

Spectrum of Thymic Neoplasms in A Tertiary Care Hospital¹Dr Silviya John, 3rd year Postgraduate, AJ Institute of Medical Sciences and Research Centre, Mangalore²Dr Purnima Rao, Professor, AJ Institute of Medical Sciences and Research Centre, Mangalore³Dr Sandhya I, Professor, AJ Institute of Medical Sciences and Research Centre, Mangalore⁴Dr Manu Mohan, Deputy Assistant Director of Health**Corresponding Author:** Dr Silviya John, 3rd year Postgraduate, AJ Institute of Medical Sciences and Research Centre, Mangalore.**How to citation this article:** Dr Silviya John, Dr Purnima Rao, Dr Sandhya I, Dr Manu Mohan, “Spectrum of Thymic Neoplasms in A Tertiary Care Hospital”, IJMACR- July - 2025, Volume – 8, Issue - 4, P. No. 108 – 112.**Open Access Article:** © 2025 Dr Silviya John, et al. This is an open access journal and article distributed under the terms of the creative common’s attribution license (<http://creativecommons.org/licenses/by/4.0>). Which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.**Type of Publication:** Original Research Article**Conflicts of Interest:** Nil**Introduction**

The thymus is a primary lymphoid organ of the immune system located in mediastinum. It undergoes several changes as a result of various factors like ageing and medical disorders ¹.

Thymic neoplasms constitute a rare group of anterior mediastinal mass which shows heterogeneity in clinical presentation and histologic appearance.

Most patients with thymic tumors are asymptomatic at the time of diagnosis.

Even though diagnosis can be made by clinical course and radiology, biopsy followed by histopathology remains the gold standard for the diagnosis.

Now a day, Immunohistochemistry (IHC) is used as a criteria for diagnosis of thymic neoplasms with ambiguous histology².

Materials and Methods

This is a 6 year retrospective study conducted in the Department of pathology on resected and biopsy specimens of thymic neoplasms from 2017 to 2023.

The present study includes 25 cases of thymic neoplasms.

Request forms of the cases were retrieved from the archives to collect available clinical details such as age, gender, symptoms and size of the lesion.

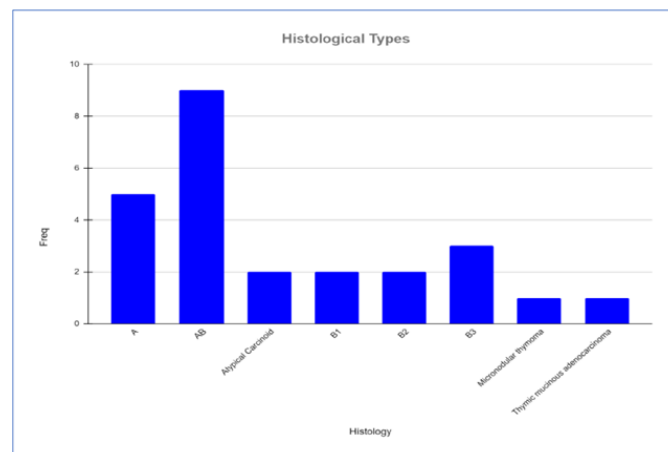
All the microscopic slides were retrieved from the archives and reviewed.

IHCs were done in 6 cases.

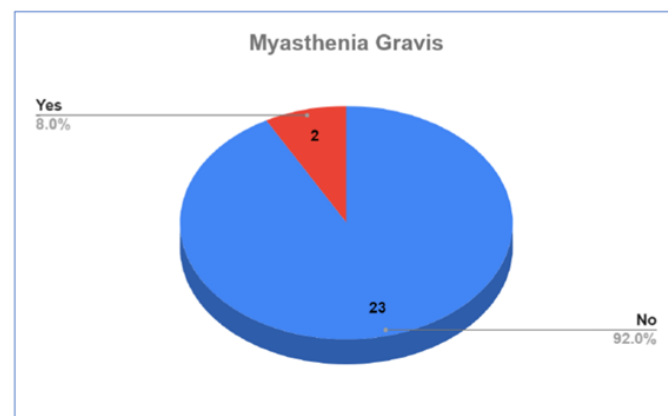
Findings were noted in Microsoft excel and analyzed using SPSS V.25.

Number	HIS number	Age	Sex	Symptoms or syndromes	size (cm)	Histology	HC	EXCISION/BIOPSY
15294/17	77M	77M	M	MUSCLE WEAKNESS	4x2.3x2	AB	Vimentin,CD3,EMA	BIOPSY
15294/17	69M	69M	M	DYSPOEA	10x7.4x3.5	AB		EXCISION
15494/17	53M	53M	M	DYSPOEA	13x5.2x4	B3		EXCISION
14332/17	57F	57F	F	CHEST DISCOMFORT	6x4x3	AB		BIOPSY
17023/17	54F	54F	F	CHEST DISCOMFORT	3.8x3.5x3	A	CK7,FE1	EXCISION
17289/17	63M	63M	M	INCIDENTAL	5x3.2x1.4	AB		EXCISION
77299/17	72F	72F	F	GOITRE	4x2.5x1.8	Atypical Carcinoid		EXCISION
9352/18	51M	51M	M	WEIGHTLOSS,HOARSENESS	18.5x8x5.5	Atypical Carcinoid		EXCISION
9353/18	64M	64M	M	CHEST PAIN	0.7x0.3x0.2	Microcystic thymoma		BIOPSY
104892/18	29M	29M	M	INCIDENTAL	12x9x8	B1		EXCISION
114732/18	23M	23M	M	BACKPAIN,ARM PAIN	5x2.6x2.5	thymic mucinous adenoma		BIOPSY
124294/18	60F	60F	F	MYASTHENIA GRAVIS	4x3x1.2	A		BIOPSY
131794/18	55M	55M	M	DYSPOEA	11.5x2.4	B3		EXCISION
143362/18	55M	55M	M	COUGH	9.5x5x3	B1		EXCISION
152834/20	58M	58M	M	WEAKNESS	5x4x1.5	AB		EXCISION
161139/21	68M	68M	M	INCIDENTAL	10.5x8x4.5	AB	Ttf1	EXCISION
17300/21	52F	52F	F	COUGH	13x7.2x1.9	A	P63,CD33,CK7,AE1	BIOPSY
18889/21	64F	64F	F	INCIDENTAL	3.8x3.5x2	B2	PAN CK, CD20	BIOPSY
191294/21	63M	63M	M	CHEST PAIN	9x7.8x3.2	B3		EXCISION
201384/21	36M	36M	M	MYASTHENIA GRAVIS	4.5x4.5x1	B2		BIOPSY
214564/21	55M	55M	M	COUGH	19x12	AB		EXCISION
224444/21	42F	42F	F	CHEST DISCOMFORT	14.5x10.5x6	AB		EXCISION
23298/22	38F	38F	F	MUSCLE WEAKNESS	10x7x3	AB		EXCISION
244305/23	73F	73F	F	COUGH	2.3x2.3x1.9	A		BIOPSY
253412/23	38M	38M	M	COUGH	6x5.6x2	A	PANCK,CD3,Ttf1,AE1	EXCISION

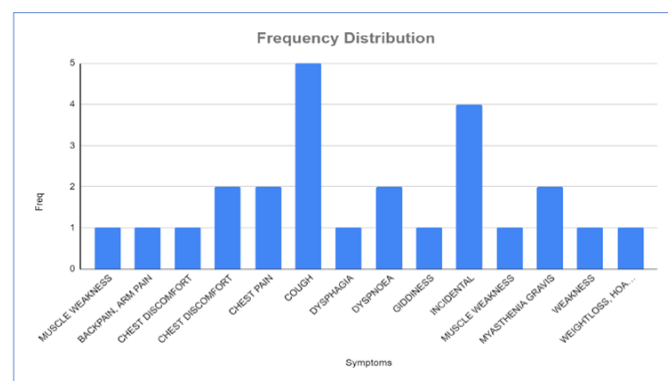
Graph 3:



Graph 4:



Graph 5:



The symptoms reported by the patients vary, with "Cough" being the most common, occurring in 20% of the cases. Other notable symptoms include "Incidental" findings (16%), "Chest Pain" (8%), "Chest Discomfort" (8%), "Dyspnoea" (8%), and "Myasthenia Gravis" (8%).

Results

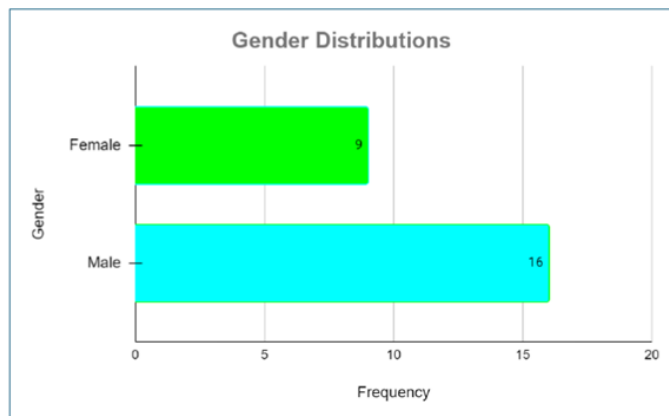
Among 25 thymic neoplasm cases, male predominance was observed which consist of 16 cases (64%) with a male-to-female ratio of 1.78:1

The most affected age group is 61-80 years (36%)

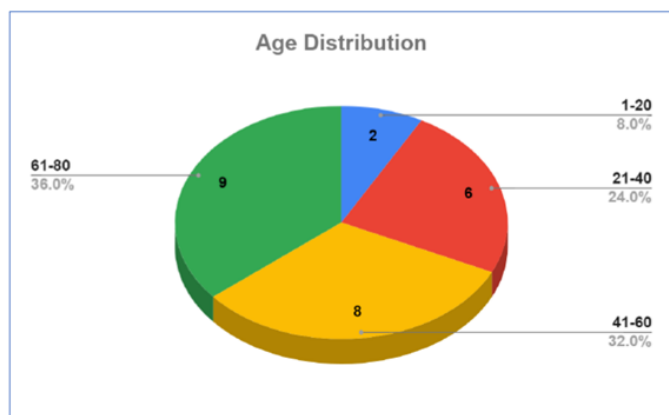
Thymomas constitute majority of the diagnoses followed by atypical thymic carcinoid tumors.

Myasthenia gravis was associated with 2 (8%) cases

Graph 1:



Graph 2:



Discussion

Thymic neoplasms are most common mass of anterior mediastinum. It frequently exhibit histologic heterogeneity, making it difficult to distinguish between benign and malignant tumours only on the basis of their medical history³

In our study, slight male predominance was observed which was consistent with literature⁴

Nearly 56% patients had compressive symptoms and two cases had history of myasthenia gravis which is commonly associated with thymoma. In our study, thymoma constituted majority (88%) of the cases followed by exceedingly rare tumors like atypical thymic carcinoid tumor (8%) and Thymic carcinoma (4%). Two cases of atypical carcinoid tumors presented with non specific symptoms. Similar observation was appreciated by shan Zhu⁵. Histology showed tumour cells in classical organoid pattern and nests separated by fibrovascular septa with occasional mitosis.

Extremely rare and aggressive variant of thymic carcinoma noted in our study was primary thymic mucinous adenocarcinoma which presented with large mediastinal mass extending to middle and posterior compartments. IHC results: Positive: CD20, CDX2 and CD5 Ki-67 labelling index: 40-50%, Negative: CK7, TTF1, Napsin A, GCDFP-15. This was similar to a study by Yu Zhang⁶

Most common histological variant among thymoma was Type AB followed by Type A, B3, B1 and B2

Type AB thymomas combine both a lymphocyte-rich (type B-like) and a lymphocyte-poor (type A-like) component.

Type A showed predominantly spindle and oval shaped cells with bland nuclei, finely dispersed chromatin and inconspicuous nucleoli with no lymphocytes.

Another histological subtype of thymoma noted was micronodular thymoma showing well-formed fibrous capsule with nodules of epithelial cells, elongated nuclei and scant cytoplasm.

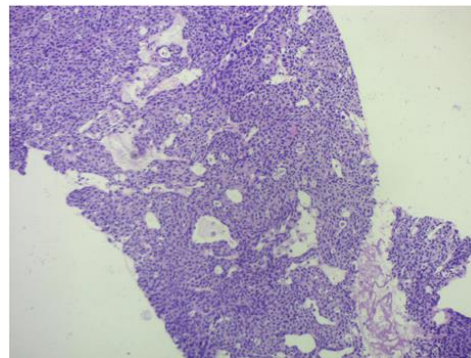


Figure 1: Thymoma Type A (H&E, 4x magnification)

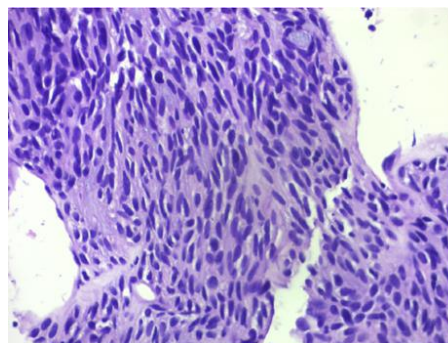


Figure 2: Thymoma Type A (H&E, 40x magnification)

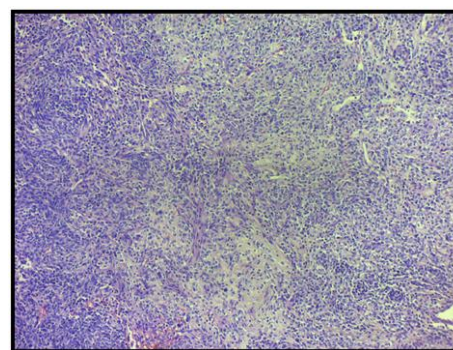


Figure 3: Type AB Thymoma (H&E, 10X magnification)

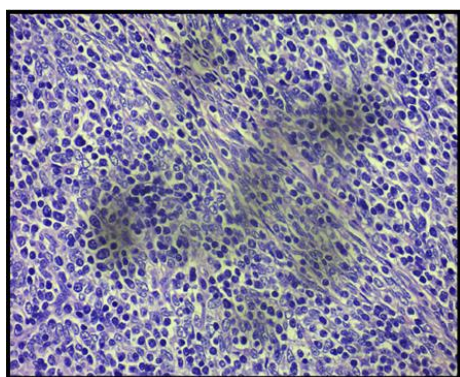


Figure 4: Type AB Thymoma (H&E, 40x magnification)

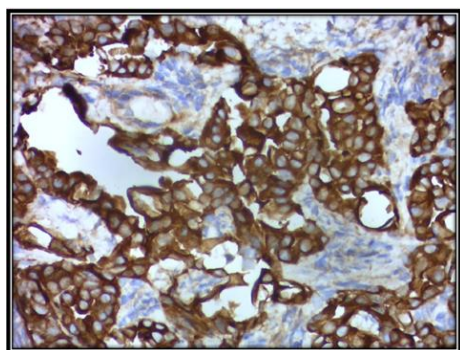


Figure 5: Pan CK (40x magnification)

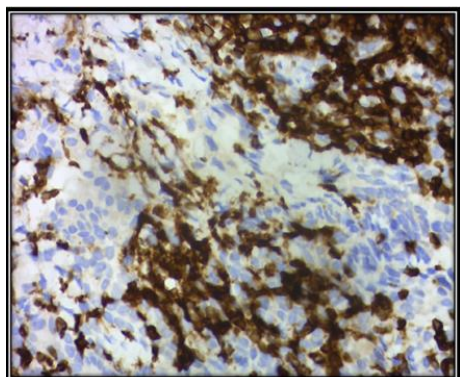


Figure 6: CD3 (40x magnification)

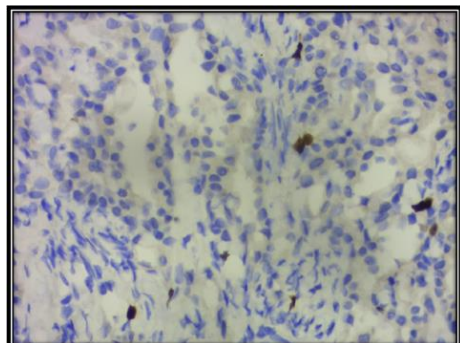


Figure 7: KI 67 (40x magnification)

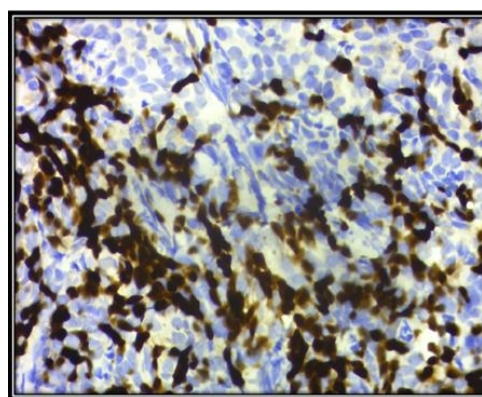


Figure 8: TDT (40x magnification)

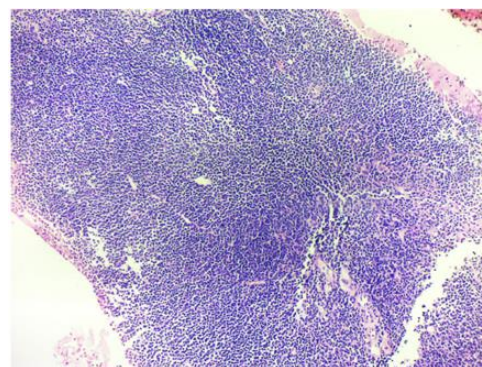


Figure 9: B2 thymoma (H&E, 10x magnification)

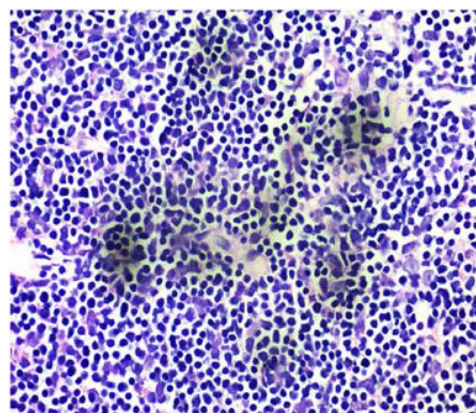


Figure 10: B2 thymoma (H&E, 40x magnification).

Conclusion

Histopathology in thymic neoplasm is an important diagnostic tool to know the spectrum of various thymic lesions, in understanding the pathogenesis, to assess the various morphological pattern of the same disease, before multimodality therapy is started. IHC can be done to distinguish classification of thymoma and thymic

carcinomas and are indispensable in academic and research purposes.

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