

## A 5 Year Prospective and Retrospective Study- Of Tumors and Tumor-Like Lesions of Testis

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### **Abstract:**

**Introduction:** Testicular tumors form an interesting group of malignant neoplasms with many exceptional and even unique epidemiological features. They occur mainly in young and middle aged adults and this suggests that their risk factors are different from those relevant to most epithelial cancers. Despite the major advances there is a gap in the knowledge in understanding the origin, pathological process and therapeutics of testicular and paratesticular tumors and tumor like conditions. This is a prospective as well as retrospective study, undertaken in the department of pathology, Siddhartha Medical College, Vijayawada; during the period from March 2014 to March 2019. Testicular specimens sent to the Department of Pathology, during this period were included in the study.

**Materials and methods:** The source of data for both prospective and retrospective study are testicular specimens received in the department of pathology, SMC, Vijayawada during the period from March 2014 to March 2019 ( 5 years). The macroscopic and microscopic findings in these specimens were tabulated and analysed. In cases of retrospective study the macroscopic findings were collected from records and sections cut from blocks were analysed

**Results:** A total of 80 cases were studied in 5 years. Out of 80 cases, retrospective component study included 41 specimens and prospective component study included 39 specimens Testicular diseases were relatively rare.

Age group most commonly involved was 21-40 years of age. Testicular lesions were rare at the extremes of age Majority of the testicular lesions were non neoplastic comprising of 62 cases (77.50%), out of total 80 cases Neoplastic lesions were found in 18 cases (22.50%), out of total 80 cases. Germ cell tumors were the most common neoplastic lesions comprising of 16 cases and forming 88.89% of all testicular neoplasms.

**Conclusion:** Testicular diseases were relatively rare. Although testicular disease is usually only encountered by the pathologist, urologist and medical oncologist of large medical centres with a special interest and experience in this field, the great advances in treatment make it imperative to provide the exact diagnosis and to use the correct therapeutic approach when one encounters such patients. Germ cell tumors were the most common neoplastic lesions comprising of 16 cases and forming 88.89% of all testicular neoplasms. Histopathological diagnosis of teratoma was made in three cases making 16.67% of all testicular neoplasms. Mixed germ cell tumor diagnosis was made in 6 cases forming 33.33% of all testicular neoplasms. THE study was done to know the diverse histological structure and variation in the incidence of lesions in the different parts of the world,--

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### **I. Introduction**

Testicular tumors form an interesting group of malignant neoplasms with many exceptional and even unique epidemiological features. They occur mainly in young and middle aged adults and this suggests that their risk factors are different from those relevant to most epithelial cancers. Moreover it is believed that the origin of the great majority of testicular tumors is the germ cell. This may be of great importance, because possible environmental risk factors in testicular cancer are liable to influence future generation.

Although testicular disease is usually only encountered by the pathologist, urologist and medical oncologist of large medical centres with a special interest and experience in this field, the great advances in treatment make it imperative to provide the exact diagnosis and to use the correct therapeutic approach when one encounters such patients. Despite the major advances there is a gap in the knowledge in understanding the origin, pathological process and therapeutics of testicular and paratesticular tumors and tumor like conditions.

The diverse histological structure and variation in the incidence of lesions in the different parts of the world, prompted us to undertake the present study.

### **Aims And Objectives**

The present study deals with tumors and tumor like lesions of testis. Duration of study is 5 years comprising of three years retrospective study and subsequent 2 years prospective study. This study was undertaken with the following objectives :

1. To study the histomorphological features of various tumors and tumor-like lesion of testis.
  2. To analyse available clinical data, for establishing clinic-pathological correlation.
- Specimens received in the department of pathology, Siddhartha Medical College, Vijayawada, during the past 3 years were included in the present study.

### **II. Materials and Methods**

The present study emphasizes on the diagnostic utility of histopathology of tumors and tumor like lesions of testis. This is a prospective as well as retrospective study, undertaken in the department of pathology, Siddhartha Medical College, Vijayawada; during the period from March 2014 to March 2019. Testicular specimens sent to the Department of Pathology, during this period were included in the study.

#### **SOURCE OF DATA:**

The source of data for both prospective and retrospective study are testicular specimens received in the department of pathology, SMC, Vijayawada during the period from March 2014 to March 2019 ( 5 years).

#### **METHOD OF COLLECTION OF DATA:**

All the patients were subjected to detail clinical examination and routine laboratory tests. Wherever possible radiological imaging techniques were employed. In the cases collected during the retrospective study their clinical records were analysed.

Testicular specimens were fixed in 10% NBF and then routinely processed to have paraffin sections and stained with Haematoxylin & Eosin routinely. Multiple blocks were taken based on size, variability of gross features and adjacent areas and sections analysed.

The macroscopic and microscopic findings in these specimens were tabulated and analysed. In cases of retrospective study the macroscopic findings were collected from records and sections cut from blocks were analysed.

#### **SAMPLE SIZE:**

A total of 80 cases were studied in 5 years. Out of 80 cases, retrospective component study included 41 specimens and prospective component study included 39 specimens.

#### **INCLUSION AND EXCLUSION CRITERIA:**

All specimens with macroscopically and microscopically detected tumors and tumor like lesions of testis mentioned in the WHO classification were included and remaining lesions like scrotal lesions and congenital anomalies were excluded

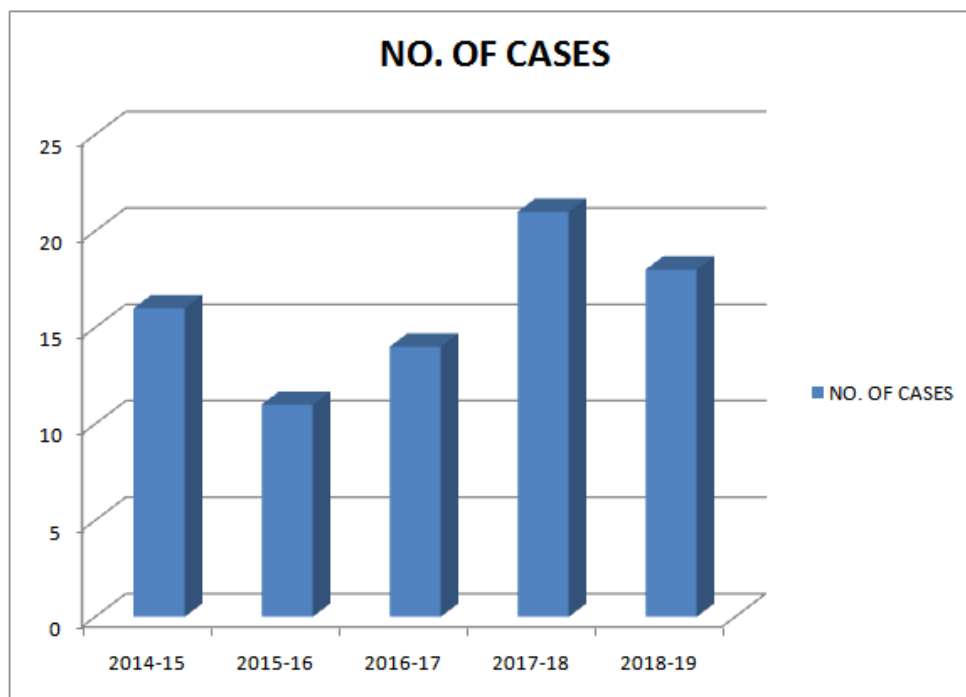
### **OBSERVATIONS**

#### **RESULTS:**

Salient observations made in this study are as follows-

**Table – 1** Number of cases in each year

SL.NO	YEARS	NO. OF CASES	PERCENTAGE
1.	2014-15	16	20%
2.	2015-16	11	13.75%
3.	2016-17	14	17.5%
4.	2017-18	21	26.25%
5	2018-19	18	22.5%
TOTAL		80	100%



**AGE DISTRIBUTION:**

Age group of patients ranged from 6 months to 85 years. Age group most commonly involved was 21-40 years of age. In the present study 36 cases (45%) were in the age group of 21-40 years. Testicular lesion were rare at the extremes of age.

SL.NO	AGE(YEARS)	NO.OF CASES	PERCENTAGE
1.	0-10	05	6.25
2.	11-20	08	10
3.	21-30	18	22.5
4.	31-40	18	22.5
5.	41-50	16	20
6.	51-60	08	10
7.	61-70	03	3.75
8.	71-80	03	3.75
9.	81-90	01	1.25
TOTAL		80	100

**GROSS:**

**1.NATURE OF SPECIMENS:**

Two types of specimens were sent to the pathology department – orchidectomy and biopsy specimens. Specimens received in our department showed variable sizes. Minimum size of 0.5x0.5 cms and maximum being 15x10x6 cms. Surface of all the orchidectomy specimens varied from smooth to bosselated to nodular. Biopsy specimens were grey white soft tissue bits.

SL.NO	SPECIMEN	NO.OF CASES	PERCENTAGE
1.	ORCHIDECTOMY	45	56.25%
2.	BIOPSY	35	43.75%

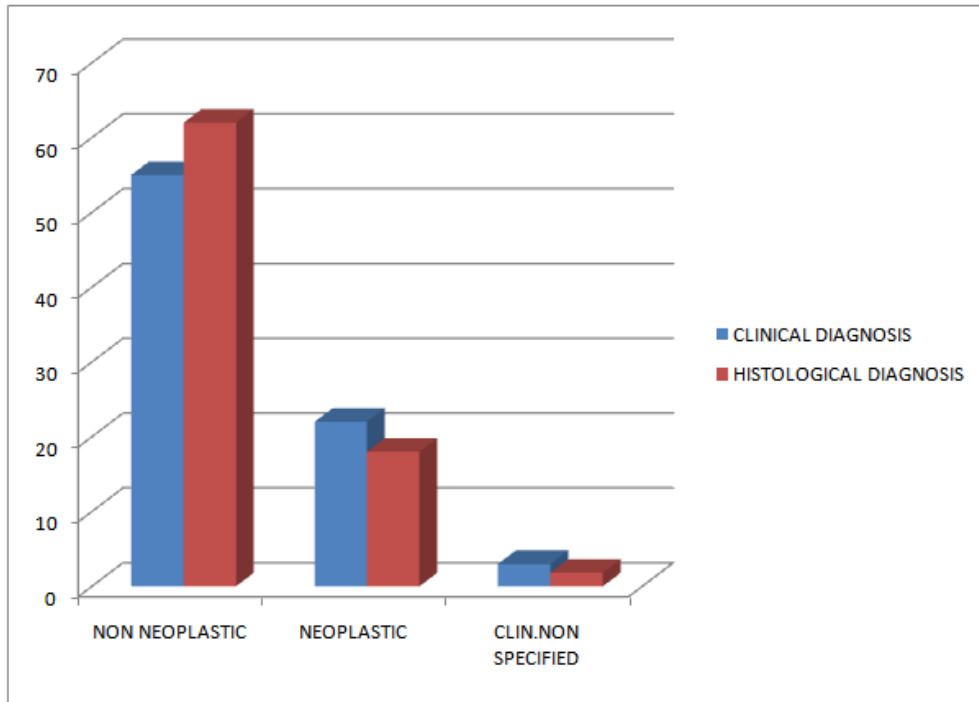
**2.CUT SECTION:**

Among 45 orchidectomy specimens studied, necrosis was the commonest finding seen in 20 cases (44.44%). Involvement of tunica was seen in 10 cases (22.22%)

**CLINICO-HISTOLOGICAL CORRELATION:**

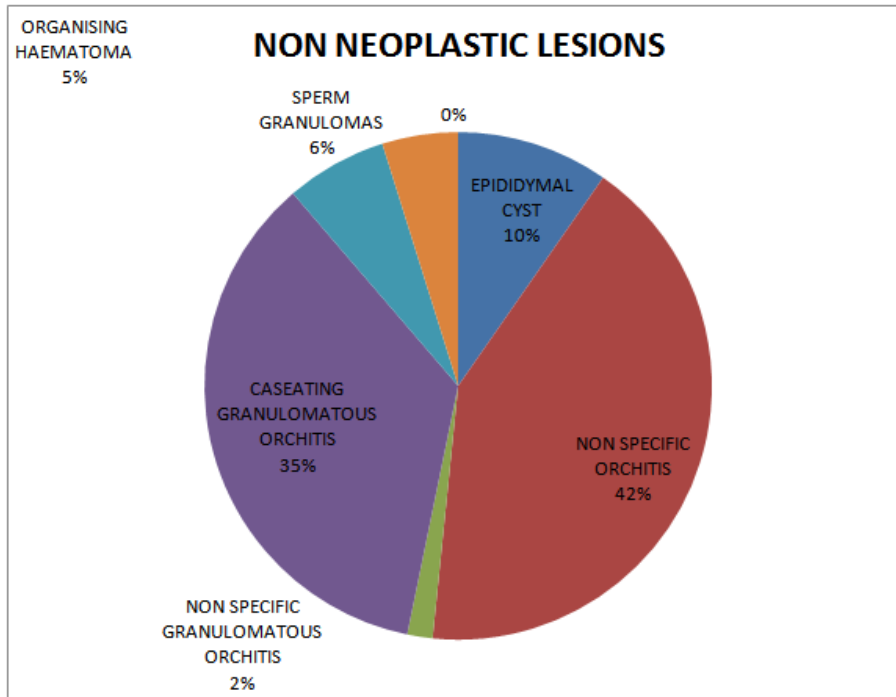
Present study showed that there was correlation between the clinical diagnosis and the histological diagnosis.

In this study, clinically 55 case were diagnosed as non neoplastic and 22 case as neoplastic, whereas histologically, 62 case were diagnosed as non neoplastic and 18 case as neoplastic.



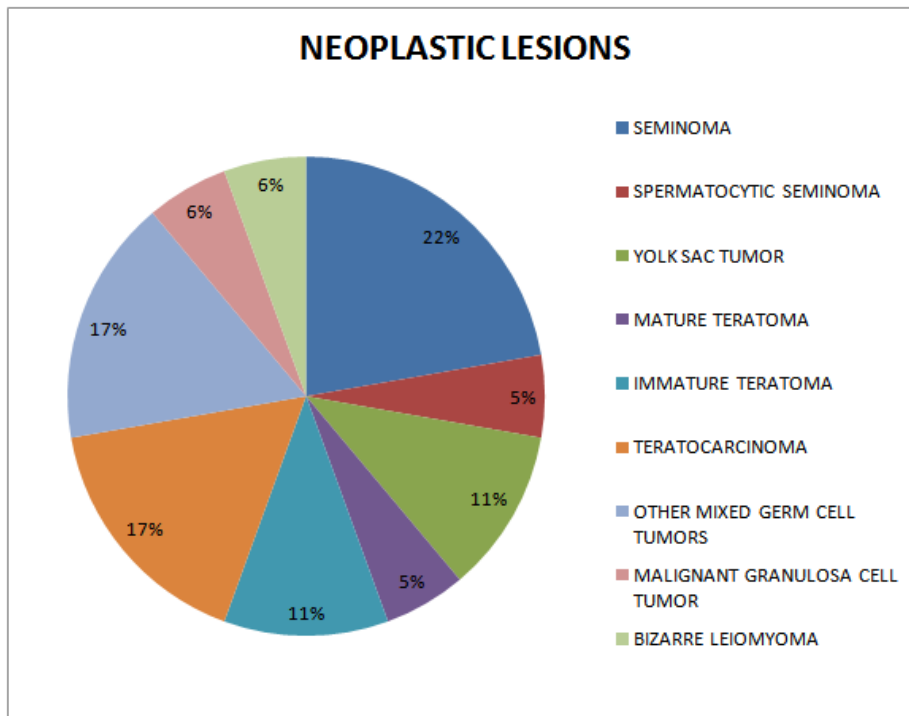
**HISTOPATHOLOGICAL DIAGNOSIS OF NON NEOPLASTIC LESIONS:**

SL.NO	HISTOPATHOLOGICAL DIAGNOSIS	NO.OF CASES	%
1.	EPIDIDYMAL CYST	6	9.67%
2.	NON SPECIFIC ORCHITIS	26	42%
3.	NON SPECIFIC GRANULOMATOUS ORCHITIS	1	1.6%
4.	CASEATING GRANULOMATOUS ORCHITIS	22	35.5%
5.	SPERM GRANULOMA	4	6.45%
6.	ORGANISING HEMATOMA	3	4.83%
TOTAL		62	100



**HISTOPATHOLOGICAL DIAGNOSIS OF NEOPLASTIC LESIONS:**

S.NO	HISTOPATHOLOGICAL DIAGNOSIS	CASES	%
1.	SEMINOMA	4	22.2%
2.	SPERMATOCYTIC SEMINOMA	1	5.5%
3.	YOLK SAC TUMOR	2	11.1%
4.	MATURE TERATOMA	1	5.5%
5.	IMMATURE TERATOMA	2	11.1%
6.	TERATO-CARCINOMA	3	16.67%
7.	OTHER MIXED GERM CELL TUMORS	3	16.67%
8.	MALIGNANT GRANULOSA CELL TUMOR	1	5.5%
9.	BIZARRE LEIOMYOMA	1	5.5%
TOTAL		18	100



**GERM CELL TUMORS:**

Among 18 neoplastic cases, 16 cases were of germ cell origin, forming 88.89% of all testicular neoplasms. Out of 16 cases, 10 cases were of single cell type and remaining cases were of mixed germ cell type. Seminoma was the most common germ cell tumor comprising of 4 cases, 22.22% of all testicular neoplasms. Average age of the patient was 50 years.

Teratoma comprised 16.67% of all testicular neoplasms. One case was of mature type and other two cases were of immature type. Teratoma component combined with other components were seen in another 5 cases making 27.78% of all testicular neoplasms.

**III. Discussion**

This study deals with tumors and tumor-like lesions of testis and based total 80 cases received in the Department of Pathology, Siddhartha Medical College, Vijayawada, during 5 years period (March 2014-march 2019).

Total number of cases received in each year are variable. Rarity and variability of testicular lesions was observed by other authors also. In the present study age distribution of patients, ranges from six months to 85 years, age group most commonly involved are 21-40 years of age. Testicular lesions are rarely seen at the extremes of age. These findings confirm well with observations of other authors.

In the present study, 45 orchidectomy specimens were sent for examination, while biopsy was done in 35 cases. Orchidectomy was done in all cases, which were clinically diagnosed as testicular tumors. Both orchidectomy and biopsy procedures were done in clinically diagnosed non-neoplastic lesions. These findings are well correlated with Mostofi's observations that biopsy generally accepted from other sites, is absolutely contraindicated in testicular tumors. If a tumor is suspected, the testis must be removed in toto.

In the present study, 16 germ cell tumors were reported accounting for 8.89% of all testicular neoplasms. Germ cell tumors of testis have accounted for 94% of all testicular tumors in the literature of FK Mostofi. In the present study 10 cases, accounting to 62.50% were single cell type this incidence of single cell type is well correlated with FK Mostofi's observations, who made single cell type incidence as 60%.

In the present study, seminoma is the commonest germ cell tumor comprising of 4 cases, that is 22.22% of all testicular tumors. EK Mostofi reported the incidence of 35-71% depending on the type of hospital population under study, where as, Anderson's study accounts for 35-45% of all testicular tumor and Juan Rosai study accounts for 30-40%. In the present study, one patient was 75 years old, another patient 45 years old, while other two patients were 40 years of age. No case of seminoma was reported before puberty. All these clinical findings are comparable to other authors. Grossly in all cases, there was an enlargement of testis, and in one case there was total replacement of testis by tumor (In half the patients the entire testis is replaced by tumor EK Mostofi).

Mostofi reported that necrosis is infrequent in seminoma, but in present study, two cases, i.e. 50% were showing necrotic areas. The microscopic features of seminoma in present study are similar to the features reported in various literatures.

In the present study, only one case of spermatocytic seminoma was reported. Age of the patient was 55 years. Tumor measures 7x6x5 cms replacing entire testis, and cut section was showing necrosis and haemorrhage.

Microscopic examination showed three types of tumour cell large intermediate and small cells. Intermediate cells have spireme type of chromatin pattern. Haemorrhage and necrosis was present. Vascular invasion and spermatic cord involvement was seen.

All these features are comparable to those described by F.K. Mostofi. Microscopic findings are also similar to the findings described by other authors, A. Talerman in 1979, Marc H. Zuckman in 1988, and E.K. Mostofi in 1973. Incidence of spermatocytic seminoma ranges from 1.7% to 12% of all seminoma in various literature. FK Mostofi in 1973 reported 9%, Juan Rosai et al. in 1969 reported 7%, and A. Talerman in 1979 reported 4.4%. In present study, the incidence of spermatocytic seminoma is 20%, out of total five seminoma cases.

In present study, two cases of Yolk sac tumor were reported. Age of both the patients was two years. Grossly, both cases were showing multiple cysts.

Microscopically, perivascular schiller-duval bodies and hyaline droplets were present in both the cases. These findings are comparable to the findings described by G. Barry Pierce et al. in 1969, Aleksander Talerman and Lawrence M Bothin 1986, F.K. Mostofi in 1973 and Ivan Damjanov in 1978.

In the present study, out of three cases of teratoma, one case was mature Teratoma and two cases were immature teratoma. Gross and microscopic findings in present study are also similar to as described in literature.

In the present study, incidence of teratoma of testis in pure form is 16.67% of all Testicular neoplasms, and in combination with other histologic type incidence is 44.44% of all testicular neoplasms. In the literature, teratoma or less in pure form has accounted for 9% of all testicular neoplasms, and in combination with other histologic type teratoma has accounted for 40% of all testicular neoplasms.

In the present study, six cases, i.e., 33.33% of all testicular tumors were histologically diagnosed as mixed germ cell tumors. Out of six cases, three cases were teratocarcinoma, one case was mixed germ cell tumor (Teratoma + chorio-carcinoma + Yolk-Sac tumor), one case was mixed germ cell tumor (Embryonal carcinoma + Endodermal-sinus tumor), and one case was mixed germ cell tumor (choriocarcinoma Endodermal sinus tumor). Present study is showing that teratocarcinoma is the commonest mixed germ cell tumor constituting 16.67% of all testicular neoplasm where as, F.K. Mostofi's study showed 24% of testicular neoplasms.

In the present study one case, i.e., 5.55% was histologically diagnosed as granulosa cell tumor, grossly, tumor measured 5x4x4cm with variable consistency. On cut section tumor was well delineated. Microscopically, tumor cells were large with scanty cytoplasm and large nucleus having nuclear grooves. These cells were arranged in sheets, cords and insular pattern. Call-Exner bodies were present. These gross and microscopic findings were also observed by other authors.

In present study, six cases were histologically diagnosed as epididymal cysts. All cases were clinically also diagnosed as epididymal cyst, so there is 100% correlation between clinical diagnosis and histological diagnosis. No malignancy was detected in any case. These findings are well correlated with other authors.

In the present study histopathological diagnosis of non-specific orchitis was made in 26 cases, i.e., 32.5% of all testicular lesions. It was the commonest histopathological diagnosis made in the study. Microscopically, all cases were showing moderate to dense infiltration of inflammatory cells.

In chronic non-specific orchitis, inflammatory cells were predominantly lymphocytes, whereas in acute non-specific orchitis, predominant inflammatory cells were neutrophils. Abscess formation was seen in some cases of acute epididymo-orchitis. According to the study of Gregor Mikuz and Ivan Damjanov, since, "The inflammation is often asymptomatic and subclinical", the real incidence and prevalence of non-specific orchitis

are unknown. Descriptive features as mentioned in the present study are perfectly correlated with the description mentioned by other authors.

One case was histologically diagnosed as non-specific granulomatous orchitis. Clinical diagnosis of this case was testicular tumor. Orchidectomy was done in this case. Gregor Mikuz and Ivan Damjanov in their review study observed that 5.1% of presumptive diagnosis of neoplasia turned out to be nonspecific granulomatous orchitis.

Microscopically, there was a diffuse inflammatory infiltrate composed of lymphocytes and macrophages. There were granulomas composed of epithelial cells, lymphocytes and fibroblasts at periphery. These findings were also observed by other authors.

Histopathological diagnosis of caseating granulomatous orchitis was made in 22 cases forming 27.75% of all testicular lesions. Microscopically, all the cases were showing presence of granulomas with central caseation bordered by epithelioid cells and in few cases Langhans type of giant cells were seen. Surrounding this, there was inflammatory infiltrate mainly consisting of lymphocytes, and few neutrophils and plasma cells. Tubercular epididymo-orchitis occurs most often in middle life and is rare in children, as reported by Gregor Mikuz and Ivan Damjanov. These microscopic findings and age incidence findings well correlate with our present study except in two cases where one patient was 6 years old and another 10 years old.

In the present study, four cases, 5% of sperm granuloma were diagnosed. Microscopic examination of sperm granuloma was showing granuloma formation with dense, chronic inflammatory infiltrate in all cases. In the center of these granulomas few neutrophils and sperms were seen. Kuffer and Buschmann have examined 250 autopsies and found sperm granulomas in 2.6% whereas in present study diagnosis of sperm granulomas was made in 5.

In the present study, three cases of organising haematoma were histologically diagnosed. All these cases were clinically diagnosed as non-neoplastic lesion. Microscopic examination shows blood clot and fatty tissue with inflammatory cell infiltrate in all three cases. This entity was included in the tumor like lesions classified according to the WHO classification.

In the present study varieties of neoplastic diseases involving the testis and paratesticular tissue were seen and results are compared with the studies of others.

#### **IV. Conclusion**

A prospective and retrospective study of tumors and tumor-like lesions of testis was undertaken during the period of March 2014-March 2019 (5 years), to evaluate histopathological features of various tumors and tumor-like lesions of the testis.

Testicular diseases were relatively rare. Age group most commonly involved was 21-40 years of age. Testicular lesions were rare at the extremes of age. Orchidectomy was done in all cases which were clinically diagnosed as testicular tumors, whereas orchiectomy and biopsy was done in clinically diagnosed non-neoplastic lesions. Majority of the testicular lesions were non-neoplastic comprising of 62 cases (77.50%), out of total 80 cases. Neoplastic lesions were found in 18 cases (22.50%), out of total 80 cases. Testicular tumors form an interesting group of malignant neoplasms with many exceptional and even unique epidemiological features. They occur mainly in young and middle aged adults and this suggests that their risk factors are different from those relevant to most epithelial cancers. Despite the major advances there is a gap in the knowledge in understanding the origin, pathological process and therapeutics of testicular and paratesticular tumors and tumor like conditions.

#### **References:**

- [1]. A. Talerman, "Spermatocytic seminoma". *Cancer*, 1980; 45:2169-2176
- [2]. Ackerman's surgical pathology, Edt. Juan Rosai, Mosby, 9th Edition
- [3]. Antonio R. Perez - Atayde, Nancy Joste and Howard Mulhern
- [4]. Hafez. "Newborn granulosa cell tumor of the testis". *JUrol*. 138
- [5]. "Juvenile granulosa cell tumour of the infantile testis", *Am J Surg. Path*, 20(1): 727-739
- [6]. Anderson's Pathology, Edt. Ivan Damjanov, 11th Edition
- [7]. Anthony A., and Borski, "Diagnosis, staging and natural history of testicular tumors". *Cancer*, 32: 1202-1205
- [8]. David J.B. Ashley, "Origin of teratomas". *Cancer*, 32: 390-394
- [9]. David T. uehling, Janet E. Smith, Richard Logan and Gholon- Reza F.B. Mahon, F Gosset. R.G. Trinity, and P.O Madsen "malignant interstitial cell testicular tumor". *Cancer*, 31: 1208-1210
- [10]. G. Barry Pierce. Weldon K. and Robert W. Huntington, "Yolk sac tumor of the testis". *Cancer*. 25: 642-657.
- [11]. George W. Kaplan et al.. "Prepubertal yolk sac testicular tumors". *Urol*, Part-2, 140: 1109-1112
- [12]. Gregor Mikuz and Ivan Damjanov, "Inflammation of the Testis epididymis, peritesticular membranes and scrotum
- [13]. Harry J. Tamoney, and Alfredo Noriega, "Malignant interstitial cell tumor of the testis" *Cancer*, 24:547-550
- [14]. Joan Austoker, "Screening for ovarian, prostatic and testicular cancers" *BMI*. 309: 315-321 Joan Austoker, "Screening for ovarian, prostatic and testicular cancers" *BMI*. 309: 315-321
- [15]. John N. Eble, "Spermatocytic seminoma" *Hum Pathol*. 25: 1035-1042
- [16]. Jozef Matoska, Dalibor Ondrus and Michal Hornak "Metastatic spermatocytic seminoma" *Cancer*, 62: 1197-1201

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- [17]. Judith A.Ferry,, Robert H. Young and Robert E.Scully Testicular andepididymalplasmacytoma. Am J Surg. Path. 21(5): 590-5981997
- [18]. Juan Rosai, Igal Silber and KavousKhodadoust"Spermatocytic seminoma" Cancer, 24: 92-11
- [19]. K.Scheiber, D.Ackermann and U.E.Studer. "Bilateral testicular germ celltumors".JUrol, 138: 73-76
- [20]. Kishor H. Shah, William C. Maxtod and Byungkyu Chun, "Epidermoidcysts of the tests". Cancer. 47: 577-582
- [21]. Manuel Nistal, Alberto Mate and Ricardo Paniagua. "Granulomatousepididymal lesion of possible ischemic origin" Am J SurgPathol21(8): 951-956
- [22]. Marc H. Zuckman. George Williams, Howard S. Levin "Mitosiscounting of seminoma" Hum Pathol, 19: 329-335
- [23]. Michael D. Melekos and Hans W.Absach,"Epididymitis, aspectsconcerning etiology and treatment" JOurnal, 138:83-86
- [24]. Michael J. Sworn, and Rose Buchanan, Malignant interstitial cell tumorof the testis" Hum Pathol, 12(1): 72-77
- [25]. Mostofi EK. "Testicular tumors. Epidemiologic,etiologic andpathologic features" Cancer, 32: 11861200
- [26]. Mostofi FK. "Pathology of germ cell tumors of testis - A progressreport" Cancer. 45: 17351754
- [27]. Mostofi FK. "Histological typing of testis tumours Geneva WHO1977
- [28]. Richard W. Grady, Jonathan H. Ross and Robert Kay, "Epidemiologicalfeatures of testicular teratoma in a prepubertal population" LofUrology, 157: 1191-1552
- [29]. Robbins. "Pathologic basis of disease", EdtCotran Kumar
- [30]. Robbins, 7th Edn
- [31]. Robert J. Fram, Marc B Garnick and Alan Retik. The spectrum ofalbuginea : Benign invasion by testicular tubules" JUrol. 138genitourinary abnormalities in patients with cryptorchidism withemphasis on testicular carcinoma. Cancer, 502243224
- [32]. Robert W Andrews Dana D.Copeland and Floyd A.Fried"Splenogonadal fusion". J. of Urology. 133: 105210531985
- [33]. S. Das, "Clinical surgery". 6th Edn
- [34]. Stanward S. Schmidt and Tate M. Minchler, "Pseudocysts of the tunica
- [35]. TalermaAleksander and Lawrence M.Roth. "Pathology of the testisand its adnexa". Churchill Livingstone, New York, 1986
- [36]. Thomas M Ulbricht, Robert Young and Robert EScully "Trophoblastic tumours of the testis other than classicchoriocarcinoma". Am J. of Surg. Path. 21(3): 2822881997
- [37]. Waun Ki Hong. Robert E. Wittes, Stenen T. Hajdu, Esteban Cvitkovicand Robert B Golbey, "The evolution of mature teratoma frommalignant testicular tumors" Cancer, 2987299
- [38]. Willis R.A., "Pathology of tumours", London: Butterworths. 1967.
- [39]. Winifred Gray, "Diagnostic cytopathology". Churchill Livingstone, 1Edn
- [40]. Y.Reinberg, J.C. Manivel, J. Llerena, G. Niehans and E.E. Fraley"Epidermoid cyst (monodermlalteratoma) of the testis". Br J ofUrology, 66: 648-651

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