OVARIAN NEOPLASMS—A STUDY OF 903 CASES

by

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Ovary is one of the common sites for neoplasms in the female. Ovarian cancer is now the leading cause of death from gynaecological malignancy and ovarian tumours comprise about 1% of all new growths in paediatric gynaecology. (Barber 1969). Being an organ with a complex structure, attempts at classification of tumours of the ovary, both by pathologists and gynaecologists. have not yielded satisfactory results so far. This paper presents a study of 903 ovarian neoplasms recorded in the department of Pathology, Medical College, Trivandrum, during a 13 year period. Non-neoplastic cysts of the ovary were excluded.

Analysis

Out of 903 ovarian neoplasms, 622 (68.98%) were benign and 281 (31.02%) were malignant. Among the benign neoplasms there were 59 cases diagnosed as haemorrhagic cysts where the primary nature of the cyst could not be made out. Table I to III gives the histological types of tumours and their age incidence.

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The common presenting complaint in these cases was mass in the abdomen. In many it was associated with pain. Dysuria and menstrual irregularities were the additional symptoms. It is interesting to note that 6 out of 25 cases of granulosa cell tumour occurred in postmenopausal women and the presenting symptom was bleeding.

Ascites was noted in 3 cases of fibroma and in one case of anaplastic carcinoma. Pleural effusion with ascites was present in one case of granulosa cell tumour and one case of papillary cystadenocarcinoma. One case of mucinous cystadenocarcinoma was associated with pseudomyxoma peritonii.

The side affected was not known in all the cases. Of 278 cases in which it was known, 128 (46.04%) involved the right ovary, 107 (38.5%) affected the left ovary and in 43 cases (15-46%) it was bilateral.

The size of the tumour varied from 3-36 cms. The maximum size was noted in a mucinous cystadenoma and the maximum weight noted was in a malignant teratoma which weighed 7 kg. Mucinous cystadenomas weighed from 0.5 to 6 kg. The maximum weight noted in a granulosa cell tumour was 6 kg., in a serous cystadenoma 4.5 kg., mucinous cystadenocarcinoma 4 kg. and dysgerminoma 1.5 kg. The single case of arrheno-

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TABLE I

Histological Classification of 903 Ovarian Neoplasms with their Percentage Incidence

	Histological type				Number of cases	Percentage
ENIGN						
Cystadenomas -	- Mucinous				165	18.27
	Serous				150	16.60
	Papillary serous				31	3.43
Teratomas -	- Benign cystic				124	13.73
	Benign solid				24	2.77
	Struma ovarii				5	0.55
	Struma with tera	toma			1	0.11
	Struma with muci	nous c	vstadeno	ma	1	0.11
	Teratoma with mu				2	0.22
Fibroma					22	2.44
Cystadenofibrom					18	1.99
Adenofibroma					3	0.33
Thecoma					6	0.67
Brenner tumour					8	0.89
	ucinous cystadenoma			- :-	3	0.33
Haemorrhagic c					59	6.54
	Total				622	68.98
	rcinoma—Papillary s	serous			64.	7.09
Mucinous					22	0.44
Papillary ad	lenocarcinoma				44	2.44
- abittary ad					61	6.76
Adenocarcin	oma					
Adenocarcin	oma oma with dermoid				61 14	6.76
Adenocarcine Adenocarcine					61	6.76 1.55
Adenocarcine Adenocarcine	oma with dermoid carcinoma with der				61 14 1 3	6.76 1.55 0.11 0.33
Adenocarcine Adenocarcine Epidermoid	oma with dermoid carcinoma with der arcinoma	moid			61 14 1 3 26	6.76 1.55 0.11
Adenocarcine Adenocarcine Epidermoid Anaplastic c	oma with dermoid carcinoma with der arcinoma	moid		::	61 14 1 3	6.76 1.55 0.11 0.33 2.87
Adenocarcine Adenocarcine Epidermoid Anaplastic c Malignant to	oma with dermoid carcinoma with der arcinoma eratoma	moid			61 14 1 3 26 10	6.76 1.55 0.11 0.33 2.87 1.11 0.22
Adenocarcin Adenocarcin Epidermoid Anaplastic e Malignant to Carcino-sarc Dysgerminor Granulosa co	oma with dermoid carcinoma with der arcinoma eratoma oma ell carcinoma	moid		::	61 14 1 3 26 10 2	6.76 1.55 0.11 0.33 2.87 1.11
Adenocarcin Adenocarcin Epidermoid Anaplastic e Malignant to Carcino-sarc Dysgerminor	oma with dermoid carcinoma with der arcinoma eratoma oma ell carcinoma	moid		::	61 14 1 3 26 10 2 36	6.76 1.55 0.11 0.33 2.87 1.11 0.22
Adenocarcin Adenocarcin Epidermoid Anaplastic e Malignant to Carcino-sarc Dysgerminor Granulosa co	oma with dermoid carcinoma with der arcinoma eratoma oma ell carcinoma ell carcinoma	moid			61 14 1 3 26 10 2 36 25	6.76 1.55 0.11 0.33 2.87 1.11 0.22 3.99 2.77
Adenocarcin Adenocarcin Epidermoid Anaplastic c Malignant to Carcino-sarc Dysgerminor Granulosa co Arrhenoblast	oma with dermoid carcinoma with der arcinoma eratoma ena ell carcinoma ell carcinoma ell carcinoma eta	moid			61 14 1 3 26 10 2 36 25 1	6.76 1.55 0.11 0.33 2.87 1.11 0.22 3.99 2.77
Adenocarcin Adenocarcin Epidermoid Anaplastic e Malignant to Carcino-sarc Dysgerminor Granulosa co Arrhenoblast Fibrosarcoma	oma with dermoid carcinoma with der arcinoma eratoma ena ell carcinoma ell carcinoma ell carcinoma ena	moid			61 14 1 3 26 10 2 36 25	6.76 1.55 0.11 0.33 2.87 1.11 0.22 3.99 2.77 0.11 0.33
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Adenocarcin Adenocarcin Epidermoid Anaplastic e Malignant to Carcino-sarc Dysgerminor Granulosa e Arrhenoblast Fibrosarcoma Mesonephror Lymphosarco	oma with dermoid carcinoma with der arcinoma eratoma oma ell carcinoma ell carcinoma a a oma oma oma	 moid 			61 14 1 3 26 10 2 36 25 1 3 2	6.76 1.55 0.11 0.33 2.87 1.11 0.22 3.99 2.77 0.11 0.33 0.22 0.11
Adenocarcin Adenocarcin Epidermoid Anaplastic e Malignant te Carcino-sarc Dysgerminor Granulosa ce Arrhenoblast Fibrosarcoma Mesonephror Lymphosarco Secondary Kruckenberg	oma with dermoid carcinoma with der arcinoma eratoma oma ell carcinoma a a oma oma a oma oma oma oma oma				61 14 1 3 26 10 2 36 25 1 3 2	6.76 1.55 0.11 0.33 2.87 1.11 0.22 3.99 2.77 0.11 0.33 0.22 0.11
Adenocarcin Adenocarcin Epidermoid Anaplastic e Malignant te Carcino-sarc Dysgerminor Granulosa ce Arrhenoblast Fibrosarcoma Mesonephror Lymphosarco Secondary Kruckenberg Secondary fr	oma with dermoid carcinoma with der arcinoma eratoma oma ell carcinoma ell carcinoma a a oma oma oma	 moid 			61 14 1 3 26 10 2 36 25 1 3 2	6.76 1.55 0.11 0.33 2.87 1.11 0.22 3.99 2.77 0.11 0.33 0.22 0.11

TABLE II

Age Incidence of 622 Benign Neoplasms

Type of Neoplasms	Less than 10	11-20	2130	31-40	41-50	51-60	61-70	Above 70	Un- known
Mucinous				- Europe		4 50 3 100			
cystadenoma	ween.	20	46	24	33	27	8	2	5
Serous cystadenoma Papillary sercus	a	11	43	43	31	14	4	3	1
cystadenoma Benign cystic	-	5	5	5	10	3	2	1	-
teratoma Benign solid	-	13	48	34	21	6	2	-	-
teratoma	2	5	10	5	2	_	-	-	
Struma ovarii Struma with	-	-	í	1	2	1	-	-	-
teratoma Struma with mucinous		-	1	-	-	-		-	-
Cystadenoma Teratoma with mucinous	-	T	-	1	-	-	-	-	-
cystadenoma		-	1				1	-	-
Cyst adenofibroma		_	5	2	6	3	3	-	-
Adenofibroma	-	-		1	2		_		
Fibroma	-	2	3	4	9	2	2	-	-
Thecoma		1	2	_	2	1			-
Brenner tumour Brenner with mucinous	-	-	2		1	1	4	-	-
cystadenoma	-	-				2	1		-
Haemorrhagic cyst	-	4	24	17	8	3	_	2	1
Total	2	61	191	137	127	63	26	8	7

blastoma weighed 0.5 kg. and measured 25 cms in size.

In a few cases histologically different tumours were noted in the same specimen. The combinations noted were, mucinous cystadenoma with teratoma, papillary cystadenoma with adenofibroma, papillary serous cystadenoma with pseudomucinous cystadenoma.

In 8 cases of papillary cystadenocarcinoma and one case of granulosa cell tumour, metastasis were present. The usual sites of metastasis were cervix, bladder, endometrium, omentum and lymph nodes which included iliac, paraaortic, inguinal and supraclavicular.

Discussion

The ovary is complex in its embryology, histology, steroidogenosis, and potential for malignancy. It is made up of germ cells, follicular cells, and mesenchymal tissue, each with its own potential to form tumour. The reported incidence of the different types of ovarian tumours varies widely. (Gault 1954; Agarwal 1962; Philips 1965; Tyagi et al 1967; Vora et al 1969).

TABLE III Age Incidence of 281 Malignant Neoplasms

Type of Neoplasms	Less than 10	11-20	21-30	31-40	41-50	51-60	61-70	Above '70	Un- known
Papillary serous									
carcinoma	*******	4	8	16	19	14	2		1
Mucinous carcinom: Papillary	a —	4	4	7	4	3	_	_	-
adenocarcinoma	1	2	6	12	20	16	4	_	_
Adenocarcinoma Adenocarcinoma	1	=	2	1	6	3	- Chinagh	-	1
with dermoid Epidermoid carcinoma with	-	*******	-	-	1		-	m100m	
dermoid				1	2				
Malignant teratoma Anaplastic	-	1	4	3	1	1	-		-
carcinoma	V-0	2	5	3	10	6		Survey	-
Carcino sarcoma		_	1	1	termine		_	_	_
Dysgerminoma Granulosa cell	1	23	6	5		1		-	_
Tumour	tom.	1	8	3	9	2	_ 1	-	1
Arrhenoblastoma	-	manua.	1	demands.	derroga,	****	~~	-	
Mesonephroma		-	_		1	-	_		1
Fibrosarcoma		1	-	1	1	Source	-	-	
Lymphosarcoma	-	1	_	_		-		hellenda	(September 1997)
Krukerberg tumour Secondary from Adeno-		-	1		-	_	***************************************	Aveal	- Allerton
carcinoma Secondary from	-	-	alasma.	3	3	2	-	_	-
leiomyosarcoma				-	1	-	-		-
Total	3	39	46	56	78	48	7	_	4

The commonest ovarian tumour encountered is the cystadenoma which forms more than 1/3rd of the total ovarian neoplasms. (345 out of 903). The serous type shows the highest incidence i.e. 20.03% of total, followed by mucinous type 18.27%. Gault et al (1954) and Tyagi et al (1967) also observed that serous cystadenomas was the commesest ovarian tumour.

70.3% of the ovarian neoplasms occurred in the reproductive period i.e. 21-50

the second decade. The youngest patient in this series was 9 years old and the oldest 77 years of age. Borges (1955) reported an age range of 14-80. In the series reported by Vora et al (1969) the youngest was 10 years and the oldest 65 years. Benign neoplasms showed a higher incidence in the younger age group i.e. 21-30, as compared to the high incidence of malignant neoplasms in the 41-50 age group.

Among the benign neoplasms, the comyears, with the maximum incidence in mon tumours were serous cystadenomas 32.1% mucinous cystadenomas 29.3% and benign teratomas 28%. The incidence of mucinous cystadenoma is comparatively less in this series (18.27% of the total) as compared to 28.3% reported by Agarwal *et al* (1962).

There were 157 cases of benign teratomas forming 17.59% of the total ovarian neoplasms. Tyagi et al (1967) noted an incidence of 12.83%. There were 2 cases of mucinous cystadenoma with teratoma, 5 cases of struma ovarii and one each of struma with teratoma and struma with mucinous cystadenoma. Gault et al (1964) found 1 case of struma ovarii in 317 ovarian neoplasms and Philips (1965) noted 2 cases in a study of 426 ovarian tumours. Combination of teratomatous elements along with struma and mucinous cystadenoma favours a common histogenesis for these as one sided development of a teratoma where one particular teratomatous element overshadows all others to form a separate tumour of its

Fibromas formed 3.9% of the benign neoplasms and 2.44% of the total. Tyagi et al (1967) reported a 5% incidence and Vora et al (1969) noted 3 cases out 188 benign neoplasms. There were 18 cases of cystadenofibromas and 3 cases of adenofibroma in this series. Gault et al (1964) noted 9 cases out of 317 ovarian neoplasms.

There were 11 cases of Brenner tumour (19%) and in 3 of these it was associated with pseudomucinous cystadenoma. Tyagi et al (1967) had only one case in their series of 120 cases, and this occurred in a 13 year old girl. The age range for Brenner tumour in this series is from 21-70 with 4 cases in the 61-70 age group. Cases of Brenner tumour with mucinous cystadenoma occurred in the older age group 51-70. Vora et al (1956) state that the

large Brenner tumours are associated with mucinous cystadenoma. However, in this series, no such association has been noted.

Incidence of malignancy in the present series was 31.12%. Table IV shows the reported incidence of malignancy. The incidence of malignancy in our series is higher than most of the reported series, except that of Patel et al (1964).

TABLE IV

Incidence of Malignancy in the Reported Series of Ovarian Tumours as Compared with the Present Series

Year	Author	Percent- age in- cidence of Ma- lignancy
1931	Mayer	14.9%
1949	Allen and Hertig	15.2%
1955	Purandare	23%
1957	Chitkara and Sharma	28%
1962	Agarwal and Saxena	22.97%
1964	Mehta and Purandare	28.1%
1964	Patil et al	42.0%
1967	Tyagi et al	24.17%
1969	Vora and Bhargava	19.01%
1971	Present series	31.12%

Carcinoma is the most common primary malignant tumour of the ovary. In the present series, there were 190 cases of primary carcinoma of the ovary among 281 malignant neoplasms which forms 21.05% of total ovarian neoplasms. Of these 7.09% were serous cystadenocarcinomas and 2.44% mucinous cystadenocarcinomas.

Of the 190 primary carcinomas of the ovary, 100 were solid tumours. This forms 11.08% of the total ovarian neoplasms. Patel et al (1964) have reported a similar incidence (12.2%) in a study of 147 ovarian tumours. But lower incidence has been reported by Agarwal et al (1962)

Mehta et al (1964) and Tyagi et al (1967). Among the solid tumours, 61 were papillary, 25 anaplastic and 14 adenocarcinomas. Meyer (1932) and Tyagi et al (1967) have reported that papillary carcinomas have a lower incidence than non-papillary type but in the present series papillary type was more common.

Malignancy in dermoid is rare. Meyer (1932) has noted 1.71% in 1268 cases of dermoid. The present series showed an incidence of 3.12% in 182 cases of dermoid. The common malignant change reported is epidermoid carcinoma. An adenocarcinoma arising from dermoid is a rarity. One such case was noted in this series.

There were 36 cases of dysgerminoma, forming 3.99% of total ovarian neoplasms and 12.8% of ovarian malignancy. This is quite high when compared to 4.5% reported by Muller et al (1950) and 3.5% by Morris and Scully (1958) and 4.3% by Kapas (1969). Vora and Bhargava (1969) noted 10.2% in 49 malignant neoplasms. However Gault et al (1954) had only an incidence of 1.8% in their series.

Granulosa cell tumours comprised 8-9% of malignant ovarian neoplasms in the present series. Vora and Bhargawa (1969) noted an incidence of 18.9%. In the series reported by Tyagi et al (1967) these formed 3.33% of total ovarian neoplasms, while in the present series it was only 2.77%. Most of the cases occurred in childbearing period, but 6 cases were seen in postmenopausal period and these presented with symptoms of bleeding. Hyperplasia of endometrium was seen in one case. One case showed evidence of secondaries and had associated pleural effusion.

Rare neoplasms of the ovary like fibrosarcoma mesonephroma and lymphosarcoma were also seen in this series. Fibrosarcomas formed .33% of total neoplasms and 3.11% of ovarian malignancy. Two of these occurred in 30-50 age group and 1 case was in a 19 years old woman.

Mesoneophroma of the ovary is an uncommon malignant tumour arising from mesometanephric rests. Two cases were diagnosed as mesonephroma in this series. These showed the typical histological appearance of tubular pattern formed by flattened or cuboidal cells with scanty clear cystoplasm and oval nuclei.

The single case of primary lymphoma of the ovary in a 17 year old girl has been reported elsewhere (Ramachandran and Gracy 1970).

Summary

The commonest ovarian neoplasm encountered in the series was cystadenoma.

70.3% of the neoplasms occurred in the reproductive period with the maximum incidence in the second decade.

The percentage incidence of malignancy observed was 31.14%.

The incidence of various neoplasms are compared with the reported series.

Rare primary ovarian neoplasms like fibrosarcoma, mesonephroma and lymphosarcoma were also observed in this series.

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