Clinical Summary

A 17 year old girl was presented with painless abdominal distention. Physical examination revealed a palpable abdominal mass. Laboratory tests were normal.

Clinical History and Imaging Procedures

The patient was admitted to the hospital with painless abdominal distention and underwent ultrasonographic evaluation, which revealed a large left abdominal mass. The lesion was heterogeneous, mostly solid with a small cystic component, demonstrating prominent vascularity. A CT scan was done, revealing a well circumscribed mass, with 18 x 17 cm, localized predominantly in the left abdomen, originating from the right ovary. This mass was heterogeneous with an irregular shaped central cystic area, a lobulated soft tissue density component and a peripheral left hypodense area. After contrast administration there was enhancement of the central mass (Fig. 1a) and prominent vessels converging to the hilum were evident (Fig. 1b). Increased caliber of right ovarian vessels was also noted (Fig. 1b). On MRI the central mass was moderately hypointense on T1WI (Fig. 2), of intermediate signal intensity on T2WI (Fig. 3 a,b), with an irregular central cystic area. It was surrounded by a smoothly marginated solid component with hypointense capsule on T1WI and T2WI, crossed by several vessels. There was enhancement of the central mass and of the tumour capsule after gadolinium administration (Fig. 4).

Surgery was done, with total resection of the mass and partial resection of the right fallopian tube (Fig. 5).

The histological diagnosis was Sex Cord Stromal Tumour of Undetermined Pattern of the right ovary,
consisting on two distinct components, one external corresponding to a fibrothecoma, and one internal of stromal origin with an undetermined pattern (Fig. 6,7).

Discussion

Sex cord stromal tumours derive from coelomic epithelium or mesenchymal cells of the embryonic gonads. This group of tumours represents approximately 8% of ovarian neoplasms and affects all age groups. The most common types are granulosa cell tumours, fibrothecomas and Sertoli-Leydig cell tumours. Fibromas and fibrothecomas belong to a spectrum of benign tumours. Lipid-rich thecoma demonstrates estrogenic activity and few fibroblasts. Fibroma is composed of spindle cells that form collagen, has no thecal cells and demonstrates no estrogenic activity.

A broad spectrum of ultrasonographic features is seen and in most cases the appearance of the tumour is non-specific. CT shows a homogeneous solid tumour with delayed enhancement.

Ovarian fibromas usually display low signal intensity on both T1WI and T2WI. High signal intensity on T2WI corresponds to regions of hyalinization and myxomatous changes. Intratumoral oedema is also a feature of larger fibromas / fibrothecomas.

Differential diagnosis in this age group should include sclerosing stromal tumour (SST). SST occur predominantly in the second and third decades of life in 80% of cases, usually appearing as a large mass with hyperintense cystic components and a heterogeneous solid component with intermediate to high signal intensity on T2WI and early peripheral enhancement with centripetal progression.

Dysgerminoma is another entity to be considered. Characteristic imaging findings include a multilobulated solid mass with prominent fibrovascular septa and areas of necrosis and haemorrhage.

Knowledge of key imaging features of sex cord stromal tumours may provide criteria to narrow the differential diagnosis of adnexal masses.

Final Diagnosis

Sex Cord Stromal Tumor of Undetermined Pattern

Figures
Figure 1 Axial CT after contrast administration

Enhancement of the central mass and capsule with prominent vascularity

Increased caliber of right ovarian vessels converging to the hilum

Figure 2 Axial T1WI

Moderately hypointense central mass surrounded by a smoothly marginated hypointense solid component.
Figure 3 T2WI

Sagittal T2WI showing a mass of intermediate signal with irregular central cystic area, surrounded by a hyperintense component, traversed by several vessels. This mass was delineated by a hypointense capsule.

Coronal T2WI showing a mass of intermediate signal with irregular central cystic area, surrounded by a hyperintense component, traversed by several vessels. This mass was delineated by a hypointense capsule.

Figure 4 Axial fat-suppressed T1WI GRE after gadolinium administration

Enhancement of the central mass and of the tumor capsule.

Figure 5 Surgical specimen

In the surgical specimen we can appreciate three sections of the right ovary.
Figure 6 Photomicrograph
Peripheral component: collagen fibres with fibroblasts and theca cells characteristic of fibrothecoma.

Figure 7 Photomicrograph
Central mass: monomorphic cell population with moderate atypia forming trabecules. Some cells show mitotic activity.

MeSH

**Ovary** [A05.360.319.114.630]
The reproductive organ (GONADS) in female animals. In vertebrates, the ovary contains two functional parts: the OVARIAN FOLLICLE for the production of female germ cells (OOGENESIS); and the endocrine cells (GRANULOSA CELLS, THECA CELLS, and LUTEAL CELLS) for the production of ESTROGENS and PROGESTERONE.

**Sex Cord-Stromal Tumor** [C04.557.475.750]
A malignant neoplasm of the ovary or testis. These tumors differentiate toward sex cords (in embryonic gonads) in the form of female (i.e., granulosa and theca) cells, male (i.e., Sertoli and Leydig) cells, or indifferent elements. In the ovary, sex cord-stromal tumors comprise 5% of all ovarian neoplasms. In the testes, Leydig and Sertoli cell tumors comprise about 5% of all testicular neoplasms, 10% of which behave in a malignant fashion. (From Segen, Dictionary of Modern Medicine, 1992)

References


Solid and Cystic Components That Mimic Malignancy. AJR Am J Roentgenol 182:1259-1265


Citation

C. Bahia1, M. O. Castro2, G. Bastos1, J. Parra1, E. Vieira1, T. M. Cunha3, F. Aleixo1
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