E-ISSN:2456-1487 P-ISSN:2456-9887 RNI:MPENG/2017/70771

Research Article

Ovarian tumors

Tropical Journal of Pathology and

Microbiology



2021 Volume 7 Number 6 November December

A prospective report on the histopathological study of ovarian tumors in a tertiary care centre

Unissa R.¹, Shameem K.^{2*}, Saleem H.³, Pramida B.⁴, Bhavani M.⁵, Chandra T.⁶

DOI: https://doi.org/10.17511/jopm.2021.i06.04

¹ Rahmath Unissa, Resident, Department of Pathology, Kamineni Institute of Medical Sciences, Narketpally, Telangana, A.P, India.

- ^{2*} Khatija Shameem, Associate Professor, Department of Pathology, Kamineni Institute of Medical Sciences, Narketpally, Telangana, A.P, India.
- ³ Hafsa Saleem, Resident, Department of Pathology, Kamineni Institute of Medical Sciences, Narketpally, Telangana, A.P, India.
- ⁴ B Pramida, Resident, Department of Pathology, Kamineni Institute of Medical Sciences, Narketpally, Telangana, A.P, India.
- ⁵ M Bhavani, Prof & HOD, Department of Pathology, Kamineni Institute of Medical Sciences, Narketpally, Telangana, A.P, India.

⁶ T Jaya Chandra, Professor, Department of Microbiology, GSL Medical College, Rajahmundry, A.P, India.

Introduction: Ovarian tumors (OTs) are the most notorious gynecological lesions in the reproductive age group. With this, a study was conducted to categorize the ovarian lesions into benign, borderline and malignant lesions and also to study their laterality, gross and microscopic patterns. Settings: The study was conducted in the Department of Pathology, KIMS, Narketpally, Telangana from January 2017 to December 2019, 36 months. Random sampling was considered. Women aged >18yrs, histologically proven OTs, sized > 5 cm diameter, were included. Abdominal hysterectomy specimen were considered. Gross examination was done. The features such as size, the color of the specimen were noted. The stained sections were examined under a light microscope for histopathological diagnosis. Results: Out of 94 participants, 79 were benign tumors (Bet), 5 were borderline tumour (BoT), and 10 were malignant tumor (MIT). Among the BeT, maximum were diagnosed in 21 - 40 years group, 21 - 40 years group in BoT and 41 - 60 years for MIT. In this research, 91% were unilateral, and Seromucinouscystadenoma was the common bilateral tumor. Among the different histological patterns, surface epithelial tumours constituted the majority. Conclusion: Maximum number of tumour cases were benign, reported in the reproductive age group, whereas the malignant neoplasms in > 40 yrs. The present study emphasizes the need for proper histopathological evaluation and screening at all ages due to the relative predominance of OTs to rule out malignancies.

Keywords: Ovarian tumors, Histopathology study

| Corresponding Author | | How to Cite this Article | | To Browse | |
|--|---|---|---|--|--|
| Khatija Shameem, Associate Professor, Department of Pathology, Kamineni Institute of Medical Sciences, Narketpally, Telangana, A.P, India. Email: drkhatija@gmail.com | | Rahmath Unissa, Khatija Shameem, Hafsa Saleem, B Pramida, M Bhavani, T Jaya Chandra, A prospective report on the histopathological study of ovarian tumors in a tertiary care centre. Trop J Pathol Microbiol. 2021;7(6):292-297. Available From https://pathology.medresearch.in/index.php/jopm/ar ticle/view/581 | | | |
| eceived 21 | Review Round 1 2021-10-23 | Review Round 2 2021-10-30 | Review Round 3 2021-11-06 | Accepted 2021-11-1 | |
| iterest | Funding | Ethical Approval | Plagiarism X-checker | Note | |
| | em, Associate Pr (amineni Institute langana, A.P, Indi a@gmail.com | em, Associate Professor, Department (amineni Institute of Medical Sciences, langana, A.P, India. (aggmail.com eccived 21 Review Round 1 2021-10-23 | em, Associate Professor, Department Rahmath Unissa, Khatija Shama fangana, A.P, India. Pramida, M Bhavani, T Jaya Cl report on the histopathologic tumors in a tertiary care co Microbiol. 2021;7(6):292-297. Available From https://pathology.medresearch. ticle/view/581 eccived Review Round 1 Review Round 2 21 2021-10-23 2021-10-30 | em, Associate Professor, Department Rahmath Unissa, Khatija Shameem, Hafsa Saleem, B ramineni Institute of Medical Sciences, Pramida, M Bhavani, T Jaya Chandra, A prospective langana, A.P, India. report on the histopathological study of ovarian ucore centre Trop J Pathol Microbiol. 2021;7(6):292-297. Available From https://pathology.medresearch.in/index.php/jopm/ar ticle/view/581 | |

Introduction

Worldwide, the mortality rate for ovarian tumors(OT) is around 4/100,000 and the seventh leading cause of death.[1] Indian reports suggest a rise in OT and the incidence rate ranging between 0.2% to 2.5%.[2] Mass in the ovary is a common complaint in the gynecology outpatient. Nulliparous nature and family history are the two risk factors being recognized to get these OT.[3]

These are notorious, exhibit mild symptoms, asymptomatic and challenging to diagnose until cause pressure such as ascites, abdominal distension due to increase in size. [2].

There is a wide range of histological differentiation, and numerous types were reported in both benign and malignancy. The majority (80%) are benign, and 20 to 45 years is the common age group affected. At the same time, malignant cases are common in older women with poor prognoses [4]. Complete evaluation before surgery form the basis for treatment and follows up.[5] The newer diagnostic tests such as ultrasonography, tumor markers are helpful for early diagnosis as well as for proper clinical evaluation. [6]

But the histological can give us a definitive diagnosis, typing and grading, which can guide the clinician for prognosis and treatment. [5]. A study was conducted to find the overall incidence, pattern and various histopathological types of ovarian neoplasms.

Materials and Methods

Settings: The study was conducted in the Department of Pathology, KIMS, Narketpally, Telangana.

Duration and type of study:Thus was a prospective study conducted from January 2017 to December 2019, 36 months.

Sampling method: Random sampling was considered.

Sample size calculation: All the eligible members who satisfy the inclusion criteria were considered in this study.

Inclusion criteria: Women aged \geq 18yrs, those histologically proven OTs, sized \geq 5 cm diameter were included.

Exclusion criteria: Women with follicular cysts, haemorrhagic inclusion cysts, endometriosis, those who were non-cooperative and didn't submit the consent were not considered.

Data collection, procedure: Specimen received as a solitary specimen, or part of total abdominal hysterectomy specimen were considered. On the receipt of the clinical specimen, the gross examination was done. The features such as size, the color of the specimen were noted. As a part of the study, the external surface features and contents were reported and recorded in the proforma.

After 24 to 48 hrs of fixation, multiple bits were taken from the representative areas, which may be prone to be OT. Tissue was processed, and paraffin blocks were made. The tissue sections of 5 microns were cut and stained using Hematoxylin and Eosin. Sections were cleared by xylene and mounted on a glass slide. Each biopsy was labelled explicitly according to the orientation of the biopsy site and sent for histopathological examination. The stained sections were examined under light microscope for histopathological diagnosis.

Special stains such as periodic acid Schiff and reticulin stains were done whenever necessary. CA 125 levels were measured for clinically and radiologically suspected to have OTs.In the case of Granulose cell tumour, IHC -Inhibin was done.

Ethical consideration and permission: The institutional ethics committee approved the study protocol. Informed written consent was taken from all the study participants. If required, consent was also taken in the presence of the witness.

Statistical analysis: SPSS21.0 was used for the analysis of the data. Various non-parametric tests were used. The data were presented in percentages.

Results

SETs constituted the majority, and among the individual tumors, serous cystadenoma was the commonest benign epithelial tumors.

During the study period, a total of 94 participants were included. Among these, 79(84%) were benign tumors (Bet), 5(5.31%) were borderline tumor (BoT) and 10(11%) cases were malignant tumor (MIT) (Table 1).

Table 1: Incidence of various OTs among thestudy participants.

| Type of tumour | Number | % |
|----------------|--------|-----|
| Benign | 79 | 84 |
| Malignant | 10 | 11 |
| Borderline | 5 | 5.3 |
| Total | 94 | 100 |

Malignant tumors are the second common, followed by borderline.

The age was ranged between 18 - 75 years. Among the BeT, the maximum was diagnosed in the 21 - 40 years group, followed by 41 - 60 and 61 - 80 years. In BoTs also, 21 - 40 years group is the commonly involved. In the MIT category, maximum cases were detected in the 41 - 60 years group (Table 2).

Table 2: Age-wise distribution of OTs amongthe study participants; n (%).

| Age | ВеТ | ВоТ | МІТ |
|----------|-----------|---------|---------|
| <20 | 2 (2.1) | 0 | 0 |
| 21 - 40 | 49 (52.1) | 3 (3.2) | 0 |
| 41 - 6 0 | 25 (26.6) | 1 (1) | 9 (9.6) |
| 61 - 8 0 | 3 (3.2) | 1 (1) | 1 (1) |
| Total | 79 (84) | 5 (5.3) | 10 (11) |
| | 94 (100) | | |

Maximum malignant tumours are detected in the 41 – 60 years group.

In this research, 91% were unilateral, and just 8.5% were bilateral, and all these were BeT; Seromucinouscystadenoma was the common bilateral tumor. In the unilateral tumors, the majority were on the right side, and most of the MIT were on the right side (Table 3).

Table 3: Site of involvement of OTs among the study members; n (%).

| Type of tumour | Unilateral | | Bilateral | |
|----------------|------------|-----------|-----------|--|
| | Right | Left | | |
| Benign | 41 (43.6) | 30 (32) | 8 (8.5) | |
| Borderline | 1 (1) | 4 (4.3) | 0 | |
| Malignant | 9 (9.6) | 1 (1) | 0 | |
| Total | 52 (55.3) | 35 (37.2) | 8 (8.5) | |
| | 94 (100) | | | |

MTs are unilateral, most of these on the right side.

In this research, most (71; 75.5%) of the OTs were grossly cystic cases, followed by mixed tumors (20; 21.27%) and solid tumors (3: 3.2%). In the

Benign group, the majority were cystic (61), followed by mixed (16) and solid (2). All the borderline (5) were cystic, and out of 8 MITs, four mixed, three were cystic, and one was solid (Table 4).

Table 4: Consistency of OTs among the studyparticipants; n (%).

| Consistency | Benign | Borderline | Malignant | Total |
|-------------|---------|------------|-----------|-----------|
| Cystic | 61 (65) | 5 (5.4) | 3 (3.2) | 71 (75.5) |
| Mixed | 16 (17) | 0 | 4 (4.3) | 20 (21.3) |
| Solid | 2 (2) | 0 | 1 (1) | 3 (3.2) |
| Total | 79 (84) | 5 (5.4) | 8 (8.5) | 94 (100) |

Among the different histological patterns, surface epithelial tumors (SETs) constituted the majority (89.2%; 83), followed by Germ cell tumors (8; 8.5%) and sex cord-stromal tumors (3; 3.2%) (Table 5). Among the individual tumors, serous cystadenoma (57.45%) were the commonest benign epithelial tumor, followed by mucinous cystadenoma (15.96%) (Table 5).

Table 5: Histopathological spectrum of OTsaccording to the WHO classification.

| Histopathological | Nature of tumour | Types | Number | % |
|--------------------|----------------------|------------|--------|-------|
| diagnosis | | | | |
| Surface epithelial | Serous | Benign | 54 | 57.45 |
| tumours | | Borderline | 2 | 2.13 |
| | | Malignant | 6 | 6.38 |
| | Mucinous | Benign | 15 | 15.96 |
| | | Borderline | 3 | 3.19 |
| | | Malignant | 1 | 1.06 |
| | Endometrioid | Malignant | 1 | 1.06 |
| | Clear cell tumor | Malignant | 1 | 1.06 |
| Sex -cord-stromal | Granulose theca cell | | 1 | 1.06 |
| tumour | tumour | | | |
| | Fibroma thecomas | | 2 | 2.13 |
| Germ cell tumor | Mature teratoma | | 8 | 8.51 |
| Total | | | 94 | 100 |

Discussion

OT is the most lethal of all the gynecologic cancers and gained importance in recent years because of the increasing pelvic examinations, sonological screening and measurement of biomarkers like CA125 in cases with symptoms concerning ovarian lesions. [7].In the present study, 94 ovarian tumors were recorded. Pathological findings of these were Analyzed and correlated with different studies. Our observation showed that 84 cases were benign, and ten were malignant. This is almost similar to the available reports in the literature that reported more common benign lesions than malignant lesions.[5, 8, 9].

Age was ranged between 18 to 75 years. The maximum number of cases were benign, reported in the reproductive age group, 20 - 40yrs. Similar findings were reported by Deepti et al. [10].BoTs were reported in the 31 – 70yrs age group and MITs in> 40yrs.

A higher median age of 60 – 65yrs for MITswas reported from western countries and India. [11].The indication towards an earlier presentation of the malignant lesion in our study compared to western countries mandates a thorough investigation of any vague abdominal complaint in this age group. The present study emphasizes the need for proper histopathological evaluation of OTs at all ages due to relative predominance at reproductive age to rule out malignancy.Screening should start at an early age to detect ovarian malignancy so that the early stage and lower grade can be diagnosed, which may help for the improved survival of women.

OTs were unilateral in 91.25% of cases and bilateral in 8.5%. Among all, mostBeTs and most of the MITs were also unilateral. In this research, right side OTs (52; 55.31%)were common than left-sided. These findings were correlated with Manoja et al. [12], where most of the bilateral tumors (8) were benign. Chandanwale SS et al. Mondal SK et al.reported that most malignant lesions are bilateral.[13, 14].The most common bilateral tumor was Seromucinous cystadenoma, one case of mucinous cystadenoma. It was suggested that bilaterality of a mucinous tumour should always suggest the possibility of a metastatic tumor to the ovaries from the appendix, pancreas, endocervix rather than primary ovarian neoplasms, so thorough investigation of such neoplasms to be done.

The external surface of the benign serous tumors was smooth as the majority of them are unilocular, nodular in mucinous as they are multilocular. The capsule was intact in all BeTs, microscopy of the capsule of all the borderline tumors was reassessed for microscopic invasion. At the same time, the malignant lesions grossly showed a variegated appearance with haemorrhage And necrosis. This was also observed in the studies done by Modepalli et al. [15].

Most of the OTs (71; 75.5%) were grossly cystic cases, followed by mixed (20; 21.27%) and solid tumors (3: 3.2%). In the benign group, the majority were cystic (61), followed by mixed (16) and solid (2). All the borderline (5) were cystic, and out of 8 MITs, four mixed, three were cystic, and one was solid (Table 4).Saha et al. reported that 70% were SET, 26% were germ cell, and 4% were sex cord-stromal tumors. [5]. Sudha et al. reported that 64% were SET, 26% were germ cell and 8% were sex cord-stromal turoms. [4].

The SETs in the present study comprised 70.25% of all tumors. Among the individual tumors, the epithelial tumors the commonest was serous cystadenoma (68.05%) followed by mucinous cystadenoma(20.21%), one clear cell carcinoma, one endometrioid carcinoma(Table5). This was in contrast to Deepti et al., where mucinous cystadenoma was reported to be the commonest. [10].Among primary malignant tumors, Serous cystadenocarcinoma was the commonest in this study, accounting for 6.3%. Whereas Deepti et al. reported this to be 6%. [10]. Germ cell tumors(8.4%)were the second major group of tumors in the present study; these were seen between 20 - 40yrs. In the present study, the incidence of germ cell tumors was relatively less than in other studies.Madhumita et al. reported 17.46 % of germ cell tumors. [16]. Whereas it was said to be 21% by Ranjana et al. [17]. In this, 1.7% were immature teratoma and dysgerminoma each. These findings may contribute significantly to the understanding of the distribution of different ovarian neoplasms among the local population, which may lead to the development of some strategic planning to investigate and treat the underlying causes of concerning neoplasms and may suggest preventive strategies.

Sex cord-stromal tumors constituted 3.19 % in this report. This was similar to the study reported byBuelaprescillaet al., Madhumita et al.,4.6% and 7.93 %, respectively. [16,17].Similar to this research, Ranjana hawaldaret al. [18].Also didn't report Sertoli leydig cell tumor. Ancillary techniques like immunohistochemistry are rarely used in the diagnosis of OTs. These can differentiate the primary ovarian mucinous tumors from malignant metastatic colorectal carcinomas.[15].

Conclusion

The maximum number of tumor cases were benign, reported in the reproductive age group,whereas the malignant neoplasms in > 40 yrs.The present study emphasizes the need for proper histopathological evaluation and screening at all ages due to the relative predominance of OTs to rule out malignancies.Surface epithelial tumors and Serouscystadenocarcinoma was the commonest primary malignant tumor.

Limitation:Genetic testing was not done in our study due to financial constraints. This is the major limitation of this research.

What this study adds to the existing knowledge?

The incidence of malignant tumors is less when compared to the benign and sexually active age is commonly affected.

Reference

01. Pachori, Geeta, et al. "Histopathological study of ovarian tumors in Ajmer region. " International Journal of Medical Science and Public Health 5. 7 (2016): 1400-1404. [Crossref][PubMed][Google Scholar]

02. Mankar, Deepti Vijay, and Gaurav K. Jain. "Histopathological profile of ovarian tumours: A twelve year institutional experience. " Muller J Med Sci Res 6. 2 (2015): 107-11 [Crossref][PubMed] [Google Scholar]

03. Prat J, Mutch DG. Pathology of cancers of the female genital tract including molecular pathology. Int J Gynaecol Obstet. 2018 Oct;143 Suppl 2:93-108. *doi:* 10.1002/ijgo.12617 [Crossref][PubMed] [Google Scholar]

04. Sudha V, Volga Harikrishnan, Sridevi. M, Padma Priya. Clinico pathological correlation of ovarian tumors in a tertiary care hospital. Ind J Of Path. *Onc.* 2018; 5(2): 332 – 7. [Article][Crossref] [PubMed][Google Scholar]

05. Manasi, Saha, Banerjee Alpana, and Datta Abhijit. "Histological patterns of Ovarian neoplasms-A five year experience in North-East India. " International Journal of Medical and Dental Sciences 7. 1 (2018): 1576-1581. [Crossref][PubMed] [Google Scholar] 06. Patel, Amita S., Jignasha M. Patel, and Kamlesh J. Shah. "Ovarian tumors-Incidence and histopathological spectrum in tertiary care center, Valsad." IAIM 5.2 (2018): 84-93 [Crossref] [PubMed][Google Scholar]

07. Doubeni CA, Doubeni AR, Myers AE. Diagnosis and Management of Ovarian Cancer. Am Fam Physician. 2016 Jun 1;93(11):937-44. [Crossref] [PubMed][Google Scholar]

08. . . Am Fam Physician. 2016 Jun 1;93(11):937-44. [Crossref][PubMed][Google Scholar] [Crossref][PubMed][Google Scholar]

09. Gurdip Kaur, Parneet Kaur, Ramesh Kumar Kundal, Jasmine Kaur. A clinicopathological study of ovarian tumors. Int J Of Cur Med Pharm Res. 2017; 3 (9): 2309 – 11. DOI: [Article][Crossref][PubMed] [Google Scholar]

10. Tamrakar, S. R., Makaju, R., Shrestha, A., & Kayastha, S. (2018). Comprehensive study of ovarian tumours in Kathmandu University Hospital. Journal of Kathmandu Medical College, 7(4), 173–179. [Article][Crossref][PubMed][Google Scholar]

11. Mankar, Deepti Vijay, and Gaurav K. Jain. "Histopathological profile of ovarian tumours: A twelve year institutional experience. " Muller J Med Sci Res 6. 2 (2015): 107-11 [Crossref][PubMed] [Google Scholar]

12. Kuladeepa, A. V. K. , et al. "Histomorphological study of 134 primary ovarian tumors." Adv Lab Med Int 1.4 (2011): 69-82 [Crossref][PubMed][Google Scholar]

13. Manoja, Vaddadi, et al. "Clinicopathological study of ovarian tumors: a 2-year study. " International Journal of Scientific Study 5. 3 (2017): 297-302. [Crossref][PubMed][Google Scholar]

14. Chandanwale, Shirish S. , et al. "Clinicopathologic study of malignant ovarian tumors: A study of fifty cases. " Medical Journal of Dr. DY Patil University 10.5 (2017): 430 [Crossref] [PubMed][Google Scholar]

15. Mondal SK, Banyopadhyay R, Nag DR, Roychowdhury S, Mondal PK, Sinha SK. Histologic pattern, bilaterality and clinical evaluation of 957 ovarian neoplasms: a 10-year study in a tertiary hospital of eastern India. J Cancer Res Ther. 2011 Oct-Dec;7(4):433-7. *doi:* 10.4103/0973-1482.92011 [Crossref][PubMed][Google Scholar] 16. Modepalli N, Venugopal SB. ClinicopathologicalStudy of Surface Epithelial Tumours of the Ovary:An Institutional Study. J Clin Diagn Res. 2016Oct;10(10):EC01-EC04.doi:10.7860/JCDR/2016/21741.8716[PubMed][Google Scholar]

17. Madhumita D. K. Neeraja, Gopal A. Histomorphological spectrum of ovarian neoplasm in tertiary care centre. *Int J of Recent Scientific Res.* 2020; 11 (05): 38498 – 502. DOI: 10.17511/JOPM.2020.I03.06 [Crossref][PubMed] [Google Scholar]

18. Beulah Priscilla M, KusarajuPyla, Sindhura Manda, Satyanarayanarao P, Vijayabhaskar R. Histopathological spectrum of ovarian tumors-A three year retrospective study. IOSR J of Den and Med Sci. 2019; 3 (2): 24 – 31. [Crossref][PubMed] [Google Scholar]

19. Hawaldar, Ranjana, Sadhna Sodani, and Ekta Patidar. "Histopathological spectrum of ovarian tumours-A two year retrospective study. " Indian Journal of Pathology and Oncology 4. 3 (2017): 450-453. [Crossref][PubMed][Google Scholar]