A rare case of sclerosing stromal tumor of ovary with Meigs syndrome presenting with torsion

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INTRODUCTION
A solid ovarian tumor with raised CA 125 levels points towards a possibility of malignant tumor. In a young patient with such clinical scenario there is always a dilemma regarding conservation of uterus and other ovary. Sclerosing stromal tumor (SST) is benign solid ovarian neoplasm of the sex cord stromal category. It is an extremely rare tumor presenting in the second and third decades of life, and they are unilateral.¹ The tumor with rare exception is hormonally inactive. All reported cases of SST have been benign.¹ This is a rare presentation of SST with Meigs syndrome with elevated serum Ca125 levels.

CASE REPORT
A 24 year old primipara, presented with pain in abdomen and abdominal distention for a month. There was no history of amenorrhoea. Clinical examination revealed a large palpable abdomino-pelvic mass, ascites. She was admitted in gynaecology ward and was investigated. Her CA 125 level was 339.20 IU/ml. Ultrasonography revealed a mass 15 cm × 14 cm × 6 cm arising from right ovary with solid and cystic areas. CT scan demonstrated large pelvic mass accompanied by ascites and right pleural effusion (Figure 1). X-ray chest showed 600-700 cc pleural effusion with collapse of underlying lung field (Figure 2).

On second day patient had breathlessness, fever, nausea and acute abdominal pain. Intercostal chest drain was kept after ruling out malignancy by diagnostic pleural tapping. Approximately 800 cc haemorrhagic pleural fluid was drained. Decision of exploratory laparotomy was taken with suspicion of torsion. On laparotomy a large bosselated right ovarian tumor with torsion involving fallopian tube was noticed with gross ascitic (Figure 3). About 550 cc ascetic fluid was drained. Ascitic fluid was sent for cytology. Left ovary, fallopian tube & uterus were unremarkable. There was no evidence of any other mass or metastasis in the peritoneal cavity.

Right salpingo-oophorectomy was performed. Frozen sections of the tumor demonstrated benign sclerosing stromal tumor of ovary (Figure 4). Ascitic fluid was negative for malignancy. Histopathology later confirmed the diagnosis. Patient had an uneventful post-operative recovery. with spontaneous resolution of ascites and pleural effusion (Figure 5).

DISCUSSION
Sclerosing stromal cell tumor is a rare benign tumor of ovarian stroma, first described by Chalvaridjian and Scully (1973),² SSTs occur in the second or third decades.¹ The most common presentation includes menstrual irregularity, pelvic pain and non-specific symptoms related to ovarian mass.³ Our patient had pelvic mass, pain but no menstrual irregularities. In 1989, Jones and Surwit reported on patients with Meigs syndrome presenting with torsion.¹ This is a rare presentation of SST with Meigs syndrome with elevated serum Ca125 levels.
Junnare, et al.: Sclerosing stromal tumor of ovary with Meigs syndrome presenting with torsion – A case report

Figure 1: CT Abdomen and pelvis – Solid ovarian mass

Figure 2: X ray chest – Right sided pleural effusion (pre operative)

Figure 3: SST with Torsion 18 cm x 14 cm x 7 cm with bosselated external surface. Cut section was solid grayish-white & variegated with yellowish cystic areas

Figure 4: Hypocellular & hypercellular areas with vagelobulation of cellular areas along with many ectatic thin walled blood vessels

Figure 5: X ray chest PA view – One week after surgery

syndrome associated with fibroma – thecoma of the ovary and elevated CA 125 levels. Since then 38 cases have been reported, including four with SSTs. We report fifth case of Meigs syndrome associated with SST of the ovary.

In our case, patient had pain, abdominal mass, ascites, pleural effusion who later developed torsion. M.V.C. de Silva et al have reported an unusual case of SST who presented as torsion.

The mechanism by which peritoneal and pleural effusion develops is still unknown. Meigs suggested that irritation of the pleural surface or leakage from edematous ovarian stroma of the tumor could stimulate peritoneal fluid production. The other mechanism includes obstruction or congestion of peritoneal lymphatics & veins by the tumor, increased vascular permeability or transudation by tumor surface.

CA 125, a high molecular weight glycoprotein is now the most commonly used serum biomarker for ovarian tumors, though it has low specificity as increased level are found in so many malignancies and non-malignant condition. In Meigs syndrome, the elevation of serum CA125 levels may result from mesothelial expression of the tumor marker. A significant linear correlation between volume of pleural effusion and serum Ca125 values has been demonstrated in both benign and malignant diseases. Similar correlation has been demonstrated between CA125 elevation and volume of ascites in Meigs syndrome. Our patient had a raised
level of CA125 of 339.20IU/ml with significant ascites and large pleural effusion of 800 cc.

On Ultrasound the appearance of SSTs may be suspected to be malignant ovarian tumors because they show mixed pattern with cystic and solid components. In our case, on USG mass was proved to be solid & cystic & measured approximately 15 cm × 14 cm × 6 cm. CT scan demonstrated large pelvic mass accompanied by ascites & right pleural effusion. Some studies have suggested that MRI findings may be more specific in distinguishing this benign neoplasm from other malignant ovarian tumors which include a large mass with hyperintense cystic components or heterogenous solid mass of intermediate-to-high signal intensity on T2 weighted MRI and thus help in preoperative diagnosis, thereby preventing excessive surgical intervention.

CONCLUSION

Due to extremely low incidence of SST, it is not always easy to predict its presence preoperatively. Clinical and ultrasonographic findings may be suggestive of malignant neoplasm. In a young patient with solid ovarian mass, possibility of SST should be kept in mind. Frozen section can help in conserving the fertility.

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REFERENCES


Authors Contribution:
JK – Operated the patient, prepared the manuscript; PT – Involved in surgery and patient management; PN – Supervised management, reviewed the manuscript.

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