Spectrum Analysis of Thyroid Lesions on Histopathology

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ABSTRACT

Introduction: Thyroid gland is an endocrine organ with two lobes and connecting isthmus, located below and anterior to the larynx. Histologically, the thyroid gland is comprised of follicles lined by follicular epithelial cells with eosinophilic colloid in the lumen.

Objective: This study aimed to analyse thyroid lesions histopathologically.

Methods: Specimens of thyroid lesions received at Department of Pathology, Government Medical College, Srinagar from January 2018 to December 2019 were studied retrospectively.

Results: Among the 180 specimens of thyroid lesions, 145 were from female patients and 35 from male patients. Age of the patients ranged from 4 years to 66 years. 31 cases (17%) were found to be non-neoplastic and 149 cases (83%) were neoplastic. Of the non-neoplastic lesions, 28 cases were documented as colloid goitre, 2 cases as a thyroglossal cyst and 1 case as Hashimoto thyroiditis. Of the neoplastic category, benign neoplasms (n=49) included 34 cases of follicular adenoma, 12 cases of Hurthle cell adenoma and 3 cases of non-invasive follicular thyroid neoplasm with papillary like nuclear features malignant category (n=100) included 90 cases of papillary thyroid carcinoma, 2 cases of follicular thyroid carcinoma and 8 cases of medullary thyroid carcinoma.

Conclusion: In our study, only 17% of thyroid surgeries were done for non-neoplastic lesions. Further reduction in the number of thyroidectomies for non-neoplastic lesions is expected with improved applicability of preoperative imaging and fine-needle aspiration cytology.

Key Words: Thyroid, Histopathology, Neoplastic, Papillary Thyroid carcinoma

INTRODUCTION

Thyroid gland is an endocrine organ with two lobes and connecting isthmus, located below and anterior to the larynx. Histologically, the thyroid gland is comprised of follicles lined by follicular epithelial cells with eosinophilic colloid in the lumen. C-cells are interspersed within the follicular epithelium with basement membrane investing both follicular as well as C-cells. Common diseases of thyroid encountered in histopathological specimens include developmental (pyramidal lobe, thyroglossal duct cyst, lingual thyroid), inflammation (thyroiditis), thyroid hyperplasia (goitre), and neoplasms. The updated classification of thyroid neoplasms (Table 1) has been introduced in the 4th Edition of the WHO classification of endocrine organs in 2011. The precise classification of thyroid neoplasms has always been onerous owing to the evolution of molecular/genetic characterisation of these tumours, introduction of new entities, reclassification of previous entities and updates in diagnostic criteria. This study was undertaken to thoroughly analyse thyroid lesions and thereby establishing the histopathological spectrum.

MATERIALS AND METHODS

This was a retrospective study conducted at the department of pathology, Government Medical College, Srinagar. The records of all thyroid specimens received in the department over 2 years (January 2018 to December 2019) were acquired.
from departmental registry and histopathology requisition forms. The clinical history including age, gender, symptoms, radiological and cytologic findings were noted. The slides were retrieved from the departmental gallery and wherever necessary, fresh 4-5micron sections were cut on a microtome and stained by Hematoxylin and Eosin stain. All the cases were studied by more than one pathologist. The data were entered in Microsoft Excel spreadsheet and was then imported into statistical package for social science (SPSS) version 20 for statistical analysis.

RESULTS

In the two years from January 2018 to December 2019, a total of 180 specimens of thyroid lesions designated thyroidectomy, lobectomy, partial excision and Sistrunk operation have been received in our department. Overall female patients outnumbered male with F: M ratio of 4.14:1 (Figure 1, Table 2). The maximum lesions predominated in the age group of 21-40 years (n=102) (Table 3).

The non-neoplastic lesions constituted 17% (31 cases) of all thyroid lesions (Figure 2). Colloid goitre was the most common non-neoplastic lesion (28/31, 90.32%). Of these cases, 25 were female (89.28%) and 3 were male (10.71%). Two cases (6.45%) of the thyroglossal cyst and 1 case (3.22%) of Hashimoto thyroiditis were found (Figure 3). In this study, 149 cases (83%) belonged to the neoplastic category. Of these cases, 49 (32.88%) were benign and 100 (67.11%) were malignant. In the benign category, 34 cases of follicular adenoma, 12 cases of hurthle cell adenoma and 3 cases of Non-Invasive Follicular Thyroid neoplasm with Papillary like nuclear features (NIFTP) were found. The female to male ratio was 5.1:1 and the majority cases were in the age group of 21-50 years (Figure 4, 5).

Among the malignant thyroid neoplasms, papillary thyroid carcinoma, PTC was found to be the most common malignant thyroid lesion (90 cases). Of these cases, 71 were female and 19 were male with F: M ratio of 3.7:1 and most of the patients was in 21-40 years of age group (n=56). Two cases of follicular carcinoma (minimally invasive) and 8 cases of medullary carcinoma were documented (Figure 4). Conventional/classic type percutaneous tranhepatic cholangiography (PTC) constituted 84.44% (76/90) of PTC cases. Also, 6 variants were identified including papillary microcarcinoma (5 cases), invasive follicular variant (2 cases), tall cell variant (3 cases), diffuse sclerosing variant (2 cases), columnar cell variant (1 case) and oncocytic variant (1 case) (Figure 5). One of the two cases of the diffuse sclerosing variant was found in association with the thyroglossal cyst.

Photomicrographs of some of the thyroid lesions are shown as hurthle cell adenoma (Figure 6), papillary microcarcinoma (Figure 7), a diffuse sclerosing variant of PTC (Figure 8), a tall cell variant of PTC (Figure 9) and medullary thyroid carcinoma (Figure 10).

DISCUSSION

Thyroid lesions are common all over the world and the histopathological spectrum of these lesions vary according to the geographical region, race, age and gender of the studied population. In concordance with the previously published literature, our study demonstrated that females are more commonly affected than males with the majority of the lesions documented in the age group of 21-40 years.

Both neoplastic and non-neoplastic lesions of thyroid are found in this part of the world and we found that neoplastic lesions are more common than non-neoplastic ones in the thyroid specimens sent for histopathological examination. Majority of other authors reported a higher percentage of non-neoplastic lesions, constituting 72.3%, 82%, 83.3% and 84% of thyroid lesions respectively in the studies by Albasri et al. Sreedevi et al., Fatima et al. Prabha et al. In a study by Ramesh, neoplastic lesions (52.5%) marginally exceeded the non-neoplastic lesions (47.5%). This deviation from the previously published literature could be because of the applicability of fine-needle aspiration cytology and increased expertise of cytopathologists in accurately diagnosing thyroid lesions, thereby reducing the burden of unnecessary surgeries for non-neoplastic lesions. Also, the increased detection of malignant lesions with the introduction of high-resolution imaging techniques may be responsible for the increased incidence of neoplastic lesions in our study.

Among the non-neoplastic lesions, colloid goitre was the most common lesion in our study similar to the previous studies. The low frequency of Hashimoto thyroiditis in our study was also noted by other authors. Of the neoplastic lesions, the most common benign tumour reported was follicular adenoma, in consonance with literature. All cases of follicular adenoma showed a single lesion except for one where multiple lesions were identified in right lobe and isthmus. Malignant lesions were found to be more common than benign paralleling the earlier studies.

The most common malignant thyroid tumour is PTC, constituting 85-90% of all differentiated thyroid cancers and females are affected four times than males. PTC was the commonest malignant lesion (91.83%) in our study supporting the increasing trend of this carcinoma documented in studies by other author. The F: M ratio for PTC in our study was 3.7. Similar to earlier studies, majority of the PTC were of conventional type (84.44%). Tall cell variant, diffuse sclerosing variant and columnar cell variant are acknowledged by WHO classification of thyroid tumours as biologically aggressive variants, all of which showed up in

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In our study, the majority of thyroid lesions and more than half cases were observes. Females constituted the majority of thyroid lesions and more than half cases were observed in the age group of 21-40 years. Colloid goitre was the most common non-neoplastic lesion, follicular adenoma, the most common benign tumour and papillary thyroid carcinoma, the most common malignant lesion.

**CONCLUSION**

This study points to the evolving histopathological spectrum of thyroid diseases with neoplastic lesions showing up as major chunk in thyroidectomy specimens. Females constituted the majority of thyroid lesions and more than half cases were observed in the age group of 21-40 years. Colloid goitre was the most common non-neoplastic lesion, follicular adenoma, the most common benign tumour and papillary thyroid carcinoma, the most common malignant lesion.

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Table 1: WHO classification of thyroid tumors

1. Follicular Adenoma
2. Hyalinising trabecular tumor
3. Other Encapsulated follicular patterned tumours:
   i. Follicular Tumor of Uncertain Malignant Potential
   ii. Well Differentiated Tumor of Uncertain Malignant Potential
   iii. Non-invasive follicular thyroid neoplasm with papillary like nuclear features
4. Papillary thyroid carcinoma
   i. Conventional/classic
   ii. Papillary microcarcinoma
   iii. Encapsulated
   iv. Follicular
   v. Diffuse sclerosing
   vi. Tall cell
   vii. Columnar cell
   viii. Cribriform-morular
   ix. Hobnail
   x. Papillary thyroid carcinoma with fibromatosis/fascitis-like stroma
   xi. Solid/trabecular
   xii. Oncocytic
   xiii. Spindle cell
   xiv. Clear cell
   xv. Warthin like the variant
5. Follicular thyroid carcinoma
   i. Follicular thyroid carcinoma, minimally invasive
   ii. Follicular thyroid carcinoma, encapsulated
   iii. Follicular thyroid carcinoma, widely invasive
6. Hurthle cell tumours
   i. Hurthle cell adenoma
   ii. Hurthle cell carcinoma
7. Poorly differentiated thyroid carcinoma
8. Anaplastic thyroid carcinoma
9. Squamous cell carcinoma
10. Medullary thyroid carcinoma
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Figure 4: Neoplastic lesions of thyroid.

Table 2: Gender distribution of thyroid lesions

| Histologic type         | Male | Female | Total |
|-------------------------|------|--------|-------|
| Thyroglossal duct cyst  | 2    | 0      | 2 (1.11%) |
| Hashimoto thyroiditis   | 0    | 1      | 1 (0.55%) |
| Colloid goitre          | 3    | 25     | 28 (15.55%) |
| Follicular adenoma      | 6    | 28     | 34 (18.88%) |
| Hurthle cell adenoma    | 2    | 10     | 12 (6.66%) |
| NIFTP                   | 0    | 3      | 3 (1.66%) |
| Papillary carcinoma     | 19   | 71     | 90 (50%) |
| Follicular carcinoma    | 1    | 1      | 2 (1.11%) |
| Medullary carcinoma     | 2    | 6      | 8 (4.44%) |
| Total                   | 35   | 145    | 180   |

Table 3: Age distribution of thyroid lesions

| Age group (years) | Thyroglossal duct cyst | Hashimoto thyroiditis | Colloid goitre | Follicular adenoma | Hurthle cell adenoma | NIFTP | Papillary carcinoma | Follicular carcinoma | Medullary carcinoma | Total (180) |
|-------------------|------------------------|-----------------------|---------------|-------------------|----------------------|-------|---------------------|---------------------|---------------------|-------------|
| <10               | 2                      | -                     | -             | -                 | -                    | -     | -                   | -                   | -                   | 2           |
| 11-20             | -                      | -                     | 3             | 2                 | 2                    | 13    | -                   | -                   | -                   | 20          |
| 21-30             | -                      | 1                     | 10            | 9                 | 1                    | 3     | 19                  | 1                   | 2                   | 63          |
| 31-40             | -                      | -                     | 6             | 9                 | 2                    | -     | 19                  | 1                   | 2                   | 39          |
| 41-50             | -                      | -                     | 5             | 9                 | 4                    | -     | 10                  | 1                   | -                   | 29          |
| 51-60             | -                      | -                     | 3             | 4                 | 2                    | -     | 8                   | -                   | 3                   | 20          |
| 61-70             | -                      | -                     | 1             | 1                 | 1                    | -     | 3                   | -                   | 1                   | 7           |

Figure 5: Papillary carcinoma and its variants.

Figure 6: Hurthle cell adenoma (10X).
Figure 7: Papillary microcarcinoma (10X).

Figure 8: Papillary thyroid carcinoma, diffuse sclerosing variant (20X).

Figure 9: Papillary thyroid carcinoma, tall cell variant (40X).

Figure 10: Medullary thyroid carcinoma (40X).