

## Juvenile granulosa cell tumour of ovary: a casr report

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### ABSTRACT

Juvenile Granulosa Cell Tumor (JGCT) is a rare ovarian neoplasm occurring predominantly in children and adolescents. We report a case of a 5-year-old girl presenting with abdominal pain, early satiety, and vaginal bleeding. Imaging revealed a large solid-cystic ovarian mass, with elevated serum Inhibin B and CA-125 levels. The patient underwent left salpingo-oophorectomy and infra-colic omentectomy. Histopathological examination confirmed JGCT. Early diagnosis and surgical management are key to a good prognosis. Regular follow-up is essential due to the tumor's potential for delayed recurrence.

**Keywords:** Juvenile Granulosa Cell Tumor, Ovarian Neoplasm, Pediatric Gynecology, Inhibin B, Pseudo-precocious Puberty, Oophorectomy.

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### INTRODUCTION

Granulosa Cell Tumor(GCT), rare entity of low-grade malignant tumor with indolent course. Here we present a case of Juvenile GCT.

### CASE PARTICULARS

A 5 year old girl of Nadia referred to MCK with primary diagnosis of abdominal mass. C/O pain in abdominal for 2 months. Child's mother observed early satiety, abdominal hardness and vaginal bleeding for few months.

O/E a solid-cystic mobile abdominal mass of 15X10cm is found. Breast corresponds Tanner stage 2. Axillary and pubic hairs absent.



Figure 1: PATIENT WITH ABDOMINAL DISTENSION, TANNER 2 OF BREAST DEVELOPMENT

USG stated solid-cystic mass (15X15cm); ovarian origin. MRI-abdomen revealed complex ovarian SOL, 9.4X14.3X13.9cm. Serum Inhibin B 1280mcg /dl, CA125 109.5 U/ml (both elevated) and CAE, AFP, HCG levels are normal. Multi-disciplinary team involved.

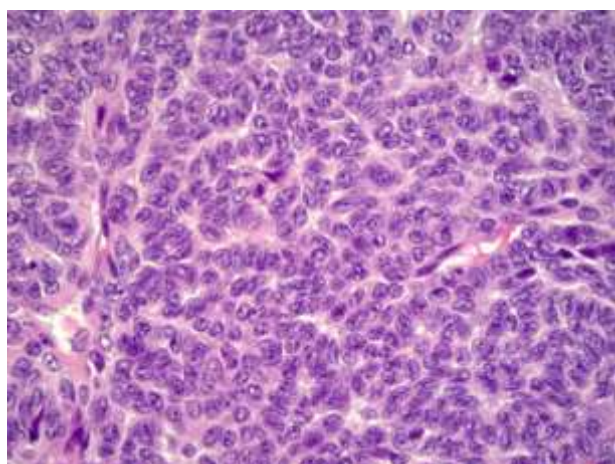


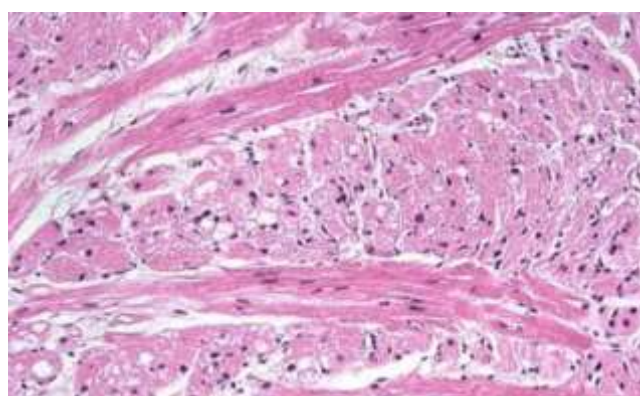
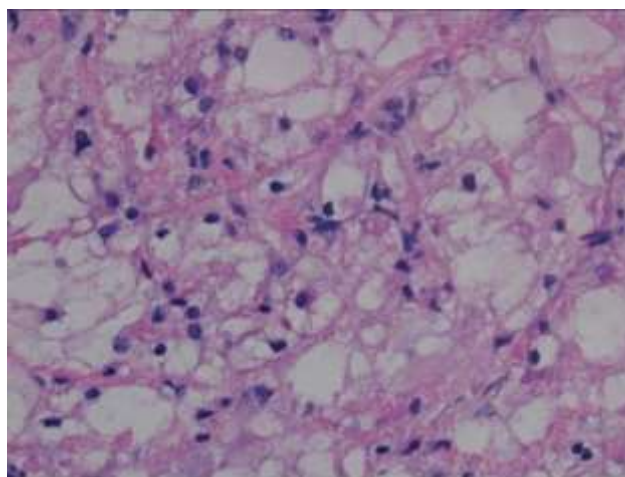
Figure 2: LEFT SIDED TUBE AND OVARIAN TUMOR

She underwent Laparotomy GA. No ascites. Left Salpingo-Oophorectomy with Infra-colic Omentectomy (thickened +areas of clumping) done. Right-sided FT and ovary normal, kept for future fertility and ovarian function. Uneventful postoperative period.



Figure 3: RESECTED OVARIAN TUMOR  
HPE confirmed the diagnosis of Juvenile GCT.





**Hematoxylin & Eosin stain (H&E) , 40 X**

## DISCUSSION

Juvenile GCT, a sex Cord- Stromal Tumor occurs in young. B/L in 2% cases. Pseudo-precocity because of estrogen secretion, vaginal bleeding is common. Serum Inhibin B & AMH produced by Granulosa cells, potential markers of GCT. Early diagnosis predicts favorable prognosis. Primary therapy is surgery; radiation and chemotherapy reserved for recurrences and metastatic cases.

## CONCLUSION

Regular follow up is necessary; indolent nature of recurrence.

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