Histopathological study of Non-neoplastic & Neoplastic ovarian lesions in a tertiary care hospital in Gujarat, India

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Abstract

Aims & Objectives: (1) To know about various histopathological types of ovarian lesions presented and diagnosed at our institute. (2) To study the incidence of ovarian lesions with respect to patient's age. (3) To study the frequency of ovarian lesions in terms of non-neoplastic or neoplastic, benign or malignant, unilateral or bilateral, etc. Materials & Methods: The present study was performed at the Department of Pathology, GMERS Medical College & Hospital-Junagadh (Gujarat, India) from January 2015 to December 2018 and includes 100 cases of ovarian lesions diagnosed on both clinical & histopathological basis. We have received ovarian specimens and performed routine grossing and H& E staining procedure. We have included parameters like Age wise incidence, Nature of Lesion, Frequency & Laterality in this present study. Results: Out of100 cases, 89% are unilateral and 11% are bilateral. 52% lesions are Benign Neoplasms, 44% lesions are Non-neoplastic Cysts and 4% lesions are Borderline & Malignant Neoplasms. Majority of cases (58%) belong to age group of 20-39 years. Among Non-neoplastic Lesions, Follicular Cyst is common & frequently bilateral while among Benign Neoplasms, Serous Cystadenoma is common & frequently bilateral. Conclusion: Ovarian Lesions both non-neoplastic and neoplastic include a variety of morphological features and show a particular age wise incidence. Role of histopathological evaluation remains always important in both diagnosis & management of such cases, particularly in cases of Malignant Lesions in order to save the patient's life.

Key words: Non-neoplastic, Benign, Borderline, Malignant, Follicular Cyst, Serous Cystadenoma

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Introduction

Ovarian enlargements (lesions), cystic or solid, may occur at any age [1]. These enlargements may be Nonneoplastic or Neoplastic in nature. Non-Neoplastic Enlargements include Simple Follicular Cysts, Corpus Leuteal Cysts, Chocolate Cysts due to Endometriosis, Twisted Hemorrhagic Cysts, Polycystic Ovarian Disease (PCOD), Various Inflammatory Lesions, etc. To define a functional non-neoplastic cyst, its size or diameter must be at least 3 cm, but not more than 7 cm [1]. Non-Neoplastic Enlargements develop almost exclusively during the childbearing years.

They may be asymptomatic or produce local discomfort, menstrual disturbances, infertility, or in rare cases cause acute symptoms due to complications like haemorrhage, rupture or torsion [1]. Neoplastic disorders or lesionsof Ovary can arise from (1) Mullerian epithelium, (2) Germ cells or (3) Sex cord

Manuscript received: 20th January 2019 Reviewed: 30th January 2019 Author Corrected: 7th February 2019 Accepted for Publication: 11th February 2019 Stromal cells [2]. Classification of Ovarian Neoplasms given by WHO is completely based upon the tissue of origin. It includes a variety of entities like Surface Epithelial Tumors, Sex cord Stromal Tumors, Pure or Mixed Germ Cell Tumors, Gonadoblastoma, Soft Tissue Tumors, Metastatic Tumors, Unclassified Tumors, etc. Surface Epithelial Tumors are further categorized into benign, borderline and malignant [1,3]. Ovarian neoplasms behave in diverse way and generally escape the detection until they attain a larger size. This is primarily due to the reason that either the symptoms are vague or most of these are asymptomatic therefore they manifest over a time period due to no definite screening program. Therefore, diagnosis of various histological patterns of ovarian tumors is very important in the treatment and prognosis. [2]

Aims & Objectives

• To know about various histopathological types of ovarian lesions presented and diagnosed at our institute.

- To study the incidence of ovarian lesions with respect to patient's age
- To study the frequency of ovarian lesions in terms of non-neoplastic or neoplastic, benign or malignant, unilateral or bilateral, etc.

Material & Method

The present retrospective studywas performed at the Department of Pathology, GMERS Medical College& Hospital-Junagadh (Gujarat, India) from January 2015 to December 2018 for a period of 4 years. It includes a total number of 100 random cases of ovarian lesions diagnosed initially on the basis of clinical findings and finally on the basis of histopathological examination.

Nature of Specimens

- Solitary Salpingooophorectomy specimens, Unilateral or Bilateral
- Hysterectomy with Unilateral Salpingooophorectomy specimens
- Hysterectomy with Bilateral Salpingooophorectomy specimens

Original Research Article Inclusion Criteria: All ovarian biopsy with abnormal findings on sonography.

Exclusion Criteria:

History of previous chemotherapy or radiotherapy
Age below 20 years.

These specimens have been received from Gynecology Department with sufficient & relevant clinical and radiological findings in a specific Performa for further Histopathological Evaluation at our Department. After a number of necessary technical steps like Fixation in 10% Formalin, Gross Examination, Surgical Dissection, Tissue Processing, Paraffin Embedding, Microtomy, staining with routine H-E method & Mounting of Slides using DPX, Microscopic Slides have been made available for further evaluation & final diagnosis.All slides have been viewed by at least two Pathologists before final reporting and all lesions have been classified using WHO guidelines. Data are plotted and statistical analysis was carried out with the help of SPSS (Statistical packages for social science) and by using X²- chi square test. P value less than 0.001 is significant.

Results

A total of 100 specimens of ovarian lesions have been received for histopathological evaluation at our department & out of them, 89 lesions are unilateral and 11 lesions are bilateral. 52% lesions are Benign Neoplasms, 44% lesions are Non-neoplastic Cysts and 4% lesions are Borderline & Malignant Neoplasms. Histopathological Categorization of Various Ovarian Lesions is given below in Table no.1 and their Age-wise incidence is given below in Table no.2.

Sr.No.	HistopathologicalDiagnosis or Category	Number of	Unilateral	Bilateral
		Cases and %	Cases	Cases
Non-ne	oplastic lesions			L
1	Follicular Cysts	20	17	03
2	Corpus Leuteal Cysts	14	11	03
3	Hemorrhagic Twisted Ovarian Cysts	06	06	
4	Endometriosis & Chocolate Cysts	04	04	
Benign	neoplasms		•	
5	Serous Cystadenoma	28	24	04
6	Mature Cystic Teratoma	13	13	
7	Mucinous Cystadenoma	06	05	01
8	Serous Cystadenofibroma	03	03	
9	Mature Teratoma with Struma Ovarii	01	01	
10	Brenner's Tumor (Benign)	01	01	
Border	line & malignant neoplasms			
11	Borderline Mucinous Tumor	01	01	
12	Papillary Serous Cystadenocarcinoma	01	01	
13	Granulosa Cell Tumor of Adult Type	01	01	
14	Dysgerminoma	01	01	
	Total	100	89	11

Table No-1: Histopathological Categorization of various Ovarian Lesions

Sr. No.	Histopathological Diagnosis or Category	20-39 years	40-59 years	60 years & above	Total
1	Follicular Cysts	14	06		20
2	Corpus Leuteal Cysts	08	06		14
3	Hemorrhagic Twisted Ovarian Cysts	06			06
4	Endometriosis & Chocolate Cysts		03	01	04
5	Serous Cystadenoma	15	08	05	28
6	Mature Cystic Teratoma	07	05	01	13
7	Mucinous Cystadenoma	04	01	01	06
8	Serous Cystadenofibroma	03			03
9	Mature Teratoma with Struma Ovarii		01		01
10	Brenner's Tumor (Benign)		01		01
11	Borderline Mucinous Tumor		01		01
12	Papillary Serous Cystadenocarcinoma			01	01
13	Granulosa Cell Tumor of Adult Type		01		01
14	Dysgerminoma	01			01
	Total	58	32	10	100

TableNo.-2: Age-wise Incidence of various Ovarian Lesions.

As mentioned above in both tables, Simple Follicular Cysts are the commonest among all Non-neoplastic Cystic Lesions and often bilateral. Non-neoplastic Cysts are seen in comparatively younger females belonging to age group of 20-39 years and quite rare after menopause because of complete cessation of normal & regular menstrual cycles.

Among various benign neoplasms, Serous Cystadenoma is the commonest & often bilateral. Next common benign neoplasm is Mature Cystic Teratoma or Dermoid Cyst of Ovary that is often unilateral. Both of these benign tumors can occur at any age but commonly seen before 40 years of age during reproductive period of life.

Other benign neoplasms like Mucinous Cystadenoma and Serous Cystadenofibroma are also seen in younger females. Brenner's Tumor is an uncommon subtype of the surface epithelial tumors of ovary and Majority of the cases are benign in nature.

This tumor is particularly seen between 40-60 years of age. Our present study includes total 4 cases of Borderline & Malignant Neoplasms namely Serous Cystadenocarcinoma (65 years old), Granulosa Cell Tumor of Adult Type (45 years old), Dysgerminoma (32 years old) and Borderline Mucinous Tumor (48 years old).



Fig 1a shows Struma Ovarii with extensive normal looking thyroid tissue. 1b shows Benign Brenner's Tumor with characteristic transitional or urothelial component giving a biphasic appearance.



Fig 2a shows Borderline Mucinous Tumor of ovary, a tumor with nuclear atypia and having low malignant potential (in-between benign & malignant). 2b shows Papillary Serous Cystadenocarcinoma of ovary with features of anaplasia and aggressiveness.



Fig no. 3a shows Granulosa Cell Tumor of Adult Type with characteristic Coffee-bean Nuclei. 3b shows Dysgerminoma of Ovary looking more or less similar to the classical seminoma of the testis having clear looking cells.

Figure no. 1a, 1b, 2a & 2b shows microscopic photographic images of various lesions under low power magnification (H & E stain, 100x) while figure no. 3a & 3b shows images under high power magnification (H & E stain, 400x).

Discussion

Our present study includes total 100 cases of Ovarian Lesions diagnosed finally on histopathological basis. Out of them, 89% cases show unilateral lesion and 11% cases show bilateral lesions. Prakash et al found similar observation in his study with 90.9% unilateral & 9.2% bilateral lesions [4], Gurung P et al [5] with 88.15% unilateral & 11.85% bilateral lesions, Prabhakar BR& Kalyani M [6]with 90.9% unilateral & 9.1% bilateral lesions and by Couto F et al[7]having 91.2% unilateral ovarian lesions & 8.7% bilateral lesions. So we can say that unilateral lesions are more frequent than bilateral ones and all the studies show comparable outcomes [4,5,6,7]. Broad categorization of these lesions includes Non-neoplastic cysts, Benign Neoplasms and Borderline-Malignant Neoplasms. Our present study shows 44% cases of Non-neoplastic Cysts. Similar studies like those done by Prakash A et al [4] and by Gurung P et al [5] shows 44.10% cases and 43.70% cases of Non-neoplastic Cysts respectively. In the study done by Zaman et al[8], 68.87% of the lesions are non– neoplastic in nature that is somewhat higher than other studies. Among all non-neoplastic cysts. Simple Follicular Cyst, unilateral or bilateral is the commonest one followed by Corpus Leuteal Cyst.

Both of them are frequently seen among women of reproductive age and perimenopausal age. Our present study shows 20% of Follicular Cysts & 14% of Corpus Leuteal Cysts respectively. Study done by Prakash A et al [4] shows 20.1% of Follicular Cysts & 10.9% of Corpus Leuteal Cysts respectively. Gurung P et al [5]reported 17.0% of Endometriotic or Chocolate cyst, 10.4% of Simple Follicular Cyst and 9.6% of Corpus Leuteal Cyst and this result or outcome is not comparable with that of our present study. Our present study shows 52% cases of benign neoplasms as well as 4% cases of borderline & malignant neoplasms. Study done by Prakash A et al [4] shows 54.14% cases of benign neoplasms as well as 1.74% cases of borderline & malignant neoplasms. Among benign neoplasms, Serous Cystadenoma is the commonest one followed by Mature Cystic Teratoma (Dermoid Cyst) or Mucinous Cystadenoma. Our present study shows 28% cases of Serous Cystadenoma, 13% cases of Mature Cystic Teratoma and 6% cases of Mucinous Cystadenoma and 19.9% cases of Mature Cystic Teratoma.

Pudasaini S et al[10] has reported 40.2% cases of Serous Cystadenoma, 15.7% cases of Mature Cystic Teratoma and 9.8% cases of Mucinous Cystadenoma. Maliheh et al[11] has reported 38% cases of Serous Cystadenoma, 30% cases of Mature Cystic Teratoma and 22% cases of Mucinous Cystadenoma. Mondal et al [12] has reported 29.9% cases of Serous Cystadenoma, 15.9% cases of Mature Cystic Teratoma and 11.1% cases of mucinous cystadenoma. Iqbal et al [13] has reported 38.5% cases of Serous Cystadenoma followed by 30.8% cases of Mature Cystic Teratoma. Yasmin et al [14] has reported 24% cases of Serous Cystadenoma followed by 18% cases of Mature Cystic Teratoma. Sowe can say that outcome of all these similar studies are comparable with that of our present study.

In our study, majority of the patients (58%) belong to age group of 20-39 years followed by age group of 40-59 years (32%) and 60 years or more (10%). In our study we have not included patients who are below 20 years of age. Comparison of this age distribution with that of other similar studies is mentioned below in table no. 3.

Nome of the Study	Age Groups in Years				
Name of the Study	0-19	20-39	40-59	60 & above	
Ramachandran et al [15]	7.9%	53%	30%	9.1%	
Pilli et al [16]	7%	58%	30%	5%	
Kar et al [17]	7.4%	41.7%	46.2%	4.4%	
Prakash A et al [4]	5.7%	53.4%	36.6%	4.3%	
Present Study		58%	32%	10%	

Table No.-3: Comparison of Age Distribution of patients with ovarian lesions.

So we can say that results of our present study are in concordance with that of other similar studies. Ovarian lesions particularly non-neoplastic cysts & benign neoplasms are commonly seen in age group of 20-39 years of age.

Our present study includes four cases of borderline and malignant lesions. All of them are Unilateral and show different age wise incidences. Borderline Mucinous Tumor is seen at the age of 48 years. Prakash A et al [4] has also reported one case of Papillary Serous Cystadenocarcinoma in 65 years old female patient. Prakash A et al [4] has also reported one similar case with age of 56 years. Gurung P et al [5] has reported five such cases with age range of 38 to 66 years. Our present study includes one case of Dysgerminoma of Ovary with age of 32 years. Modi D et al [2] has reported one case of Dysgerminoma with age less than 20 years. Our present study also includes one case of Adult type Granulosa Cell Tumor who is 45 years old. Modi D et al [2] has reported three such cases out of which, two belongs to age group of 31-40 years and one belongs to age group of 41-50 years. Hatwal D et al [18] has reported one case of Granulosa Cell Tumor of Adult type who is 45 years old. So we can say that age wise incidence of Borderline & Malignant Neoplasms of our present study also correlate with that of other similar studies. Dysgerminoma is a malignant neoplasm of younger females while Borderline tumors and Granulosa cell tumor of Adult type are commonly seen during middle age or perimenopausal age. Serous Cystadenocarcinoma occurs at comparatively older age, usually beyond 60 years.

Conclusion

Ovarian Lesions or Enlargements, both non-neoplastic and neoplastic include a variety of morphological features and show a particular age wise incidence. Among non-neoplastic lesions, simple follicular cyst is common while among benign tumors, serous cystadenoma is common. Both are frequently bilateral. Malignant Neoplasms of Ovary are rare as compared to Benign Neoplasms and Non-neoplastic Lesions, but require a specific attention during diagnosis on both clinical and pathological basis in order to save the patient's life.

Role of histopathological evaluation remains always important in diagnosis and management of such cases

along with clinical and radiological evaluations. Histopathological study is useful to predict nature and course of ovarian lesions so that future worse outcome can be prevented with early intervention and marker study.

Contribution from authors

- Dr. Alpesh Maru: Data collection, analysis and preparation of manuscript.
- Dr. Chirag Menpara: Analysis and preparation of manuscript & critical revision.

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