VOL15, ISSUE 02, 2024

HISTOPATHOLOGICAL SPECTRUM OF THYROID LESIONS IN TERTIARY CARE CENTER – A DESCRIPTIVE CROSS SECTIONAL STUDY

Rubina Hitawala¹, Praveen Rachakatla², Prakash Roplekar³, C P Bhale⁴

¹Senior Resident, Department of Pathology, MGM medical college and Hospital, Chhatrapati Sambhajinagar, Maharashtra, India.

²Assistant Professor, Department of Pathology, MGM medical college and Hospital, Chhatrapati Sambhajinagar, Maharashtra, India.

³Retired Professor and Head of Department, Department of pathology, Padmashree Dr. D.
Y. Patil Medical College, Hospital and Research Centre, Navi Mumbai, Maharashtra, India.
⁴Professor and Head of Department, Department of Pathology, MGM medical college and Hospital, Chhatrapati Sambhajinagar, Maharashtra, India.

Corresponding Author: Dr Rubina Hitawale, 403, B1, Pride Phoenix Apartment, MIDC

Area, Chikalthana, Chhatrapati Sambhajinagar, Maharashtra, India.

Email: ruby.hita.91@gmail.com

Abstract

Background: Thyroid carcinoma closely resembles its benign counterpart in physical characteristics, measurable physiological parameters such as serum T3/T4 levels and ultrasound features. Therefore, the surgical excision of the nodule and its histological examination is important to differentiate between the more frequent benign and much less frequent malignant nodules. The highest incidence of thyroid lesions was seen between the 3rd and 4th decades of life with female preponderance. Colloid goiter was overall the most common lesion seen. Papillary carcinoma was the most common malignant lesion. Familiarity with the diverse morphology of neoplastic lesions of thyroid helps in providing quality care for patients as they have prognostic significance. Consequently, the diagnosis and management of thyroid lesions is dependent on both the clinician and pathologist for better patient care. Therefore, detailed histopathological analysis of thyroid lesions contributes significantly to the diagnosis, management and prognosis of the patient as small lesions in thyroid may present with dual pathology which can be picked up on detailed grossing and microscopic examination for accurate diagnosis and management.

Keywords: Thyroid gland, papillary carcinoma, colloid goiter, thyroid function tests, histopathology.

Introduction

Thyroid lesions are common worldwide and are routinely encountered in clinical practice. Thyroid lesions may be developmental, inflammatory, hyperplastic or neoplastic.

Thyroid carcinoma closely resembles its benign counterpart in physical characteristics, measurable physiological parameters such as serum T3/T4 levels and ultrasound features. Therefore, the surgical excision of the nodule and its histological examination is important to differentiate between the more frequent benign and much less frequent malignant nodules.

VOL15, ISSUE 02, 2024

There are different diagnostic modalities used to evaluate and diagnose efficiently thyroid nodules. These include clinical examination, thyroid function tests (TFT), ultrasonography (USG), fine needle aspiration cytology (FNAC) and histopathological examination. However clinical assessment, TFT and USG have been poor parameters for assessing thyroid nodules. Final diagnosis requires morphological examination of lesions and for this FNAC or histological examination becomes mandatory.

Despite many advantages, FNAC has certain limitations which include specimen adequacy, cytological interpretation, as the sampling is variable and not always representative, and lesions may be small in enlarged thyroid glands and may not be sampled by FNAC.

Hence, histopathological examination remains gold standard for diagnosis.

Materials and Method

This was a prospective study of all the patients who presented with thyroid lesions in Department of Surgery, D. Y. Patil Hospital.

All clinical details, laboratory tests, FNAC in cases which were subjected for evaluation along with USG/CT/MRI Radiological investigations were included in the present study.

A total of 128 cases were studied over a span of two years from September 2017 to September 2019 were included in the study after clearance from Institutional Ethical Committee.

Inclusion Criteria

All Thyroid resected specimens from surgery department for histopathological examination in the Department of Pathology, D. Y. Patil School of Medicine.

Exclusion Criteria

Congenital anomalies including thyroglossal cysts were excluded from the study.

- The specimens were properly labelled, and their detailed gross examination was done followed by fixation in 10% formalin for 24 48 hours.
- Representative sections were taken.
- The tissue sections were processed for paraffin block preparation and sections of 4-5 µm thickness were taken which then underwent hematoxylin and eosin staining and studied under the microscope.

The data was compiled, and statistical analysis was done.

The present study was further compared with other similar studies.

Results

The highest incidence of thyroid lesions was seen between the 3rd and 4th decades of life (78.91%) while the lowest was seen in 5th and 6th decade of life. (Table 1&2) (Figure 1 & 2)

Table 1: Mean Age (yrs.) of the patients in benign and malignant thyroid lesions

	Benign lesions	Malignant lesions
N	104	24
Mean age	42.44	40.25
Standard deviation	11.29	14.25
95% CI for the mean	40.25 to 44.64	34.23 to 46.27

VOL15, ISSUE 02, 2024

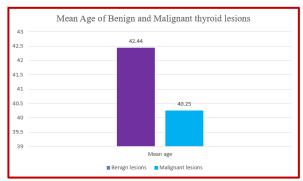


Figure 1: Mean age of the patients in benign and malignant thyroid lesions (%)

Table 2: Age-wise prevalence of thyroid lesions

Tuble 2. Fige wise prevalence of myrota resions						
	Benign (n=104)		Malignant (n=24)		Tota	l (n=128)
	No.	%	No.	%	No.	%
<= 20 yrs.	12	11.54%	5	20.83%	17	13.28%
20 to <40 yrs.	24	23.08%	9	37.50%	33	25.78%
40 to <60 yrs.	62	59.62%	6	25.00%	68	53.13%
>= 60 yrs.	6	5.77%	4	16.67%	10	7.81%
Chi-square						
test						
χ2	10.204					
P	0.0169					

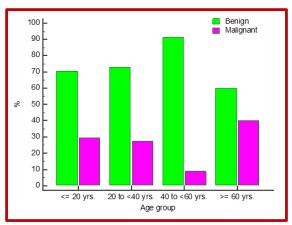


Figure 2: Age distribution of the patients (%)

Female to male ratio was found to be 1:5.5. (Table 3) (Figure 3.1 & 3.2)

Table 3: Gender distribution of the patients in benign and malignant thyroid lesions

Gender	Benign (n=104)	Malignant	Total (n=128)
		(n=24)	
	No. (%)	No. (%)	No. (%)
Male	14 (77.8%)	4 (22.2%)	18 (14.1%)
Female	90 (81.8%)	20 (18.2%)	110 (85.9%)
Total	104 (81.2%)	24 (18.8%)	128 (100.0%)
Chi-square test	Benign vs Malignant		
χ2	0.164		
P	0.6851		

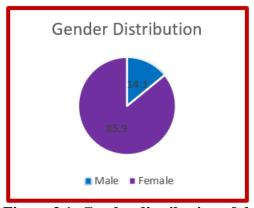


Figure 3.1: Gender distribution of the patients in benign and malignant lesions (%)

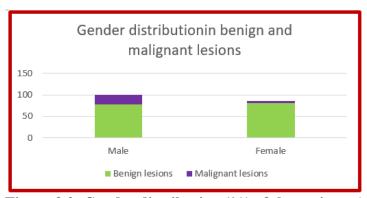


Figure 3.2: Gender distribution (%) of the patients (stacked)

Of the 128 thyroid specimens studied, 16 cases were non-neoplastic lesions and 88 and 24 were benign and malignant neoplastic thyroid lesions respectively. (Table 4) (Figure 4)

Table 4: Benign thyroid lesions NIFTP – Non-invasive Follicular Thyroid Neoplasm with Papillary-Like Nuclear Features

	No.	% of	% of
		benign lesions	total lesions
		(n=104)	(n=128)
Colloid goitre	35	33.65%	26.92%
Multinodular colloid Goitre	23	22.12%	17.69%
Follicular Adenoma	16	15.38%	12.31%
Hashimoto's thyroiditis	10	9.62%	7.69%
Lymphocytic Thyroiditis	6	5.77%	4.62%
NIFTP	6	5.77%	4.62%
Adenomatoid Goitre	3	2.88%	2.31%
Hurthle Cell Adenoma	2	1.92%	1.54%
Hyalinizing Trabecular	1	0.96%	0.77%
Tumor			
Papillary Hyperplastic	1	0.96%	0.77%
Nodule			
Thyroid Hamartoma	1	0.96%	0.77%

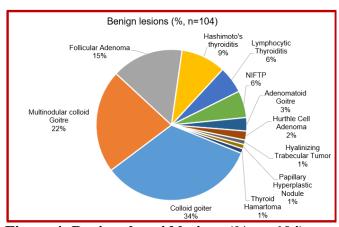


Figure 4: Benign thyroid lesions (%, n=104)

Colloid Goiter (26.92%) was the commonest lesion seen in the present study.

Among the non-neoplastic lesion of thyroid, Hashimoto's Thyroiditis was the most commonly occurring entity. (Table 4) (Figure 4)

Among the Neoplastic lesions of thyroid, Colloid goiter was the most commonly occurring entity followed by Multinodular goiter in benign category and Papillary Carcinoma of thyroid occurred most in malignant category. (Table 5) (Figure 5)

Table 5: Malignant thyroid lesions

	No.	% of malignant	% of total
		lesions	lesions
		(n=24)	(n=128)
Papillary Carcinoma	12	50.00%	9.23%
Follicular variant of Papillary Ca (FVPTC)	8	33.33%	6.15%
Medullary Carcinoma	1	4.17%	0.77%
Poorly Diffrentiated Thyroid Carcinoma	1	4.17%	0.77%
Follicular Carcinoma	1	4.17%	0.77%
Follicular Tumor of Uncertain Malignant	1	4.17%	0.77%
Potential (FTUMP)			

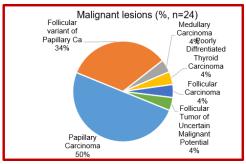


Figure 5: Malignant thyroid lesions (%, n=24)

T3 T4 and TSH values showed no association with occurrence of benign or malignant lesions and hence had no positive predictive value, (p value = 0. 3425, 0.4856 and 0.2215 respectively.)

Most of the lesions were found to be Euthyroid (93.8%). (Table 6)

Table 6: Functional status of thyroid in 128 cases.

Thyroid function tests	No. of cases	Percent of cases
------------------------	--------------	------------------

Journal of Cardiovascular Disease Research

ISSN: 0975-3583,0976-2833

VOL15, ISSUE 02, 2024

Euthyroid	120	93.8%
Hypothyroid	2	1.6%
Hyperthyroid	6	4.6%

Table 7: Comparison of Age-wise distribution of Thyroid lesions

	< 20 years	20 –39 years	years (% of	> 60 years (% of cases)
	(% of cases)	(% of cases)	cases)	
Singh P et al. (6) (2000) (n = 108)		62.4	20	13.6
Md R. Islam <i>et al.</i> (2009) (n	3.3	81.3	15.4	00
=118)				
Aahana Gupta <i>et al.</i> (1) (2016) (n	12	51	31	06
= 100)				
A. Kumar et.al (7) (2017)	9.1	55.2	28.8	06.9
(n=248)				
Present Study ($n = 128$)	13.28	25.78	53.13	7.81

Table 8: Sex predilection of thyroid lesions

Tuble of Sea predicetion of thy	Female (%)	Male (%)
Singh P et al. (6) (2000) (n = 92)	82.4	17.6
Md R. Islam <i>et al</i> ⁽²⁾ (2009) (n =	67.7	32.3
118)		
Aahana Gupta <i>et al.</i> (2016) (n	77	23
= 100)		
A. Kumar et.al $^{(7)}$ (2017) (n =	85.1	14.9
248)		
Present Study (n = 128)	85.9	14.1

Table 9: Comparison of Histopathological spectrum of thyroid lesions

	Benign (%)	Malignant (%)
Singh P et al. (6) (2000) (n = 92)	86.9	13.1
Md R. Islam <i>et al.</i> ⁽²⁾ (2009) (n =	81.4	18.6
118)		
Aahana Gupta <i>et al.</i> ⁽¹⁾ (2016) (n	72	28
= 100)		
A. Kumar et.al $^{(7)}$ (2017) (n =	81.01	18.99
248)		
Present Study (n = 128)	81.2	18.8

Table 10: Comparison of thyroid function test and thyroid lesions

tube 10. Comparison of mytola function test and mytola lesions				
Thyroid hormone profile	Euthyroid	Hypothyroid	Hyperthyroid	
(in percentage)				
Godinho- Matos L <i>et al.</i> ⁽⁸⁾ (1992) (n =	88	3	9	
144)				
Md R. Islam <i>et al.</i> ⁽²⁾ (2009) (n = 118)	100	00	00	
Aahana Gupta <i>et al.</i> ⁽¹⁾ (2016) (n =	81	08	11	
100)				

VOL15, ISSUE 02, 2024

A. Kumar et.al ⁽⁷⁾ (2017) (n = 221)	85.07	9.5	5.43
Present Study	93.8	1.6	4.6

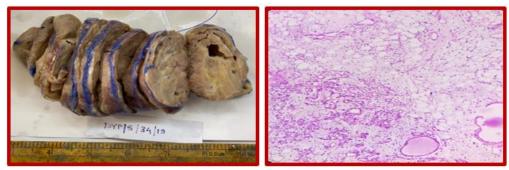


Figure. 6: Thyroid Hamartoma A. Gross B. Microscopy

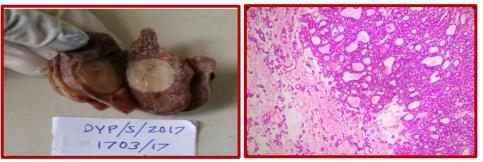


Figure 7: Follicular Adenoma A. Gross B. Microscopy

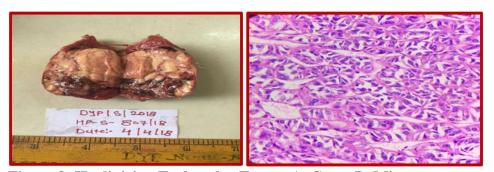


Figure 8: Hyalinizing Trabecular Tumor A. Gross B. Microscopy

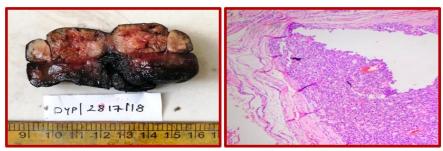


Figure 9: Invasive Encapsulated Follicular Variant of Papillary Carcinoma A. Gross B. Microscopy

VOL15, ISSUE 02, 2024

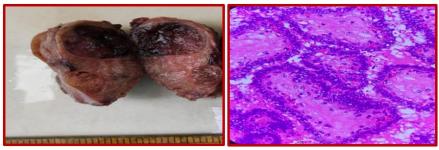


Figure 10: Papillary Carcinoma A. Gross B. Microscopy

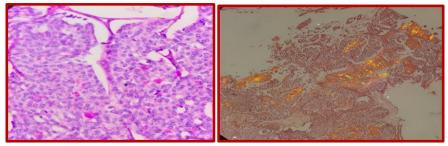


Figure 11: Medullary Carcinoma microscopic pictures - A. H&E stain B. Congo Red Stain

Discussion

In the present study, the highest incidence of thyroid lesions was seen between the 3rd and 4th decades of life (78.91%) while the lowest was seen in between 5th and 6th decade of life. (Table 7)

These findings compare favorably with the findings of Aahana Gupta *et al.*⁽¹⁾. However, Md R. Islam *et al.*⁽²⁾ found a higher incidence of cases in the 4th decade of life itself.

There is lower incidence of thyroid lesions in fifth and sixth decade of life in all studies including the present one and hence younger age group should be evaluated for thyroid lesions much more than older age group. (Table 8)

In this series, out of 128 patients, female (104) outnumbered the males with female-male ratio of 5.5:1. This female preponderance is reflected in all other studies also.

Studies done by Sangall G, *et al.* 2006⁽³⁾, female:male was 4.21:1, Mandal S, *et al.* ⁽⁴⁾ female:male was 5:1. Kilopatric, *et al.* ⁽⁵⁾ found a female to male ratio of 4:1 in nonendemic area. It is due to fact that thyroid disorder is female prone owing to the presence of estrogen receptors in the thyroid tissue.

The present study compared favorably with all the studies when comparing the percentage of benign and malignant cases of thyroid. (Table 9)

Md R. Islam *et al.*⁽²⁾, Singh P *et al.*⁽⁶⁾ and A. Kumar et.al⁽⁷⁾ showed similar percentage of benign lesions while Aahana Gupta *et al.*⁽¹⁾ showed a slightly lesser benign lesions.

However, overall these figures indicate the higher frequency of occurrence of benign lesions of thyroid and the importance of histopathology in identifying them.

All cases of our study underwent thyroid hormone profile and found to be, Euthyroid 93.8%, Hyperthyroid 4.6% and 1.6% Hypothyroid. In the study by Godlnho-Matos L, *et al.* there were 144 cases out of them 88 cases were euthyroid, 9 cases were hyperthyroid and 3 cases were hypothyroid ⁽⁸⁾ (Table 10). Also, in a study by Kraiem Z *et al.* multinodular goiter can be associated with hyperfunction⁽⁹⁾, but the patient is usually found to be euthyroid.

Hence, TFTs do not shed much light on the nature of the lesion and a thyroid mass should be evaluated even in case of normal thyroid profile.

Journal of Cardiovascular Disease Research

ISSN: 0975-3583,0976-2833

VOL15, ISSUE 02, 2024

Most common non neoplastic thyroid lesion was colloid goitre (Nodular Hyperplasia) followed by multinodular Colloid goitre.

Most common malignant thyroid lesion seen in the present study was Papillary Carcinoma of thyroid (15.38%) including its follicular variant among all lesions occurring in thyroid. This is in accordance with a study done by Sherman SI *et al.*⁽¹⁰⁾ and Jemal A *et al.*⁽¹¹⁾ which found Papillary Carcinoma as the most common malignant tumor of the thyroid gland in countries having iodine-sufficient or iodine-excess diets. (10) These common tumors tend to be biologically indolent and have an excellent prognosis (90% survival at 20 years) (10).

Rare cases like thyroid hamartoma and anaplastic carcinoma of thyroid, one each, were also reported and show the extremes of the spectrum.

All the facts and figures mentioned here may considerably vary from those of large series covering wide range of time, but still then, as the cases of this study are collected from a tertiary level hospital in our country, this study has some credentials in reflecting the facts regarding distribution and type of various thyroid lesions.

Conclusion

The incidence of diseases involving the thyroid gland has been increasing over the years. However, early diagnosis can assist in early treatment and have better outcomes.

The availability of newer diagnostic modalities has increased the detection rate of early presenting patients. However, clinical examination, thyroid function tests, radiological examination and fine needle aspiration cytology remain only provisional diagnostic modalities. Histopathology remains the gold standard not only for the confirmation of clinical diagnosis but also for planning the management and predicting the prognosis.

Histopathology holds a prime significance in malignant lesions not only to identify and grade the tumors but also to pick up lesions that are treated as benign, as there is an overlap in the clinical features of both.

A wide spectrum of diseases can be diagnosed on histopathology, as they may be too small to be detected on clinical presentation or radiology. Therefore, every resected specimen should be subjected to a detailed histopathological examination for a surgico-pathological correlation and to ensure proper management of the patient.

Our study highlights the fact that the thyroid is a site for more than one pathology and histopathologists should strive to identify and diagnose each of these lesions.

Grading and staging are important, and pathologists should ensure that they use standard criteria which clinicians can interpret correctly.

Familiarity with the diverse morphology of neoplastic lesions of thyroid helps in providing quality care for patients as they have prognostic significance. Consequently, the diagnosis and management of thyroid lesions is dependent on both the clinician and pathologist for better patient care.

Therefore, detailed histopathological analysis of thyroid lesions contributes significantly to the diagnosis, management and prognosis of the patient as small lesions in thyroid may present with dual pathology which can be picked up on detailed grossing and microscopic examination for accurate diagnosis and management.

Acknowledgment- None. Conflict of Interest - None

VOL15, ISSUE 02, 2024

References

- 1. Gupta A, Durgashankar Jaipal, Sunita Kulhari *et al.* Histopathological study of thyroid lesions and correlation with ultrasonography and thyroid profile in western zone of Rajasthan, India. Int J Res Med Sci. 2016 Apr;4(4):1204-1208
- 2. Islam R, Ekramuddaula AFM, Alam MS, Kabir MS, Hossain D, Alauddin M. Frequency & pattern of malignancy in solitary thyroid nodule. Bangladesh J of Otorhinolaryngology. 2009;15(1):1-5.
- 3. Sangalli G, Sergio G, Zampatti C, Bellotti M, Lomuscio G. Fine needle aspiration cytology of the thyroid: A comparison of 5469 cytological and final histological diagnosis. Cytopathology. 2006;17(5):245-50.
- 4. Mandal S, Barman D, Mukherjee A, Mukherjee D, Saha J, Sinha R. Fine needle aspiration cytology of thyroid nodules-evaluation of the role in diagnosis and management. J Indian Med Assoc. 2011;109(4):258-61.
- 5. Ashraf SA, Matin ASM. A Review of thyroid diseases in Bangladesh. Journal of BCPS. 1996;2(1):6-10.
- 6. Singh P, Chopra R, Calton N, Kapoor R. Diagnostic Accuracy of Fine Needle Aspiration Cytology of Thyroid lesions. Journal of Cytology. 2000;17(3):135-9.
- 7. Kumar A *et al.* Comparative study of FNAC and histopathology of thyroid swellings, diagnostic accuracy and role in its management Int J Otorhinolaryngol Head Neck Surg. 2017 Oct;3(4):885-892
- 8. Godinho-Matos L, Kocjan G, Kurtz A. Contribution of fine needle aspiration cytology to diagnosis and management of thyroid disease. J Clin Pathol. 1992;45(5):391-5
- 9. Kraiem Z, Glaser B, Yigla M, *et al.* Toxic multinodular goiter: a variant of autoimmune hyperthyroidism. J Clin Endocrinol Metab 1987;65:659–664.
- 10. Sherman SI, Angelos P, Ball DW, et al. Thyroid carcinoma. JNatl Compr Canc Netw 2007;5:568–621.
- 11. Jemal A, Siegel R, Ward E, et al. Cancer statistics, 2007. CA Cancer J Clin 2007;57:43–66.