Introduction

Different types of thyroiditis may share some parallel clinical and biochemical features. Timely intervention can significantly reduce morbidity and mortality. Aim. Aim of this study is to find the frequency of various thyroiditis, study the cytomorphological features and correlate with clinical findings including radiological findings, thyroid function test, and anti-thyroid peroxidase antibodies (Anti-TPO antibodies). Materials and Methods. The study included consecutive 110 cases of thyroiditis. Detailed cytomorphological features were studied and correlated with ultrasonography findings, thyroid function test, anti-thyroid peroxidase antibodies (anti-TPO) and histopathological features where thyroidectomy specimens were received for histopathological examination. Results. The majority were Hashimoto’s thyroiditis (n = 100) and females (n = 103). Other forms of thyroiditis were Hashimoto’s thyroiditis with colloid goiter (n = 5), De Quervain’s thyroiditis (n = 3), and one case each of postpartum thyroiditis and Hashimoto’s thyroiditis with associated malignancy. The majority of patients were in the age group of 21–40 (n = 70) and the majority (n = 73) had diffuse enlargement of thyroid. The majority of patients were hypothyroid (n = 52). The serum anti-TPO antibodies were elevated in 47 patients out of 71 patients. In the 48 patients who underwent ultrasonography, 38 were diagnosed as having thyroiditis. The most consistent cytomorphological features seen in fine-needle aspiration smears of Hashimoto’s thyroiditis were increased background lymphocytes, lymphocytic infiltration of thyroid follicular cell clusters, and Hurthle cells. Conclusion. The diagnostic cytological features in Hashimoto’s thyroiditis are increased background lymphocytes, lymphocytic infiltration of thyroid follicular cell clusters, and Hurthle cells. FNAC remains the "Gold Standard" for diagnosing Hashimoto’s thyroiditis. Clinical history, thyroid function, and biochemical parameters are the key for diagnosis of other forms of thyroiditis.

1. Introduction

The prevalence of these thyroid disorders varies widely according to geographical distribution, diet and nutrition, and patient population [1, 2]. Thyroiditis a diverse group of disorder characterized by thyroid inflammation and have different etiologies [3]. It can be categorized as acute, subacute or chronic forms. The most common form of thyroiditis is chronic lymphocytic thyroiditis (Hashimoto’s thyroiditis [HT]) [4–9]. Other common types include postpartum thyroiditis (subacute lymphocytic thyroiditis), silent sporadic thyroiditis, subacuate granulomatous thyroiditis (De Quervain’s thyroiditis), supplicative thyroiditis, and fibrous thyroiditis (Riedel’s thyroiditis) [3].

Thyroiditis patients may be euthyroid, hyperthyroid, or hypothyroid and may evolve over the time [10]. Chronic high iodine intake has been associated in various studies with increased frequency of autoimmune thyroiditis [11]. It is well-known that HT can coexist with other lesions such as follicular neoplasm, Hurthle cell neoplasm, papillary carcinoma, and goitrous nodules [12–15]. Incidence of coexisting thyroid neoplasia with HT range between 3 and 14%. Coexistence of thyroiditis and colloid goiter is also seen in significant number of cases [12, 14, 16].
Table 1: Correlation of goiter, thyroid function, and anti-TPO in 110 cases of thyroiditis.

<table>
<thead>
<tr>
<th>Type of thyroiditis (𝑛 = number)</th>
<th>Goiter</th>
<th>Thyroid function</th>
<th>Anti-TPO titer</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Diffuse</td>
<td>Uneven</td>
<td>Hypo</td>
</tr>
<tr>
<td>Hashimoto thyroiditis (𝑛 = 100)</td>
<td>71</td>
<td>27</td>
<td>02</td>
</tr>
<tr>
<td>HT with colloid goiter (𝑛 = 5)</td>
<td>01</td>
<td>04</td>
<td>00</td>
</tr>
<tr>
<td>de Quervain thyroiditis (𝑛 = 3)</td>
<td>00</td>
<td>03</td>
<td>00</td>
</tr>
<tr>
<td>Postpartum thyroiditis (𝑛 = 1)</td>
<td>01</td>
<td>00</td>
<td>00</td>
</tr>
<tr>
<td>HT with malignancy (𝑛 = 1)</td>
<td>00</td>
<td>00</td>
<td>01</td>
</tr>
<tr>
<td>Total-I10</td>
<td>73</td>
<td>34</td>
<td>03</td>
</tr>
</tbody>
</table>

HT: Hashimoto's thyroiditis; SN: solitary nodule; Hyper: hyperthyroidism; Hypo: hypothyroidism; SCH: subclinical hypothyroidism; Euth: euthyroidism.

Different types of thyroiditis may share same clinical and biochemical features. Conventional ultrasonography is the most commonly applied modality in the evaluation of thyroiditis [17]. Fine-needle aspiration cytology is practiced worldwide and is the investigation of choice in thyroid enlargement [18–21]. This study has been undertaken to find out the frequency of various thyroiditis and study the cytomorphological features and correlate with clinical findings including radiological features, thyroid function test, and anti-thyroid peroxidase antibodies (Anti-TPO antibodies).

2. Materials and Methods

A prospective cohort study was carried out. The study includes consecutive 110 cases of thyroiditis. Inclusion criteria were thyroiditis patients diagnosed on fine-needle aspiration and or ultrasonography, thyroid function tests, and or anti-TPO antibodies. Exclusion criteria were noninflammatory and neoplastic lesions of thyroid.

All the patients underwent fine-needle aspiration (FNA) in cytology clinic. Aspiration/nonaspiration techniques were used. After prior written consent, FNA was done with standard technique and aseptic precautions by using 10cc disposable syringe and 23–25-gauge needle. Aspiration was done from more than one site. Material obtained was smeared on glass slides and smears were stained with Leishman’s and Hematoxylin and Eosin (H and E) stains. Detailed cytomorphological features were studied.

Detailed clinical history, ultrasonography findings, serum T3, serum T4, and thyroid stimulating hormone (TSH) and anti-TPO antibodies were noted and correlated with FNA features. The surgical specimens of thyroid were received for histological examination and were formalin-fixed and paraffin-processed. The 3-4 thick micron sections were cut and stained with Hematoxylin and Eosin (H & E).

3. Results

Out of 110 thyroiditis patients, majority (𝑛 = 103) were females and the remaining seven were males. The majority of patients (𝑛 = 37) were in the age group of 21–30 years followed by 33 patients in the age group of 31–40 years, 17 in 41–50, and 12 in 51–60 years. Four patients were in the age group of 51–60 years and one patient was in the age group of 61–70 years. The majority of the patients in our study were diagnosed as HT (𝑛 = 100), followed by HT with colloid goiter (𝑛 = 5) and De Quervain’s thyroiditis (𝑛 = 3) and one case each of postpartum thyroiditis and HT with associated malignancy (Table 1).

All the patients presented with goiter. The 73 patients presented with diffuse enlargement of thyroid, 34 patients with uneven enlargement, and 03 patients with solitary nodule. Table 1 shows detailed correlation of nature of goiter, thyroid function test, and anti-TPO antibodies in 110 patients of thyroiditis.

Ultrasonography findings of thyroid were available in 48 patients. The 38 patients were diagnosed as thyroiditis and showed features such as diffusely altered parenchyma with heterogeneous echogenicity, hypovascular goiter, micronodules, ill-defined hypoechogenic regions, and marked fibrosis. One case showed dominant nodule which turned out to be papillary carcinoma of thyroid in HT on FNA and histopathological examination. The 10 patients were diagnosed as multinodular goiter.

FNA cytology smears in all cases showed increased background lymphocytes. The lymphocytic infiltration of thyroid follicular cell clusters was seen in 79 cases and Hurthle cells in 64 cases. Other associated features such as mild anisonucleosis, giant cells, histiocytes, scanty colloid, epithelioid cells, plasma cells, fire flares, and eosinophils were seen in few cases. Table 2 shows the frequency of cytomorphological features in 100 cases of HT. Based on severity of lymphocytic infiltrate and presence or absence of Hurthle cells in cytology smears, we categorized HT cases into three groups (Table 3).

We came across three patients of De Quervain’s thyroiditis in our study. All three patients presented with painful uneven enlargement of thyroid; thyroid function was variable and, in all patients, anti-TPO titers were normal (Table 1). FNA...
In all cases, anti-TPO antibody titers were elevated (Table 1). FNA smears in all cases improved the preoperative assessment of thyroid lesions due to its high positive and negative predictive value [22–24]. HT and De Quervain's thyroiditis are the most commonly encountered although overlap of both these conditions is documented [25–27]. In our study the majority of patients were of HT (n = 100).

The male-to-female ratio in our study was 1:14.7. Thyroiditis was more common in females (n = 103) than males (n = 7). The majority of our patients (n = 100) had HT. Maximum number of patients (n = 68) were in between the ages of 21 and 40 years. Our results were in accordance with many studies [6, 7, 9, 12, 28–31]. In contrast, in Italian study, male-to-female ratio was only 1:3.2 [32]. The only case of De Quervain's thyroiditis in our study was a 45-year-old female.

Similar age group was observed by Nishihara et al. [33]. In contrast Vanderpump et al. [34] found a man aged 59 years with HT at the time of diagnosis. A greater number of younger patients in our study can be explained by the fact that the area they live in is a noncoastal area which can be iodine-deficient. In these areas, HT is known to occur in young patients [26]. Other reason can be early diagnosis.

The majority of patients in our study (n = 73) had diffuse enlargement of thyroid. Discrete nodular presentation is well known in HT [12]. Higher number of discrete nodularities (n = 34) in our study can be due to younger age of the patients and early stage of the disease at the time of diagnosis. Out of three solitary nodules, one case was eventually turned out to be HT with papillary carcinoma of thyroid.

Thyroid function tests were done in all patients. Large number of hypothyroid patients (n = 52) in our study can be explained by the fact that majority of thyroiditis patients had HT and possibly the advanced stage of the disease at the time of diagnosis. Substantial number of patients had hyperthyroidism (n = 27). This can be explained by the fact that Hashitoxicosis is a transient hyperthyroid phase of HT.

Anti-TPO antibody titers were done in 71 cases out of which 47 patients had elevated titers. Autoantibodies against thyroglobulin and thyroid peroxidase antigens are clinically most important for diagnosis. Elevated titers have been in 95% of the patients [6]. There is a controversy whether anti-TPO alone is sufficiently reliable to diagnose HT. Anti-TPO negative cases are known to occur. Up to 20% adult females with no clinical disease have detectable Tg/TPO antibodies, raising the question about their pathologic significance.

Thyroid imaging is nowadays an essential part of thyroid disease evaluation and it may aid in establishing proper diagnosis. The most commonly applied modality in evaluation of thyroiditis is conventional ultrasonography [17]. Similar observations were made in our study. The 10 patients diagnosed as MNG turned out as HT on FNA. The most consistent cytological features seen in FNA smears of HT were increased background lymphocytes, lymphocytic infiltration of thyroid follicular cell clusters, and Hurthle cells. HT shows two patterns in cytology smears which correspond to different phases of the disease: (1) classic HT, in which smears show increased background lymphocytes and infiltration of follicular cell clusters by lymphocytes (Figures 1(a), 1(b), and 1(c)); (2) florid lymphocytic pattern in which smears show dominant lymphoid cell population in stages of maturation. Epithelial cells may be inconspicuous

Table 2: Frequency of various cytological features in Hashimoto's thyroiditis.

<table>
<thead>
<tr>
<th>Cytomorphological features</th>
<th>Number of cases</th>
<th>Percentage (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Increased background lymphocytes</td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>Lymphocytic infiltration of follicular cell clusters</td>
<td>79</td>
<td>79</td>
</tr>
<tr>
<td>Hurthle cells</td>
<td>64</td>
<td>64</td>
</tr>
<tr>
<td>Mild anisonucleosis</td>
<td>19</td>
<td>19</td>
</tr>
<tr>
<td>Few giant cells</td>
<td>18</td>
<td>18</td>
</tr>
<tr>
<td>Histiocytes</td>
<td>12</td>
<td>12</td>
</tr>
<tr>
<td>Scanty colloid</td>
<td>8</td>
<td>8</td>
</tr>
<tr>
<td>Epithelioid cells</td>
<td>6</td>
<td>6</td>
</tr>
<tr>
<td>Plasma cells</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>Fire flares</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>Eosinophils</td>
<td>1</td>
<td>1</td>
</tr>
</tbody>
</table>
Figure 1: Hashimoto’s thyroiditis: (a) Group 1 smears showing follicular cell clusters infiltrated by few lymphocytes (Leishman’s stain ×100); (b) Group 2 smears showing follicular cell clusters infiltrated by moderate amount of lymphocytes (Leishman’s stain ×400); (c) Hurthle cells and increased background lymphocytes (Leishman’s stain ×400); (d) dense lymphocytes in follicular cell clusters (Haematoxylin & Eosin ×100).

Figure 2: Hashimoto’s thyroiditis with colloid goiter: smears showing thin background colloid and follicular cell cluster infiltrated by lymphocytes (Leishman’s stain ×400); (b) Hashimoto’s thyroiditis with colloid goiter (Haematoxylin & Eosin ×400).
Table 3: Cytomorphological features of three groups of HT patients.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Number of cases</th>
<th>Cytomorphological features</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>46</td>
<td>Mild lymphocytic infiltrate in follicular cells with or without Hurthle cells</td>
</tr>
<tr>
<td>2</td>
<td>40</td>
<td>Moderate lymphocytes, follicular cell destruction, and Hurthle cells</td>
</tr>
<tr>
<td>3</td>
<td>14</td>
<td>Dense lymphocytes/lymphoid cells in various stages of transformation with very few follicular and Hurthle cells</td>
</tr>
</tbody>
</table>

Figure 3: De Quervain’s thyroiditis: (a) smears showing follicular cell clusters with many giant cells (thick arrow, Leishman’s stain ×100); (b) epithelioid cells and lymphocytes (Leishman’s stain ×400).

Figure 4: Hashimoto’s thyroiditis with thyroid carcinoma: (a) smears showing many large atypical cells (Leishman’s stain ×400); (b) atypical follicular cells with intranuclear cytoplasmic inclusion; (c) tissue section showing papillary thyroid carcinoma (Haematoxylin & Eosin ×400).

(Figure 1(d)). Similar observations were made in our study. Florid lymphocytic thyroiditis at times can be difficult to differentiate from lymphoma in cytology smears especially in adults. But careful search for other cytological features of HT reduces the diagnostic errors. It is now widely accepted that florid lymphocytic thyroiditis and HT represent different manifestations of autoimmune thyroiditis.

HT patients were grouped into Group 1, Group 2, and Group 3 based on severity of lymphocytic infiltrate and were correlated with thyroid function and anti-TPO titers. Group 1, Group 2, and Group 3 suggested milder early stage, more severe form and late stage of the disease, respectively. Other cytomorphological features listed in Table 2 were seen in variable numbers in FNA smears.

Scanty colloid is not an usual feature in FNA smears of HT (n = 8). But careful search for lymphocytic infiltrate and repeat aspiration from other area establish the diagnosis of HT. Other associated features such as mild anisonucleosis
(n = 19), few giant cells (n = 19), and histiocytes (n = 12), though were not diagnostic, were seen in variable numbers in cytology smears and this is known to occur. Fire flares were noted in few patients of HT (n = 3). Similar frequency was noted by Rathi et al. [29]. In contrast, some studies noted slightly more frequency [26, 35]. It is known that fire flares can be seen in variety of benign and malignant thyroid lesions. Ekambaram et al. [36] found eosinophils in 84% patients of HT in thyroid smears. In contrast, we found eosinophils in only one cases.

Other autoimmune disorders may coexist with HT [37]. We did not find any significant evidence of any other autoimmune disorders in any patients with the available history.

FNA diagnosis of HT with colloid goiter was made in few cases (n = 5). Clinical findings and radiological findings suggested colloid goiter. FNA smears in these patients showed moderate to severe background colloid and lymphocytes infiltrating follicular cell clusters and Hurthle cells (Figure 3(a)). Anti-TPO titers were done in all five cases out of which titers were elevated in two cases (Table 2). Surgical specimens of partial thyroidectomy were received in three cases for histopathological examination and confirmed the diagnosis of colloid goiter with HT (Figure 3(b)).

Neoplastic and nonneoplastic lesions are known to be associated with HT like colloid goiter, follicular adenoma, Hurthle cell neoplasm, papillary thyroid carcinoma (PTC), non-Hodgkin’s lymphoma (NHL), and follicular carcinoma [38]. In various surgical series, the prevalence of malignancy in HT ranges from 0.4% to 28% [39, 40]. The most commonly encountered neoplasms in association with HT are papillary thyroid carcinoma and primary thyroid lymphoma [41]. Similar observations were made in our study. We came across only one case of HT with PTC in our study. We missed definitive diagnosis of HT with PTC because neoplastic component was scanty and did not show definitive features of PTC.

We came across three patients of De Quervain’s thyroiditis. They were between the ages of 31 and 50 years. All three patients complained of painful goiter. FNA features, painful goiter, and elevated erythrocyte sedimentation rate and C-reactive protein established the definitive diagnosis. Etiology is probably viral. In majority of the cases, there is compete resolution [42].

Postpartum thyroiditis is a variant of autoimmune thyroiditis. It occurs during the postpartum period in up to 5% of women. The 1/3 of cases can evolve into overt hypothyroidism over a period of 10 years. Most of the patients have elevated circulating anti-TPO. Similar observation was made in our study (Table 1). We conclude that HT is the most common form of thyroiditis and it occurs most commonly in young and middle age women. The diagnostic cytological features seen in FNA smears of HT are increased background lymphocytes, lymphocytic infiltration of thyroid follicular cell clusters, and Hurthle cells. Clinical history, anti-TPO, thyroid function test, and ultrasonography are very important and useful adjuncts in the diagnosis of HT. FNAC remains the “Gold Standard” for diagnosing Hashimoto’s thyroiditis. Clinical history, thyroid function, and biochemical parameters are key for diagnosis of other forms of thyroiditis.

**Disclosure**

The work should be attributed to Department of Pathology, Dr. D. Y. Patil Medical College, Pimpri, Pune 411018.

**Conflicts of Interest**

The authors declare that there are no conflicts of interest.

**References**


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