## Clinical Profile of Neoplastic and Non-Neoplastic Lesions of Ovary: A Prospective Observational Study from Central Indian population

## Somya Saxena<sup>a</sup>, Rinku Bhagora<sup>a</sup>, Priyanka Solanki<sup>b</sup>, Harshul Patidar<sup>a\*</sup>

<sup>a</sup>Department of Pathology, Nandkumar Singh Chouhan Government Medical College, Khandwa-450001, Madhya Pradesh, India.

<sup>b</sup>Department of Pathology, MGM Medical College, Indore, Madhya Pradesh, India.

#### Abstract

**Background:** Most Indian population-based cancer registries have reported a gradual rise in the ovarian cancer incidence over the years. These neoplasms exhibit a spectrum of genetic background, much more varied than any other gynecological condition and present a big challenge to a gynecological oncologist. Therefore, proper recognition and classification of such pelvic masses is important for appropriate therapy and better prognosis.

**Objectives** This study aimed to look at the demographics and clinical profile of various ovarian lesions in the local population of the central India.

**Patients and Methods:** A prospective observational study was carried out on the surgically resected ovarian samples that were referred to the Pathology department over two and half year. A total of 100 ovarian cases were included. Relevant clinical information regarding age, bleeding, pain in abdomen, menstrual history, histopathological examination reports were recorded.

**Results:** Out of 100 cases of ovarian lesions, majority were neoplastic lesions. Most of the cases of non-neoplastic ovarian lesions belonged to 31-40 years' age group, whereas most cases of neoplastic ovarian lesions belonged to 41-50 years' age group. Most common presenting symptom was abnormal uterine bleeding in non-neoplastic cases. But neoplastic cases presented mainly with abdominal pain.

**Conclusion:** Majority of the ovarian lesions in central India population present after second parity, are benign in nature and present with abnormal uterine bleeding, whereas malignant ovarian lesions mainly present with abdominal pain and after 40 years of age.

Keywords: Ovarian Neoplasms; India; Abdomen; Parity; Oncologists.

#### DOI: 10.21608/svuijm.2023.195005.1533

\*Correspondence: harshul.patidar@gmail.com

Received: 25 Januray, 2023.

Revised: 8 February, 2023.

Accepted: 8 March, 2023.

Published: 19 April, 2023

**Cite this article as**: Somya Saxena, Rinku Bhagora, Priyanka Solanki, Harshul Patidar. (2023). Clinical Profile of Neoplastic and Non-Neoplastic Lesions of Ovary: A Prospective Observational Study from Central Indian population. *SVU-International Journal of Medical Sciences*. Vol.6, Issue 2, pp: 198-205.

Copyright: © Saxena et al (2023) Immediate open access to its content on the principle that making research freely available to the public supports a greater global exchange of knowledge. Users have the right to Read, download, copy, distribute, print or share link to the full texts under a Creative Commons BY-NC-SA 4.0 International License

## Introduction

Ovary is an important female reproductive organ. It consists of totipotent gametocytes and multipotent mesenchymal tissue. So almost any type of tumor can arise from ovary (Sikdar et al.,1981). Ovarian carcinoma stands out as the third most prevalent cancer in Indian women and eighth in the world, contributing to 3.44% of all cancer cases (GLOBOCAN, 2008). It is also an important cause of deaths in Indian cancer women, constituting about 3.34% of all cancer deaths in India. If diagnosed early in Stage I, the five-year survival from ovarian cancer good is (about 94%) but unfortunately only 15% of cases are diagnosed in this Stage I. While most (about 62%) of cases are diagnosed in Stages III and IV, when five-year survival is merely 28% (Misra et al., 1991). Ovarian cancers in advanced stage have worst prognosis, having the highest case mortality ratio amongst all gynecological cancers globally (ICMR, 2019).

. Most Indian population-based cancer registries have reported a gradual rise in the ovarian cancer incidence over the years. They have reported that ageadjusted incidence of ovarian cancer varies from 0.9 - 8.4 per 100,000 women (Murthy et al., 2009). There is age related increase in the incidence of ovarian cancer. The ASIR (age specific incidence rate) starts increasing from 35 years of age and peaks between 55-64 years of age. However, many western countries have documented a decreasing trend of incidence and mortality. This could be due to wider increase in preventive measures like use of oral contraceptives, reduction in post-menopausal HRT (hormone replacement therapy) and increased

application of risk-reduction surgeries (ICMR, 2019).

It is a well-known fact that neoplastic conditions of ovaries are a complicated and interesting subject in the history of oncology. These neoplasms inherit a spectrum of genetic background, much more varied than any other gynecological condition (Misra et al., 1991), and present a big challenge to a gynecological oncologist. Even, certain non-neoplastic ovarian lesions frequently present with a pelvic mass and mimic an ovarian tumor. Therefore, proper recognition and classification of such pelvic masses is important for appropriate therapy and better prognosis. This study aimed to look at the demographics and clinical profile of various ovarian lesions in the local population of the central India.

## **Patients and methods**

This research study was conducted after getting approval from the institutional ethics committee of L.N. Medical College & Research Centre, Kolar Road, Bhopal, Madhya Pradesh. A prospective observational study was carried out on the surgically resected ovarian samples that were referred to the Pathology department of L.N. Medical College & Research Centre over two and half year starting from May 2019.

All resected samples with diagnosis of ovarian lesion of all age groups were included, while autolyzed specimens were excluded from study. A non-probability convenient sampling technique was used and a total of 100 ovarian cases were included. Relevant clinical information regarding age, bleeding, pain in abdomen, menstrual history, histopathological examination reports were recorded.

#### Statistical analysis

Data analysis was performed with the help of Microsoft Excel 20087 and SPSS version 20 software. Frequency distribution and cross tabulation were used to make tables. All the qualitative data were expressed as number and percentage. p value < 0.05 was taken to be statistically significant.

#### Results

Out of 100 cases of ovarian lesions studied, majority (n=64) were neoplastic lesions. Amongst the neoplastic lesions majority (n=58) were found to be benign in nature (**Table. 1**).

Lesion Type		on Type Number of cases				
Non-Neopla	astic	36	36			
N l	Benign	58	58			
Neoplastic	Malignant	6	6			
Total		100	100			

Table 1.Distribution of ovarian cases

Most (44%) of the cases of nonneoplastic ovarian lesions belonged to 31-40 years' age group (**Table. 2**), whereas most (36%) of the cases of neoplastic ovarian lesions belonged to 41-50 years' age group (**Table. 3**).

Age	Number of cases	%
< 30	3	8.33
31-40	16	44.44
41-50	13	36.11
51-60	3	8.33
> 60	1	2.77
Total	36	100

#### Table 3. Age wise distribution of Neoplastic ovarian cases

A go	Benign		Mali	gnant	Total	
Age	Ν	%	Ν	%	Ν	%
< 20	2	3.45	1	16.67	3	4.69
21-30	13	22.41	2	33.33	15	23.44
31-40	17	29.31	1	16.67	18	28.13
41-50	22	37.93	1	16.67	23	35.94
51-60	3	5.17	0	0	3	4.69
> 60	1	1.72	1	16.67	2	3.13
Total	58	100	6	100	64	100

Majority of the non-neoplastic as well as neoplastic cases presented with second parity (**Table. 4** and **Table.5**).

#### Table 4. Parity distribution of Non-Neoplastic ovarian cases

Parity	Ν	%
Nulliparous	0	0
P1L1	3	8.33
P2L2	20	55.56
P3L3	12	33.33
P4L4	1	2.78
Total	36	100

Parity	Benign		Mali	gnant	Total		
	Ν	%	Ν	%	Ν	%	
Nulliparous	7	12.07	2	33.33	9	14.06	
P1L1	6	10.34	1	16.67	7	10.94	
P2L2	26	44.83	2	33.33	28	43.75	
P3L3	10	17.24	1	16.67	11	17.19	
P4L4	6	10.34	0	0	6	9.38	
Unmarried	3	5.17	0	0	3	4.69	
Total	58	100	6	100	64	100	

Most common presenting symptom was abnormal uterine bleeding (42%) in non-neoplastic cases followed by abdominal pain (22%). But neoplastic cases presented mainly with abdominal pain (36%), and lump in abdomen being the second most common (27%) presenting symptom (Table. 6 and Table.
7).

Table 6. Clinical	presentation	of Non-Neo	plastic	ovarian	cases
Table 0. Chincar	presentation	01 11011-1100	plastic	0 v al 1 all	cases

Clinical presentation	Ν	%
Asymptomatic	7	19.44
Abdominal mass	3	8.33
Abdominal pain	8	22.22
Abnormal Uterine Bleeding	15	41.67
Abdominal mass with adenomyosis	2	5.56
Abdominal mass with fibroid	1	2.78
Total	36	100

 Table 7. Clinical presentation of Neoplastic ovarian cases

Clinical presentation		Benign		Malignant		Total	
		%	Ν	%	Ν	%	
Abdominal mass (LA)	13	22.41	4	66.67	17	26.56	
Abdominal pain (AP)	22	37.93	1	16.67	23	35.94	
Abnormal Uterine Bleeding	15	25.86	0	0	15	23.44	
Ascites	4	6.90	0	0	4	6.25	
Asymptomatic	1	1.72	0	0	1	1.56	
LA, PA & Urinary complaints	0	0	1	16.67	1	1.56	
LA, PA with Ascites	3	5.17	0	0	3	4.69	
Total	58	100	6	100	64	100	

#### Discussion

Ovarian neoplasms are a challenging problem in recent days' gynecology, mainly due to variable, and many a times undifferentiated, pathologic subtypes. Its mortality rate exceeds the combined mortality of both endometrium and cervical malignancy. We studied the clinical spectrum and associated findings that is of huge clinical significance for pathologists, radiologists, and gynecologists for a good knowledge and better prognosis of the disease and

planning proper management	(Gadducci et al.,2019).						
Name of Authors	Non-	Neoplastic					
Name of Authors	neoplastic	Benign	Malignant				
Kanthikar (2014)	51.72% (n=75)	42.75% (n=62)	5.51% (n=8)				
Tejani et al. (2020)	32.1% (n=122) 60% (n=24)		7.9% (n=18)				
Priya et al. (2017).		78% (n=97)	21% (n=16)				
Prakash et al. (2017)	44% (n=110)	45.8% (n=100)	9.2% (n=19)				
Laul et al. (2020)	27% (n=27)	60.8% (n=59)	11.2% (n=11)				
Gaikwad et al. (2020)	54.6% (n=101)	45.4 % (n = 84)					
Present study	36% (n=36)	58% (n=58)	6% (n=6)				

The above table shows the different types of the study performed all over India different parts and expresses the incidence of ovarian neoplasm in all the areas. In our study, we found out, in a total of 100 cases the non-neoplastic cases (36%) are less than the total of neoplastic cases 64 (64%) including both benign and malignant cases. Similar, the finding was seen with **Laul et al. (2020)** with 27% of nonneoplastic cases and a total of 73% cases were of neoplastic origin. similar distribution was seen in the study of **Prakash et al. (2017)** and **Kanthikar** (2014)

Ovarian tumors are common in all age groups and no age is excluded. The age range in the present study was 9 to 65 years. The maximum number of cases included in our study were in the age group of 41-50 years.

Age in	Tejani et al.	Prakash et	Pradhan et	Mondal	Present
years	(2020)	al. (2017)	al. (2018)	et al.	study
				(2011)	( <b>n=100</b> )
<20	3.16%	5.7.%	8.3%	6.8%	4.69%
21-30	21.84%	6.2%	18.3%	30.04%	23.44%
31-40	30.26%	47.2%	25%	27.6%	28.13%
41-50	33.95%	20.12%	29.1%	22.6%	35.39%
51-60	8.42%	15.88%	15%	10.1%	4.69%
>60	2.37%	4%	4.1%	2.1%	3.13%

The above table shows comparison of various ovarian tumors according to age with other studies. The most common age group in the present study was 41-50 (35.39%) followed by 31-40 years (30.26%) with the mean age in our study being 36.43 years. This highlights that ovarian neoplasms were more prevalent in those living in the second to fifth decade of their life. Similar study done by **Tejani** 

et al. (2020), Pradhan et al. (2018) reveal that the maximum age group of presentation was 41-50 year. However, discordant results were reported by Prakash et al. (2017), which revealed ovarian neoplasms in 31-40 years (47.2%) of age because maximum cases of germ cell tumor reported commonly in middle age group. Increasing parity is associated with a reduction in the risk of ovarian cancer, but it is not clear whether this association applies to all the histopathological types and borderline tumors. nulliparity and unmarried females were associated with the increased relative risk of ovarian tumors (**Priya et al.,2017**).

Parity	Kanthikar (2014), (n=70)	Pradhan et al. (2018), (n=230)	Present Study (n=100)
Nulliparous	20% (n=14)	14.16% (n=34)	14.06% (n=9)
P1L1	8.57% (n=6)	17.50% (n=42)	13.94% (n=7)
P2L2	27.14% (n=19)	27.50 % (n=66)	43.75% (n=28)
P3L3	17.14% (n=12)	20 % (n=48)	17.19% (n=11)
P4L4	20% (n=14)	13.6 % (n=32)	9.38 % (n=6)
Unmarried	7.14% (n=5)	3.46 % (n=8)	8.69 % (n=3)

The above table shows comparison of various studies showing parity status in various neoplastic tumor. In our study it is found that ovarian neoplastic lesion are more in number with women having two issues or parity status two 43.75%, followed by cases of women having gravida three 17.09% (n=11). However, while considering the malignant lesion, status of parity is not very clear because of paucity of cases. The status of parity in our malignancy cases study for were maximum in women having gravida two which include two cases 33.3%, one case was presented who as nulliparus. Similar results were seen in study of Kanthikar (2014) and Pradhan et al. (2018) that women having issue two and three are minimally having the risk of development of ovarian neoplasm.

Some of the ovarian tumors may be incidentally diagnosed on ultrasound

whereas others may be symptomatic. The present study reveals that the presentation of ovarian tumors is variable. If a patient presented with more than one complaint, then the predominant symptom was considered as the presenting symptom. In the present study, the most common clinical presentation among the patients with benign lesions was abdominal pain 22(37.93%) followed by abnormal uterine bleeding 15(25.86%) and abdominal mass 13(22.41%). There were 4 (6.90%) patients presented with benign lesions who had ascites and two case 5.17% had a abdominal mass with pain in the abdomen with ascites. Only one patient was asymptomatic. Out of 6 patients with malignant lesions, mostly patient present with abdominal mass (66.67%) followed by 16.67% abdominal pain and lump, rest presented with pain with urinary complaints.

Kanthikar (2014)	Tejani et al. (2020)	Priya et al. (2017)	Present Study
10.67%	6%	12%	1.56%
10.67%	13%	11%	26.56%
	Kanthikar           (2014)           10.67%           10.67%           29.33%	Kanthikar (2014)Tejani et al. (2020)10.67%6%10.67%13%29.33%27%	Kanthikar (2014)Tejani et al. (2020)Priya et al. (2017)10.67%6%12%10.67%13%11%29.33%27%57%



Abnormal				
Uterine	36%	18%	22%	23.44%
Bleeding				
Ascites	8%	3%	3%	6.25%
Abdominal				
mass with other	16.6%	9%	8%	6.25%
complaints				

The above table shows comparison of mode of presentation of Neoplastic lesions of ovary with other studies. However, Kanthikar (2014) study report of 70 cases of neoplastic neoplasm, the most common presentation was abnormal uterine bleeding or vaginal bleeding with 36% cases followed by pain in the abdomen 29.33% of cases the total of 7.1 % of malignant cases are presented with a lump in the abdomen this discordance seen because majority of patient presented with complex feature of cyst. Tejani et al. (2020) stated that out of 301 cases maximum number of women presented with Pain in the abdomen around 27% of cases followed by 18% of women presented with abnormal uterine bleeding with asymptomatic women were 18 cases in number (6%) (Tejani et al. ,2020). Margaret H Priya et al80 in this study most of the patients with ovarian mass, almost 57% presented with abdominal pain and 12% were asymptomatic. Abdomen pain, bleeding per vaginum on and off were also the most common symptoms Priya et al. (2017).

# Conclusions

Clinical presentation of ovarian lesions varies in central India population with majority presenting after second parity. They are benign in nature and present with abnormal uterine bleeding, whereas malignant ovarian lesions mainly present with abdominal pain and after 40 years of age.

**Conflicts of interest:** None reported **Sources of funding:** Nil **References** 

- Sikdar K, Kumar P, Roychowdhary NN. (1981). A study of ovarian malignancy: A review of 149 cases. J Obstet Gynaecol India,30:478–80.
- GLOBOCAN (2008). Estimated cancer Incidence, Mortality, Prevalence and Disability-adjusted life years (DALYs) Worldwide in 2008. http://globocan.iarc.fr/.
- Misra RK, Sharma SP, Gupta U, Gaur R, Mishra SD. (1991). Pattern of ovarian neoplasm in eastern UP. J Obstet Gynecol India, 30:242–46.
- ICMR. (2019). Consensus Document for Management of Epithelial Ovarian CANCER. https://main.icmr.nic.in/sites/default/files

/guidelines/Ovarian\_Cancer.pdf.

- Murthy NS, Shalini S, Suman G, Pruthvish S, Mathew A. (2009). Changing trends in incidence of ovarian cancer - the Indian scenario. Asian Pac J Cancer Prev,10(6):1025-30.
- Gadducci A, Guarneri V, Peccatori FA, Ronzino G, Scandurra G, Zamagni C, et al. (2019). Current strategies for the targeted treatment of high-grade serous epithelial ovarian cancer and relevance of BRCA

mutational status. J Ovarian Res, 28;12(1):9.

- Kanthilkar A M N. (2014). Clinicohistopathological analysis of neoplastic and non-neoplastic lesions of the ovary: a 3-year prospective study in dhule, north maharashtra, India. J Clin Diagn Res. 2014;8(8):FC04-FC7. doi:10.7860/JCDR/2014/8911.4709.
- Tejani AS, He L, Zheng W, Kanupriya k. (2020). Concurrent, Bilateral Presentation of Immature and Mature Ovarian Teratomas with Refractory Hyponatremia: A Case Report. J Clin Imaging Sci. 2020;10:23.
- Priya MHF, Vanusha, Kirubamani NH. (2017). Clinical correlation of ovarian mass with ultrasound findings and histopathology report. Int J Reprod Contracept Obstet Gynecol ,6:5230-4.
- Prakash A, Akiana S, Duraiswami R, Indira.V. (2017). Histopathological study of ovarian lesions in a tertiary care center in Hyderabad, India-a retrospective fiveyear study. Int J Adv Med ,4:745-9.
- Laul P , Miglani U, Srivstava A , Sood N , Miglani S. (2020). Correlation of clinical, biochemical and radiological characteristics with histopathology of ovarian masses: hospital based descriptive study. International Journal of Reproduction, Contraception, Obstetrics and Gynecology, 9(11):4449-4454.
- Gaikwad SL, Badlani KS, Birare SD. (2020). Histopathological study of ovarian lesions at a tertiary rural hospital. Trop J Pathol Microbiol. 2020;6(3):245-252.
- Pradhan HK, Singh P , Ravikumar MS , Gothwal M. (2018). Study of

risk factors and tumor markers in ovarian malignancy in western part of Odisha: a prospective observational study. International Journal of Reproduction, Contraception, Obstetrics and Gynecology ,7(4):1571-1578.

• Mondal SK, Banyopadhyay R, Nag DR, Roychowdhury S, Mondal PK, Sinha SK. (2011). 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,7(4):433-437.