Case Report

A 48-year-old female (P1L1) came to Obstetrics and Gynaecology OPD of K.S Hegde Charitable Hospital, Mangalore, India with history of pain abdomen which was dull aching and non radiating. There were no history of dysmenorrhoea, menorrhagia, burning micturition, bowel and bladder disturbances. With proper patient consent, per abdomen examination revealed a nodular, soft to firm ill defined mass measuring 10x5 cm was present in the left iliac fossa reaching the hypogastrium. CT pelvis showed a left ovarian mass lesion with predominantly cystic with focal solid areas [Table/Fig-1a]. Uterus, right ovary and other abdominal organs were normal. A clinical diagnosis of complex ovarian cyst was made. CA-125 was 52.5 U/ml (normal value <35U/ml). Other routine biochemical and haematological parameters were within normal limits. Laparotomy was performed and entire specimen was sent to pathology department for histopathological examination. Postoperative period was uneventful. The patient on follow up is doing well without any recurrence or metastasis as of six months after surgery.

Grossly, we received a specimen of total abdominal hysterectomy with bilateral salpingo-oophorectomy, omental tissue and left and right iliac lymph nodes. The left ovarian tumour measured 16x9x6.5 cm. Cut section revealed well encapsulated, multiloculated cystic tumour with focal solid areas [Table/Fig-1a]. Uterus, right ovary and other abdominal organs were normal. A clinical diagnosis of complex ovarian cyst was made, CA-125 was 52.5 U/ml (normal value <35U/ml). Other routine biochemical and haematological parameters were within normal limits. Laparotomy was performed and entire specimen was sent to pathology department for histopathological examination. Postoperative period was uneventful. The patient on follow up is doing well without any recurrence or metastasis as of six months after surgery.

Microscopically, solid areas showed proliferating glands lined by tall columnar cells having pleomorphic nuclei with stratification, prominent nucleoli and hyperchromatism [Table/Fig-2a,b]. Plenty of intracellular mucin was also seen. Focal areas showed cyst wall lined by stratified squamous epithelium associated with osteoid and keratinous material [Table/Fig-3a,b]. No evidence of immature...
teratoma component was seen. Sections from the capsule showed no tumour infiltration. Omentum, right ovary, uterus, cervix, right and left iliac lymph nodes were free from the tumour. The final diagnosis of mucinous cystadenocarcinoma (grade 2) co-existing with mature cystic teratoma, stage la (FIGO) of the left ovary was made.

**DISCUSSION**

Mucinous cystadenocarcinomas are only one-third as common as serous cystadenocarcinomas and occur in the age group of 45-65 years. They are typically large, unilateral, multicystic cystic masses containing watery or viscid secretions with smooth white capsules and have average size of 18-22 cm. They may have solid areas with foci of haemorrhage and necrosis [1-3]. Microscopically they show complex papillary areas or back to back glands lined by pleomorphic cells with stromal invasion. Mucinous cystadenocarcinomas may arise de novo or transformation from a benign mucinous cystadenoma which is estimated around 12%-23% [2,3]. Approximately 70% of these patients present with advanced stages i.e. would have metastasised to upper abdomen and beyond abdominal cavity because these patients mistake the initial symptoms as other minor GIT disorders.

Mature cystic teratoma of the ovary (dermoid cyst) comprises 27%-44% of all ovarian tumours. It is the most common germ cell neoplasm occurring in reproductive age group and usually presents with a mass, but 25% are detected incidentally. It presents as multiloculated masses and contain cheese like cystic component composed of keratin, sebum and hair [2,3].

Mucinous tumours can rarely co-exist with other tumours like mature cystic teratoma, serous tumours, Brenner tumours, thecoma etc. The incidence of mucinous tumours co-existing with teratoma is just 3%-8% [4]. However, mucinous tumours can be surface epithelial origin or teratomatous origin. In our case, since mucinous cystadenocarcinoma component is a predominant tumour with only focal teratomatous component, we made a diagnosis of co-existence of these tumours. However, immunohistochemistry helps to differentiate the exact origin of mucinous tumours. CK 7 is positive and CK 20 negative in surface epithelial origin; whereas CK7 is negative and CK 20 positive in teratomatous origin [4-8]. Immunohistochemistry in our case revealed strong membrane positivity for CK7 and negativity for CK 20, hence confirming the histogenesis of mucinous cystadenocarcinoma from surface epithelium of the ovary [Table/Fig-4.a,b].

On reviewing literature, we found Okada et al., in 2004 reported only 2 cases (out of 11) of mucinous cystadenocarcinoma co-existing with mature cystic teratoma [9]. Russel Van et al., reported 44 cases of ovarian mucinous tumours associated with mature cystic teratoma out of which six were malignant [4]. Jesse K Mc Kenney et al., reported 5 out of 42 cases of ovarian mucinous cystadenocarcomas associated with mature cystic teratoma [5].

**CONCLUSION**

Majority of these studies and our experience showed that, the distinction between benign, borderline, and malignant tumour are generally not possible by radiological studies and only diagnosed by histopathological examination. Hence, histopathological examination and immunohistochemistry plays a significant role in accurate diagnosis and management of these patients. So, we should be aware of these rare co-existent tumours and meticulous dissection should be done to look for any synchronous tumours or malignant areas; since management and prognosis vary significantly depending upon the microscopic type and stage.

**REFERENCES**


**PARTICULARS OF CONTRIBUTORS:**

1. Postgraduate, Department of Pathology, KS Hegde Medical Academy, Mangalore, India.
2. Associate Professor, Department of Pathology, KS Hegde Medical Academy, Mangalore, India.
3. Professor, Department of Pathology, KS Hegde Medical Academy, Mangalore, India.
4. Postgraduate, Department of Pathology, KS Hegde Medical Academy, Mangalore, India.
5. Professor and Head, Department of Pathology, KS Hegde Medical Academy, Mangalore, India.

**NAME, ADDRESS, E-MAIL ID OF THE CORRESPONDING AUTHOR:**

Prachi Kukreja, Postgraduate, Department of Pathology, KS Hegde Medical Academy, Nithyananda Nagar Post, Deralakatte, Mangalore- 575018, India.

Email: prachikukreja@gmail.com

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