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Title: DYSGERMINOMA MASQUERADING AS NEOPLASTIC LESION "A DIAGNOSTIC DILEMMA".





INTRODUCTION

A Dysgerminoma is a germ cell tumor ,it is rare and typically slow growing.

Most commonly occurring in young women, particularly between the ages of 10 and 30. Presents with abdominal pain, swelling, abnormal



CASE STUDY

A 32yrs P2L2 with previous 2 LSCS with c/o pain abdomen since2 years which was insidious in onset, dull aching, intermittent ,non radiating . On examination: smooth ,irregular mass of 20*15cm in right iliac fossa ,non tender, firm in consistency, limited mobility from side to side . On pelvic examination: mass corresponds to 24-26 weeks of gestation ,uterus could not be made separately,Anterior and Right Forniceal fullness , No Forniceal tenderness . Tumor markers like CEA,CA 19 -9,alpha Feto protein ,LDH found to be normal,

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MRI: right sided malignant solid ovarian tumor (10*15*17), minimal ascites ,no evidence of any pelvic lymphadenopathy is seen.

Diagnosis of Right side malignant solid ovarian tumor was made and was taken for staging laparotomy

Laparotomy findings: lobulated right ovarian mass approximately 15*10*12 cm ,right fallopian tube normal ,left tube and ovary normal.

Specimen sent for histopathological examination and was found to be dysgerminoma with no focal capsule invasion, Grade 2 FIGO stage 1A. pT1aNx.

Postoperatively patient was stable and discharged on day 7.

CONCLUSION

- Dysgerminoma can present in early 3rd decade of life as solid cystic masses and hence high degree of clinical suspicion is required to investigate, treat and follow up patient.
- Rarely present with elevated CA 125 levels and Beta HCG in the absence of choriocarcinoma, since choriocarcinoma is aggressive tumor proper evaluation of tumor is required.
- In present case we were able to withhold chemotherapy and patient was asked to follow up.

REFERENCES

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