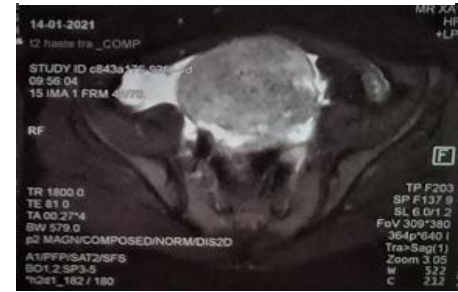
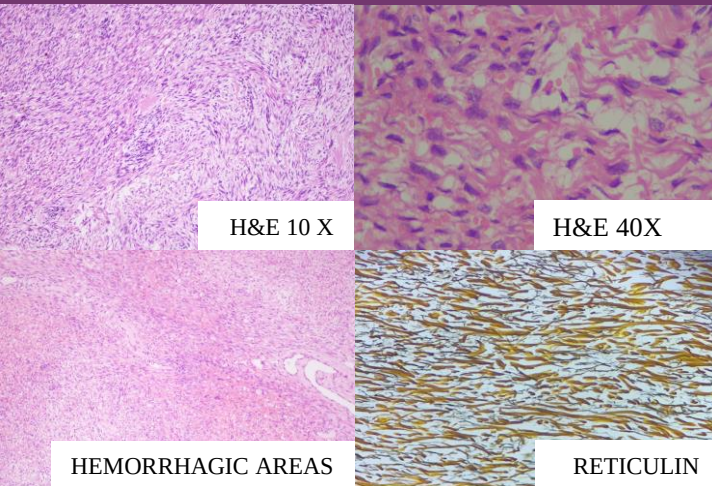


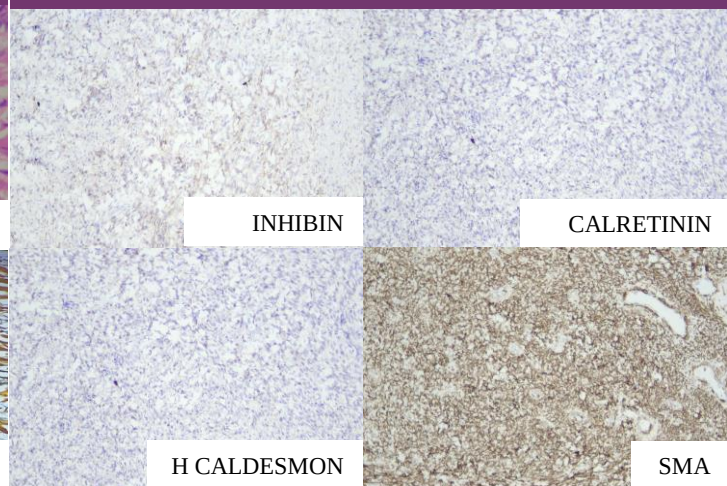
CT IMAGE - OVARIAN MASS



HISTOPATHOLOGICAL EXAMINATION



IMMUNOHISTOCHEMISTRY



- ❖ **Gross examination** showed an encapsulated ovarian mass with tan white firm cut surface & areas of congestion.
- ❖ **Histopathological examination** showed a variably cellular tumor consisting of fascicles & sheets of bland fibroblastic cells with scant eosinophilic cytoplasm & spindled to fusiform nuclei. These cells showed indistinct cell borders & were seen to blend with extracellular collagen/surrounding stroma. Areas of fresh hemorrhage noted. No evidence of significant nuclear atypia or mitosis identified.
- ❖ **Uterus, cervix, other ovary, bilateral fallopian tubes, all pelvic lymph nodes** were unremarkable. Peritoneal wash cytology, omental & peritoneal biopsy were within normal limits.
- ❖ **Special histochemistry & IHC** were done to confirm the diagnosis and to exclude remote possibility of **leiomyoma/ granulosa cell tumor**.

FINAL DIAGNOSIS - OVARIAN FIBROMA WITH MEIGS SYNDROME

The ascites and pleural effusion resolved post surgery, confirming the final diagnosis.

DISCUSSION

MEIGS SYNDROME

- ❖ Meigs syndrome is a rare benign condition, occurring in less than 1% of ovarian tumors, with a classical triad of ovarian fibroma, ascites and pleural effusion & is uncommon before 30 years of age.
- ❖ The pathophysiology behind Meigs syndrome is due to seepage of fluid from ovary into the peritoneal cavity and then into one or both pleural cavities via lymphatics.
- ❖ Recent studies on the pathogenesis of Meigs syndrome suggest role of VEGF, which increases vascular permeability and thus causes fluid accumulation.
- ❖ Its clinical presentation, often associated with a raised CA-125 levels, mirrors that of an ovarian malignancy.
- ❖ Our index case is 30 yr/F with a triad of ovarian mass, ascites & right hydrothorax but normal CA-125 levels.