estimated at 1 in 1,500,000 women, this condition remains exceedingly rare [3,4]. Accurate diagnosis frequently occurs only during exploratory surgery, and delayed intervention can lead to serious complications such as necrosis, irreversible damage to the fallopian tube, and infection [1,5].

IFTT is a rare gynecological emergency with significant fertility implications. Its non-specific symptoms and rarity often lead to misdiagnosis and delayed treatment. This case report highlights a pediatric patient with IFTT and a large hematosalpinx initially mistaken for ovarian cyst torsion, emphasizing the need for increased clinical awareness to ensure timely and accurate diagnosis.

CASE REPORT

The present case report was conducted after informed written consent from the patient’s mother (as the patient was a minor). A 13-year-old unmarried virgin girl presented to the Gynecological Outpatient Department with acute abdominal pain

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Correspondence to: Naina Kumar, Department of Obstetrics and Gynecology, All India Institute of Medical Sciences, Bibinagar - 508126, Hyderabad Metropolitan Region, Telangana, India. E-mail: drmainakumar@gmail.com

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lasting 5 days, accompanied by nausea and vomiting for 3 days. She also reported experiencing a fever over the past 2 days. The patient attained menarche at age 12 and has had regular menstrual cycles since (4–5 days every 28–30 days), with her past menstrual period occurring on September 17th. She denied any history of dysmenorrhea, irregular cycles, or heavy bleeding.

On examination, the patient was alert and oriented to time, place, and person. She had a thin build, with a temperature of 99.8°F, blood pressure of 120/70 mmHg, and a pulse rate of 110 beats/min. Abdominal examination revealed tenderness, rigidity, and guarding, with a large tender mass palpable in the left lower abdomen, comparable in size to a 12–14 weeks gestational uterus. Local examination showed healthy labia majora and minora, with no vaginal bleeding. A per rectal examination revealed a large, tender cystic lesion in the left adnexa with restricted mobility, though its exact size was difficult to ascertain due to pain.

An urgent transabdominal ultrasound demonstrated a normally sized uterus, deviated to the left, with an endometrial thickness of 6 mm. The right ovary appeared normal, while the left ovary was slightly enlarged (4 × 5 cm) and displaced medially, showing a “whirlpool sign” indicative of vascular pedicle twisting, suggestive of left ovarian torsion. In addition, a large exophytic lesion measuring 8 × 6 cm was observed in the left adnexa extending toward the midline, with no solid components or septations, raising the possibility of a left exophytic cyst or paraovarian cyst (Fig. 1a-d).

Given the acute abdominal presentation and ultrasound findings consistent with left ovarian torsion, the patient was taken for emergency laparotomy. Intraoperative findings revealed minimal hemoperitoneum (50–80 mL), a normally sized uterus deviated to the left, and the left fallopian tube was found to be tightly twisted 3 times, with a large hematosalpinx measuring 8 × 6 cm, distended with clots (Fig. 2). The left ovary appeared enlarged and hemorrhagic due to the torsion of the fallopian tube.

Due to complete damage to the left fallopian tube, a left salpingectomy was performed, along with resection of a part of severely necrosed ovarian tissue. The gross appearance of the tube was dark red to black (Fig. 3), and upon sectioning, it was filled with dark reddish-black clots.

Microscopic examination revealed extensive ischemic necrosis of the fallopian tube with extensive hemorrhage throughout, congested blood vessels, and a barely discernible outline of the fallopian tube with no viable tubal tissue (Fig. 4a and b). The part of ovarian tissue also revealed extensive ischemic necrosis with near-total loss of ovarian parenchyma.

The patient experienced an uneventful post-operative recovery and was discharged in satisfactory condition following stitch removal on the 8th day.

DISCUSSION

The incidence of IFTT is estimated at just 1 in 1.5 million women of reproductive age, making it exceedingly rare in adolescent girls [6,7]. While the precise cause remains unclear, research suggests several associations, including prior tubal ligation, hydrosalpinx, pelvic inflammatory disease, a long or congested

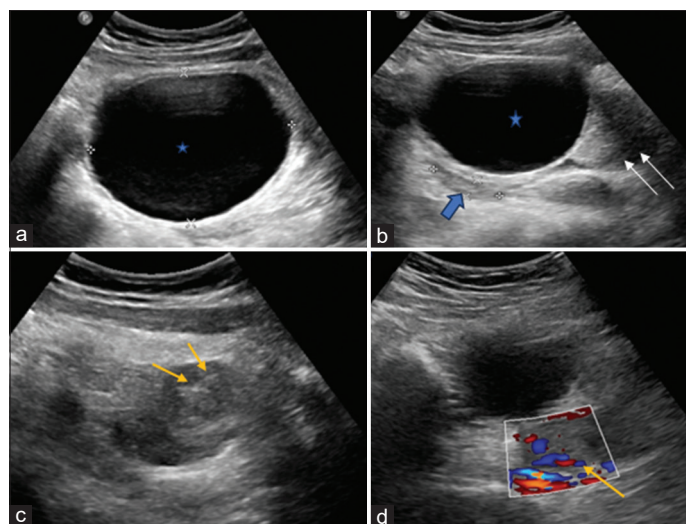


Figure 1: Ultrasonographic image revealing- (a) A thin-walled cystic lesion (asterisk) in the midline arising from left adnexa with few dependent internal echoes. (b) The enlarged left ovary (arrows) was seen abutting the lesion. Possible para ovarian/exophytic ovarian cyst was considered. (c and d) a whirl-like echogenic lesion in left adnexa (arrows) with minimal peripheral vascularity - whirlpool sign s/o twisted pedicle

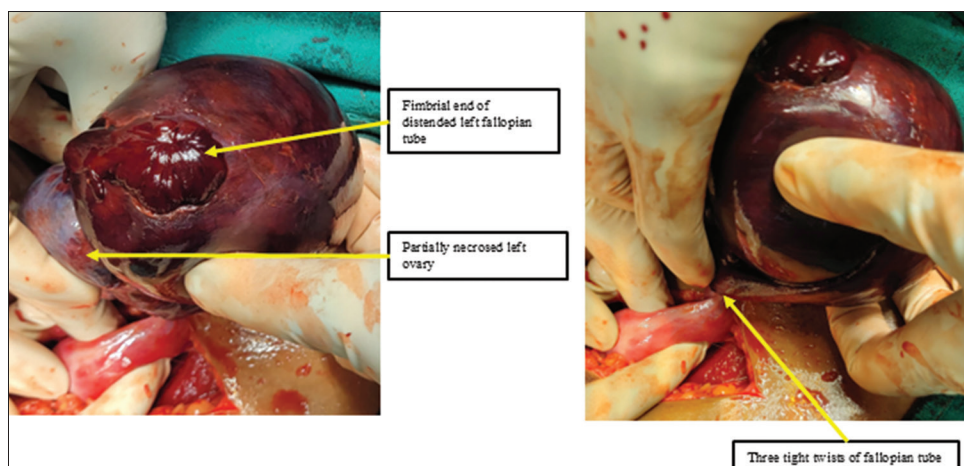


Figure 2: Intraoperative image showing left fallopian tube with three tightly twisted turns, with a large hematosalpinx

Table 1: Comparison of IFTT cases across different age groups, their clinical presentation, and management [15-22]			
Author and study year	Presentation	Final diagnosis	Management
Ziogas <i>et al.</i> , 2020 [15]	A 47-year-old with acute abdomen	Left-sided IFTT without ovarian involvement with left hydrosalpinx	Left salpingectomy with preservation of ovaries
Qian <i>et al.</i> , 2021 [16]	A 13-year-old girl with a 10-day history of right lower abdominal pain	Triple torsion of the right fallopian tube with a 6-cm paraovarian cyst	Tubal conservation surgery
Ali <i>et al.</i> , 2022 [17]	A 32-year-old with 4 days history of severe right lower abdominal pain	Twice-fold twisted ischemic right fallopian tube with hydrosalpinx	Right salpingectomy
Almandeel <i>et al.</i> , 2023 [18]	A 25-year-old newly married woman with acute abdomen	Right-sided IFTT	Right salpingectomy with preservation of right ovary
Thanasa <i>et al.</i> , 2023 [19]	A pregnant multiparous woman at 40 weeks gestation immediately after vaginal delivery presented with acute lower abdominal pain, nausea, dizziness, and vomiting	Left-sided IFTT with left para-tubal cyst	Left salpingectomy with the removal of a para-tubal cyst
Pignataro and Schindler, 2023 [20]	A 15-year-old girl with acute lower abdominal pain, and nausea	IFTT and ipsilateral hemorrhagic ovarian cyst	Unilateral salpingectomy with cystectomy
Jajoo <i>et al.</i> , 2024 [21]	A 33-year-old with a history of bilateral tubal ligation with acute abdomen	Right-sided IFTT with hydrosalpinx	Right salpingectomy with preventive left salpingectomy. Both ovaries were preserved
Toumi <i>et al.</i> , 2024 [22]	A 27-year-old primigravida woman at 37 weeks gestation with severe right lumbar pain	Right-sided IFTT with distal right hydrosalpinx	Cesarean section with right salpingectomy, and ovarian suspension

IFTT: Isolated fallopian tube torsion



Figure 3: Gross appearance of a large hematosalpinx distended with hemorrhage

mesosalpinx, tubal tumors, Morgagni hydatids, and trauma [1,8]. In pubertal girls, the condition may also be linked to congenitally elongated fallopian tubes or developmental anomalies [9]. Moreover, IFTT is predominantly found on the right side, likely due to anatomical considerations. The presence of the sigmoid

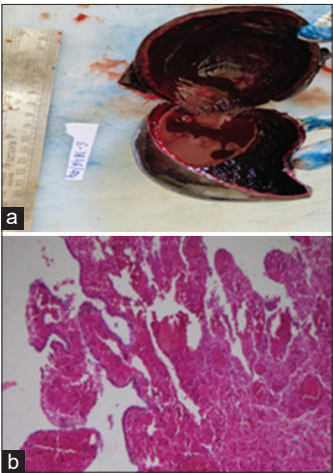


Figure 4: (a) Hematosalpinx on the cut surface of the fallopian tube, filled with hemorrhage. (b) Complete ischemic necrosis of fallopian tube with outlines of luminal plicae (black arrow) (hematoxylin and Eosin stain, ×40)

colon in the left pelvic cavity may restrict mobility in that area, making the right fallopian tube more susceptible to torsion [1]. Common presenting symptoms of IFTT include an acute onset of lower abdominal pain, reported in all cases. This pain may be accompanied by nausea, vomiting, and signs of localized peritoneal irritation. Fever is uncommon, occurring only in the presence of complications such as infection or necrosis [1,10].

IFTT may, in rare instances, be associated with a significant hematosalpinx. Both partial and complete torsion of the fallopian tube can lead to the formation of a hydrosalpinx. If the arterial supply is compromised, this can result in hemorrhagic infarction and subsequently hematosalpinx [11]. In addition, while

exceedingly rare, autoamputation of the fallopian tube may occur as a complication of this condition [12].

Typical ultrasound findings in cases of IFTT include a dilated tube with thickened echogenic walls, the presence of internal fluid with debris, and surrounding inflammation; however, these indicators alone are insufficient for a definitive diagnosis. A key feature is the “beak sign,” which presents as a dilated tubular structure with internal debris and a tapered end. In addition, the “whirlpool sign” may be observed by moving the endovaginal ultrasound transducer over the twisted vascular pedicle. Identifying a normal ovary through ultrasound can further aid in the diagnosis. Ultimately, laparoscopy remains the gold standard for both diagnosing and managing these cases [3,10,11].

Similar to the present case report, a study reported a rare case of hematosalpinx with torsion at its pedicle, accompanied by hemoperitoneum, in a 28-year-old female who presented with acute abdominal pain that closely resembled ovarian torsion [13]. A recent case study described an unusual presentation of hematosalpinx with torsion in a 13-year-old virgin girl who presented with acute abdominal pain accompanied by nausea and vomiting [3]. Another recent case study highlighted an instance of IFTT associated with hydrosalpinx in a 12-year-old girl presenting with acute abdominal pain. She underwent a left salpingectomy, and the histopathological analysis revealed a fallopian tube affected by hemorrhagic infarction [14].

IFTT currently lacks distinct symptoms, clinical findings, imaging characteristics, or laboratory results for definitive identification [10,13]. Imaging studies frequently present non-specific findings, complicating the pre-operative diagnosis and often mimicking other conditions, such as ovarian torsion, paratubal or paraovarian cyst torsion, and ruptured ectopic pregnancy. Consequently, many cases are diagnosed late, delaying essential interventions that can compromise tubal function. Torsion can lead to serious complications, including hematosalpinx, hemoperitoneum, and necrosis of the tube, all of which necessitate urgent surgical management [3,13]. Management typically involves the surgical removal of the affected fallopian tube. If the ovary cannot be preserved, the tubo-ovarian mass may also need to be excised. With timely surgical intervention, the overall prognosis is favorable [3]. Table 1 compares IFTT cases across different age groups, their clinical presentation, and management [15-22].

CONCLUSION

Hence, IFTT though uncommon, must be considered in the differential diagnosis of acute abdomen in young girls and women of reproductive age. Early recognition and timely surgical intervention are vital to preventing complications and preserving tubal function, as partial or complete torsion can lead to serious issues such as hematosalpinx, hemoperitoneum, and necrosis of the tube. This underscores the critical need for clinicians to maintain awareness of this rare yet significant condition.

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