January 2017

Thyroid ultrasound: State of the art part 1 - Thyroid ultrasound reporting and diffuse thyroid diseases

Manjiri Dighe
University of Washington

Richard Barr
Northeastern Ohio Medical University

Jörg Bojunga
Goethe University Hospital

Vito Cantisani
University Sapienza

Maria Cristina Chammas
University of São Paulo

See next page for additional authors

Follow this and additional works at: http://ecommons.aku.edu/eastafrica_fhs_mc_imaging_diagn_radiol

Part of the Radiology Commons

Recommended Citation
Available at: http://ecommons.aku.edu/eastafrica_fhs_mc_imaging_diagn_radiol/18
Thyroid Ultrasound: State of the Art
Part 1 – Thyroid Ultrasound reporting and Diffuse Thyroid Diseases

Manjiri Dighe1, Richard Barr2, Jörg Bojunga3, Vito Cantisani4, Maria Cristina Chammas5, David Cosgrove6, Xin-Wu Cui7, Yi Dong8, Franziska Fenner9, Maija Radzina10, Sudhir Vinayak11, Jun-Mei Xu12, Christoph F Dietrich13,14

1Department of Radiology, University of Washington, Seattle, USA, 2Department of Radiology, Northeastern Ohio Medical University, Rootstown, Ohio, USA and Southwoods Imaging, Youngstown, Ohio, USA, 3Department of Internal Medicine 1, Endocrinology and Diabetology, Goethe University Hospital, Frankfurt am Main, Germany, 4Department of Radiological Sciences, Oncology and Pathology, Policlinico Umberto I, University Sapienza, Rome, Italy, 5Ultrasound Division, Department of Radiology, Hospital das Clinicas, School of Medicine, University of São Paulo, São Paulo, Brazil, 6Division of Radiology, Imperial and Kings Colleges, London, UK, 7Department of Medical Ultrasound, Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, China, 8Department of Ultrasound, Zhongshan Hospital, Fudan University, Shanghai, China, 9Department of Surgery, Caritas-Krankenhaus Bad Mergentheim, Germany, 10Diagnostic Radiology Institute, Paula Stradins Clinical University Hospital, Riga, Latvia, 11Department of Imaging and Diagnostic Radiology, Aga Khan University Hospital, Nairobi, Kenya, 12Department of Medical Ultrasound, Shanghai Tenth People’s Hospital, Ultrasound Research and Education Institute, Tongji University School of Medicine, China, 13Department of Internal Medicine 2, Caritas Hospital, Bad Mergentheim, Germany, 14Sino-German Research Center of Ultrasound in Medicine, The First Affiliated Hospital of Zhengzhou University, Zhengzhou, China

Abstract
Accurate differentiation of focal thyroid nodules (FTL) and thyroid abnormalities is pivotal for proper diagnostic and therapeutic work-up. In these two part articles, the role of ultrasound techniques in the characterization of FTL and evaluation of diffuse thyroid diseases is described to expand on the recently published World Federation in Ultrasound and Medicine (WFUMB) thyroid elastography guidelines and review how this guideline fits into a complete thyroid ultrasound exam.

Keywords: thyroid; ultrasonography; elastography; color Doppler; contrast enhanced ultrasound (CEUS); point of care (POC).

Introduction

Ultrasound (US) of the thyroid is well established for a variety of indications and applications. US is the most sensitive imaging test available for the examination of the thyroid gland, to detect and characterize lesions, accurately calculate their dimensions, identify the internal structure and vascularization using color Doppler imaging (CDI) and evaluate diffuse changes in the thyroid parenchyma. Thyroid US is able to confirm the presence of a thyroid nodule when the physical examination is equivocal and differentiate between thyroid nodules and cervical masses from other origins [1,2]. In this setting, the accurate differentiation of benign from malignant lesions is critical to ensure that the patient undergoes the appropriate diagnostic and therapeutic decision.

In the present two part papers, the role of ultrasound techniques in the evaluation of diffuse thyroid disease (DTD) (Part 1) and characterization of focal thyroid lesions (FTL) (Part 2) is described to accompany the recently published WFUMB thyroid elastography guide-
lines and review how the guidelines fit into a complete thyroid ultrasound exam.

**Clinical indications according to guidelines**

Thyroid ultrasound is recommended in [1,2]:
1. All patients with a palpable thyroid nodule or with multinodular goiter.
2. High-risk patients for thyroid malignancy: history of familial thyroid cancer, multiple endocrine neoplasia (MEN) type II and irradiated neck in childhood.
3. Patients with palpable cervical adenopathy suspicious for malignancy.
4. Follow-up and monitoring thyroid nodules.

Thyroid ultrasound is not recommended in [1,2]:
1. Patients with a normal thyroid on palpation and low risk of thyroid cancer.
2. As a screening test in the general population.

**Examination technique**

The recommended protocol for thyroid US is as follows [3,4] (Table I):
1. Patient is scanned in supine position with neck extended with a small pillow or rolled towel behind the shoulders.
2. Axial scans of the whole gland at the upper, mid, lower poles and the isthmus, and side-by-side images of each lobe, to compare echogenicity and size of both lobes. Each lobe width and AP diameters are measured.
3. Longitudinal scans through each lobe, on medial, mid and lateral planes. The length of the lobes is measured.
4. Identify focal lesions, measure the main lesions and identify the dominant one (according to size).
5. Evaluation of vascularization using CDI of both lobes and any lesions if detected.

Nodules with any malignant potential (essentially all nodules except pure cysts) should be identified [3,7].

<table>
<thead>
<tr>
<th>Thyroid gland</th>
<th>Thyroid nodules</th>
</tr>
</thead>
<tbody>
<tr>
<td>Position</td>
<td>Size / Location</td>
</tr>
<tr>
<td>Shape</td>
<td>Margins</td>
</tr>
<tr>
<td>Size</td>
<td>Composition (solid, cystic, spongiform, proportion)</td>
</tr>
<tr>
<td>Content</td>
<td>Presence and type of calcifications</td>
</tr>
<tr>
<td>Echogenicity</td>
<td>Echogenicity iso/hyper/hypo</td>
</tr>
<tr>
<td>Vascular pattern</td>
<td>Vascular pattern</td>
</tr>
</tbody>
</table>

**The value of thyroid ultrasound examination**

The value of thyroid ultrasound examination is as follows:
1. To detect FTL;
2. To differentiate between benign and malignant FTL;
3. To evaluate cervical lymphadenopathy;
4. To guide interventions, e.g., biopsy and percutaneous treatment;
5. To assess change in size of thyroid nodules with follow up US.

In all patients with palpable thyroid nodules, US should be performed to accomplish the following [3]:
1. Detect US features suggestive of malignant growth and select the lesions to be recommended for fine-needle aspiration (FNA);
2. Look for coincidental thyroid nodules;
3. Help with the diagnosis in difficult cases (as in Hashimoto’s thyroiditis);
4. Choose the gauge and length of the biopsy needle;
5. Obtain an objective measure of the baseline volume and characteristics of the lesions that will be assigned to follow-up or medical or minimally invasive therapy – Radiofrequency ablation (RFA), Microwave ablation (MWA), Laser.

Indications for thyroid US, following the American Association of Clinical Endocrinologists (AACE) are as follows [8]:
1. To confirm presence of a thyroid nodule when physical examination is equivocal.
2. To characterize a thyroid nodule(s), i.e. to measure the dimensions accurately and to identify internal structure and vascularization.
3. To differentiate between benign and malignant thyroid masses, based on their sonographic appearance.
4. To differentiate between thyroid nodules and other cervical masses like lymphadenopathy, thyroglossal cyst and cystic hygroma.
5. To evaluate diffuse changes in thyroid parenchyma.
6. To detect post-operative residual or recurrent tumor in thyroid bed or metastases to neck lymph nodes.
7. To screen high-risk patients for thyroid malignancy like patients with history of familial thyroid cancer, MEN type II and irradiated neck in childhood.
8. To guide diagnostic (FNA cytology/biopsy) and therapeutic interventional procedures.

**TIRADS (thyroid imaging reporting and data system) – Classification**

In the last few years, assessment concepts called as ‘grading system’ or ‘reporting system’ have emerged. In
Table II. TIRADS classification [9].

<table>
<thead>
<tr>
<th>Category</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>6</th>
</tr>
</thead>
<tbody>
<tr>
<td>Definition</td>
<td>Normal thyroid gland</td>
<td>Benign</td>
<td>Probably benign</td>
<td>Suspicious for malignancy</td>
<td>Probably malignant</td>
<td>Biopsy proven malignancy</td>
</tr>
<tr>
<td>Number of suspicious features</td>
<td>4a</td>
<td>4b</td>
<td>4c</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Fig 1. Normal thyroid gland. Transverse ultrasound through the neck shows that the normal thyroid consists of two lobes and a bridging isthmus with a homogenous appearance on ultrasound.

In case of a positive answer to any of the above questions, FNA must be performed. Recommendations for FNA of thyroid nodules based on US features were published by the American Thyroid Association (ATA) in 2015 [2].

The TIRADS classification is shown in Table II [9].

Normal size, shape, echogenicity

The normal thyroid gland consists of two lobes and a bridging isthmus. Thyroid size, shape, and volume vary with age and sex. The normal thyroid gland (fig 1) weighs approximately 30 g but the size and shape vary according to age [14].

When the gland enlarges, it may extend inferiorly into the superior mediastinum and is then described as a retrosternal thyroid. The volume of the thyroid gland is quite important for clinical practice since it identifies the enlargement of the thyroid gland (goiter) and its response to suppressive therapy. For the treatment of Graves’ disease it allows a rigorous calculation of the appropriate radio-iodine dose. Diminution of the thyroid volume (thyroid atrophy) can be detected in some cases, but has less clinical importance. Normal thyroid lobe dimensions are: 18-20 mm longitudinal and 8-9 mm antero-posterior (AP) diameter in newborn; 25 mm longitudinal and 12-15 mm AP diameter at one year age; and 40-60 mm longitudinal and 13-18 mm AP diameter in adult population. Measurement of the thyroid lobe involves three measurements: the width, depth and length, then the volume is calculated by the formula $V = W \times D \times L \times 0.523$ (V-volume, W-width, D-depth, L-length). The total volume
results from the sum of the two volumes; the isthmus is omitted, unless its thickness is over 3 mm. The limits of normal thyroid volume are 10-15 ml for females and 12-18 ml for males. The width and depth are measured on transverse section of the lobe: the width is the distance between the most lateral point of the lobe and the acoustic shadowing of the trachea and the depth is the maximum antero-posterior distance in the middle third of the lobe (fig 2) [15].

In some cases, on the upper margin of the isthmus there is an accessory lobe (Lalouette pyramid), which must be identified and measured, and the volume must be added to that of the lobes [16].

The structure of the thyroid gland is granulated (ground glass appearance) and the echogenicity is similar to that of the parotid glands, and of increased echogenicity compared to that of the adjacent muscles [15]. The relationships with surrounding structures are: sternocleidomastoid and strap muscles anteriorly; trachea/esophagus and longus colli muscles posteriorly; and common carotid arteries and jugular veins bilaterally (fig 2) [17,18].

**Congenital anomalies**

Congenital agenesis or hypoplasia of the thyroid gland may include the whole gland or just one of the lobes [19]. Congenital hypothyroidism is a relatively common endocrine disorder [20]. Approximately 80% of congenital hypothyroidism is caused by thyroid dysgenesis due to absence, hypoplasia or ectopia of the gland, which is almost always sporadic in nature [21]. Ectopic thyroid tissue is derived from incomplete migration of thyroid gland and can be found anywhere along the migration course of the thyroid primordium [22] which is between the base of tongue and pretracheal region. The most common presentation is a euthyroid neck mass [23]. Ectopic thyroid tissue most commonly appears in the midline in the cervical region (90% of the cases) [24]. Appearance varies on US, but the tissue is generally located close to the hyoid bone (fig 3) [20]. Its prevalence is approximately 1/100,000 – 1/300,000 [25].

**Tips and tricks:** Scintigraphy is more sensitive than US for detecting an ectopic thyroid. Scintigraphy provides functional information of the thyroid which is not available based on ultrasound examination alone. The thyroid gland appears normal on US in patients with hypoplasia; however, scintigraphy reveals decreased isotope uptake [20].

**Diffuse thyroid disease (DTD)**

A wide spectrum of DTDs affect the thyroid gland with the most common being autoimmune thyroid disease (AITD). Hashimoto’s thyroiditis and Graves’ disease are the most common AITDs. US is not generally required for the diagnosis of DTD and diagnosis is based on presenting symptoms, laboratory analysis of thyroid function, immunology, and occasionally radioactive iodine uptake scans [26]. Radioiodine scans are neither needed nor recommended to diagnose Hashimoto or Graves’ disease. In some cases, like Hashimoto’s thyroiditis, the disease is primarily subclinical and US is helpful in detecting these patients. In addition, US helps exclude focal thyroid disease in these patients and in assessing the size of the thyroid gland.

Ultrasongraphy findings indicative of DTD are characterized by a diffusely enlarged or normal size thyroid gland or small size in later stages of DTD, decreased or increased diffuse heterogeneous parenchymal echogenicity, a coarse echotexture, and micronodulation [27,28].

In theory, differentiation with ultrasound elastography is feasible because of the different pathologic fea-
tures and stiffness. Yang et al in their study found that the strain ratio (SR) values (calculated by comparing the strain in the thyroid nodule to the sternocleidomastoid muscle in the same image) ranked in ascending order, control group < hyperthyroidism group < Hashimoto’s thyroiditis group < subacute thyroiditis group with statistically significant difference (p<0.05) between groups and that it is feasible to assess diffuse thyroid disease with strain ratios obtained with ultrasound elastography [29]. In clinical practice, elastography is usually not used for these indications.

**Goiter, Struma diffusa**

Generalized enlargement of the thyroid is called goiter and can be diffuse or nodular. It may be within a range from simple diffuse nontoxic non-nodular thyroid enlargement to multinodular goiter in a euthyroid patient. The cause of simple goiter is multifactorial and involves complex interactions between environmental (iodine intake), genetic, and endogenous (female gender) factors [30]. Insufficient thyroid hormone input is the most frequent cause worldwide except for the United States; it begins as reaction with follicular epithelium compensatory hypertrophy which generates ordinary, polyclonal follicles alternating with scarring caused by hemorrhagic necrosis over the course of goiter growth in an attempt to maintain a euthyroid state [31,32]. At conventional B-mode US and CDI multinodular goiter frequently appears as thyroid enlargement with focal or diffuse replacement of the thyroid parenchyma by strictly adjacent, sometimes, not distinctive, variable echo structure nodules containing variable amount of cystic degeneration, vascularization and dystrophic calcifications, without or with minimal normal remaining parenchyma (fig 4). In the case of multiple sonographically similar appearing nodules, representative nodules, or nodule clusters when involvement is focal, can be identified for measurement to obtain more reproducible follow-up [33]. The main aim of ultrasound is to identify nodules that have malignant sonographic features (see TIRADS) to be submitted to ultrasound-guided FNA [34].

**Elastography**

To date, no specific papers, neither specific recommendation, to the best of our knowledge have been published focusing on diffuse goiter. For the specific features regarding single nodules please see below.

**Contrast enhanced ultrasound**

To date, no specific papers, to the best of our knowledge have been published focusing on diffuse goiter. For the specific features regarding single nodules please see below.

**Tips and tricks**

Thyroid scintigraphy is often used in cases where the cause of the hyperthyroidism is unclear and to evaluate for thyroid nodules before treatment [35]. Society of Radiologists in Ultrasound [36] recommend that sonographic features of individual nodules in a multinodular gland should be the primary criteria for suggesting biopsy rather than using nodule size. In their recent guidelines published in 2015 ATA [2] recommends that patients with multiple nodules >1cm should be evaluated in the same fashion as patients with a solitary nodule >1cm with each nodule >1cm carrying an independent risk of malignancy. If multiple nodules >1cm are present, those with a suspicious sonographic pattern should be aspirated preferentially and if none of the nodules have a high or moderately suspicious sonographic pattern, nodules greater than 2 cm should be aspirated and the rest should undergo follow up with serial US examinations [2]. The ATA guidelines use size criteria not since there is an increased risk of cancer in larger nodules but because lesions with different size have different prognosis.

**Thyroiditis**

Different causes of thyroiditis have been identified including: infection, autoimmune processes, medication, and ionizing radiation. As a result of the inflammatory process, temporary or permanent impairment of thyroid function usually follows. The different forms of thyroiditis may be clearly distinguished on the basis of clinical findings, a few laboratory values, and the ultrasound results [37]. US guided FNA with cytological examination is rarely needed. Scintigraphy of the thyroid is usually of no added diagnostic value. Impairment of thyroid function is divided into hyper- and hypothyroidism [38].

The commonest types of thyroiditis encountered are Hashimoto’s thyroiditis and de Quervain’s thyroiditis. This section will restrict itself to discussion of these two conditions in addition to a discussion of Graves’ disease [39].

---

![Fig 4. Multinodular goiter. A 43 year old patient with multinodular goiter and hypothyroidism. Sagittal ultrasound the right lobe showed an enlarged thyroid gland with multiple iso- to hyperechoic nodules of variable size, however similar in appearance.](image)
Hashimoto’s thyroiditis or chronic lymphocytic thyroiditis (CLT) is the most common form of organ-specific autoimmune diseases [20]. It is the commonest cause of hypothyroidism in the United States [27]. Its annual incidence is estimated to be between 0.3 and 1.5 cases per 1000 persons, with no significant race-related predominance [27]. Hashimoto’s thyroiditis affects 1.3% of children and has a female predominance [40], with an 8 to 9:1 female to male ratio [41]. The disease usually develops in young or middle aged women and leads to progressive thyroid failure [42]. It is characterized by diffuse interstitial lymphocytic infiltration and a variable degree of fibrosis [43]. Diagnosis is made by detecting anti-thyroid antibodies, including anti-thyroid peroxidase and anti-thyroglobulin antibodies [20]. Patients with the acute form present with painless, lobular, diffusely enlarged thyroid gland (classical Hashimoto’s) or – much more common – an atrophic thyroid (historically named as Ord’s thyroiditis) [27].

Conventional B-mode ultrasound

Hashimoto’s thyroiditis is primarily a subclinical disease, and US can detect this subset of patients before they come to clinical attention when typical US findings are present [44].

US is useful for measuring thyroid size and assessing echotexture. Sonographic appearances vary depending on the degree of gland involvement, which include severity of follicular disruption, lymphocytic infiltration, and chronicity of disease and extent of thyroid involvement. Initially the parenchyma is heterogeneous and coarsened compared with normal thyroid (fig 5). In some patients, innumerable hypoechoic solid micro nodules are seen ranging in size from 1 to 7 mm surrounded by echogenic rim of fibrosis. This appearance is highly specific with a positive predictive value of 95% [28]. As the disease progresses, thyroid parenchyma is progressively destroyed and develops echogenic linear bands of parenchymal fibrosis which can become confluent and thicker (fig 6). There can be asymmetric involvement with preference for the anterior part of the gland. Eventually the gland becomes atrophic with a hypoechoic appearance similar to that of strap muscles.

Color Doppler imaging (CDI)

CDI in Hashimoto’s is variable with either normal or increased vascularity seen in the early disease. The increase in vascularity seems to be associated with development of hypothyroidism [45]. Later in the disease course, the gland shows decreased vascularity.

On CDI, the hypertrophic form (Hashimoto’s thyroiditis) presents itself with slightly increased vascularity (hyperthyroid phase) in the early stages (fig 7a), whereas the stage of atrophy is characterized by reduced or absent vascularization (fig 7b). During the phase of clinically manifest hypothyroidism, vascularization may increase once more. Peak systolic velocity (PSV) in the afferent arteries is normal in all stages (slightly increased in the case of manifest hypothyroidism), thereby facilitating the differentiation from Graves’ disease [38].

Elastography

Sporea et al reported a cut-off value >2.53 m/s using shear wave elastography (SWE) for differentiation between normal thyroid and diffuse thyroid diseases, with a positive predictive value >90% [46]. They also found a statistically significant difference in velocity values in autoimmune pathology with a value of 2.07±0.44 m/s in...
Graves’ disease compared with 2.68±0.50 m/s in chronic autoimmune thyroiditis [47].

Kim et al found a cut-off mean value of the EI (elasticity index) using carotid artery as the internal compression source, in cases of diffuse thyroid disease, of 27.6 kPa and a maximum value of 41.3 kPa, with a sensitivity of 40.9% and a specificity of 82.9% [48]. Magri et al studied the EI in patients with chronic autoimmune thyroiditis and found that the elasticity of extra nodular tissue is decreased according to the thyroid antibody titer and the degree of thyroid function damage [49].

Fukuhara et al evaluated the utility of Acoustic Radiation Force Impulse (ARFI) SWE for diagnosing chronic autoimmune thyroiditis (CAT) and to verify the effect of fibrotic thyroid tissue on SWV. The SWV for CAT (2.47±0.57 m/s) was significantly higher than that for controls healthy subjects (1.59±0.41 m/s) (p<0.001). AUROC for CAT was 0.899, and the SWV cut-off value was 1.96 m/s. The sensitivity, specificity, and diagnostic accuracy were 87.4%, 78.7%, and 85.1%, respectively. Levels of anti-thyroperoxidase antibodies and thyroid isthmus thickness were correlated with tissue stiffness in CAT. However, there was no correlation between levels of anti-thyroglobulin antibodies and tissue stiffness. Quantitative SWE is useful for diagnosing CAT, and it is possible that SWE can be used to evaluate the degree of fibrosis in patients with CAT [50].

Magri et al indicated that SWE correctly defines the elasticity of thyroid nodules independently from the coexistence of autoimmune thyroiditis, always being able to differentiate nodular tissue from the surrounding parenchyma. In Hashimoto’s thyroiditis the stiffness of extra-nodular tissue increases in relation to both the thyroid antibody titer and the degree of impairment of thyroid function [49]. For a cut-off value of 22.3 kPa, which resulted in the highest sum of sensitivity and specificity, the EI assessed by SWE had a sensitivity of 59.6% and a specificity of 76.9% (AUROC=0.71; p<0.001) for predicting the presence of autoimmune thyroid disease [51].

However, in clinical practice, elastography is usually not needed for these indications.

**Contrast enhanced ultrasound**

To date, no specific papers, to the best of our knowledge have been published focusing on Hashimoto’s thyroiditis. For the specific features regarding single nodules please see below.

**Tips and tricks**

US guided FNA is an appropriate means of establishing tissue diagnosis and should be used in association with appropriate laboratory studies, FNA, demonstrating diffuse lymphocytic infiltration, as well as high-titer antibodies against thyroid peroxidase (anti-TPO) (sensitivity 70-90%) might be helpful in the diagnostics of chronic Hashimoto’s thyroiditis in the early stages. The atrophic form is characterized by an elevated basal TSH level and requires no further laboratory tests or cytological diagnostics. Scintigraphy (mottled accumulation pattern and low nuclide uptake) provides no additional information [38].

**b) Graves’ disease**

Hyperthyroidism is rare in childhood and is most commonly caused by Graves’ disease. It affects 0.02% of children or 1 in 5000 [40]. The peak incidence occurs from 11-15 years of age with a female predominance. A positive family history is common. Graves’ disease is an autoimmune disorder that results in hyper-functioning of the thyroid. It is caused by thyroid receptor antibodies binding to the thyrotropin receptor (TSH-receptor). The typical biochemical thyrotoxic profile is matched by a diffuse enlargement of the thyroid gland with rounding of the normal angular outline [52].

**Conventional B-mode ultrasound**

No specific findings are present, however findings suggestive of the disease include diffuse enlargement, convex bowing of the anterior gland margin and mild textural coarsening (fig 8). The echogenicity is often decreased due to the increased blood flow, increased cellularity and decreased colloid content. Compared to Hashimoto’s thyroiditis, the appearance of the thyroid in Graves’ disease is less heterogeneous and the contour is lobulated [53]. Features of increased volume, hypervascularity, and heterogeneous reduced echogenicity have been shown in Graves’ disease [35].

**Color Doppler imaging**

Normal thyroid parenchyma shows occasional spots of flow on color Doppler; peak systolic velocities between 15 and 30 cm/s in the inferior thyroid artery and 3 to 5 cm/s in the intrathyroid arteries are considered within the range of normal [54]. Increased vascularity and arteriovenous shunting is called “thyroid inferno” which is suggestive of Graves’ disease (fig 9). Studies have shown that
thyroid and intra-thyroid arteries [56]. Flow on spectral Doppler in the medium-sized perithyroidal arteries was approximately 15-fold higher TBF measured in mL/min compared with normal and high peak systolic velocities in Graves’ disease [55]. Patients with Graves’ disease show markedly increased vascularity in Graves’ disease than in painless thyroiditis, subacute thyroiditis, or normal controls. The TBF of patients with painless thyroiditis, subacute thyroiditis, or normal controls. The TBF of patients with Graves’ disease was consistently >4%, and all other patients had TBF <4%, indicating that 4% is the cutoff for distinguishing destruction-induced thyrotoxicosis and Graves’ disease [55]. Patients with Graves’ disease show an approximately 15-fold higher TBF measured in mL/min compared with normal and high peak systolic velocity flow on spectral Doppler in the medium-sized perithyroid and intra-thyroid arteries [56].

English et al described the typical sonographic features of the thyroid gland in patients with Graves’ hyperthyroidism after radiiodine therapy (RIT). The sonographic features of the post-RIT gland included a significantly reduced mean total volume of 4.2 mL, hypovascularity, coarse echotexture, and hyperechojenicity [35]. Currently, thyroid imaging does not play a routine role in the post-treatment follow-up of RIT patients, but thyroid ultrasound would be employed if there was a change in the size (enlargement) or nodularity of a treated gland [35].

Elastography

Sporea et al found that thyroid stiffness (TS) assessed by means of ARFI SWE in healthy subjects (2±0.40 m/s) was significantly lower than in Graves’ disease (2.67±0.53 m/s) (p=0.0001) and CAT patients (2.43±0.58 m/s) (p=0.0002), but the differences were not significant between Graves’ disease and CAT patients (p=0.053). The optimal cut-off value for the prediction of diffuse thyroid pathology was 2.36 m/s. For this cut-off value, TS had 62.5% sensitivity, 79.5% specificity, 87.6% predictive positive value, 55.5% negative predictive value, and 72.7% accuracy for the presence of diffuse thyroid gland pathology (AUROC=0.804). There were no significant differences between the TS values obtained with linear vs convex probes and when 5 vs 10 measurements were taken in each lobe (median values). ARFI SWE seems to be a useful method for the evaluation of diffuse thyroid gland pathology and is able to predict with sufficient accuracy the presence of diffuse thyroid diseases (AUROC=0.80) [46,47].

Contrast enhanced ultrasound

To date, no specific papers, to the best of our knowledge, have been published focusing on Graves’ disease.

Tips and tricks

Thyroid scintigraphy is often used in cases where the cause of the hyperthyroidism is unclear and to evaluate thyroid nodules before treatment [35].

c) Subacute granulomatous thyroiditis

(De Quervain or granulomatous thyroiditis)

The clinical scenario differs in that the patient presents with a painful swelling in the lower neck, fever, and sometimes lethargy; typically following a viral illness. Typically no hyperthyroidism is observed. The biochemistry in the acute phase might appear as thyroid toxicity, sometimes followed by a period of hypothyroidism. Typically (after a period of 6 – 18 months from acute onset) the patient recovers and becomes euthyroid. Subacute granulomatous thyroiditis (SGT) is a self-limiting subacute inflammatory disease of the thyroid [57]. It typically occurs in the area of the gland in mid-aged hyperthyroid women complaining of pain, tenderness, fatigue and mild fever [58] and constitutes nearly 3-6% of all thyroid diseases. Generally appears 2 weeks after a viral upper respiratory tract infection and regresses spontaneously within 2-3 months. Although its etiology remains unknown, it is believed have a viral origin, similar to the mumps virus, hepatitis B and C viruses, cytomegalovirus, entervirus, and type A and B coxsackie viruses. Clinically, patients present with localized anterior neck pain associated with glandular tenderness and diffuse pain in the ears and the jaw, which might be accompanied by fatigue, weight loss, low-grade fever, elevated C-Reactive Protein (CRP), elevated erythrocyte sedimentation rate, suppression in the TSH level and occasionally dysphagia [59].

Conventional B-mode ultrasound

Characteristic US findings include ill-defined, moderately or markedly patchy hypoechoic areas in the thyroid. Hypoechoic areas tend to elongate along the long axis of
the thyroid. In severe disease, the capsule can be expanded in the affected regions [60]. Generally, an increase in the size of the thyroid and heterogeneous, diffuse, hypoechoic and confluent areas with negative margins, characteristically defined as “lava flow,” are observed on US (fig 10) [59,61,62]. Hypoechoic appearance can also be seen in malignancy, however other features like calcifications and taller than wide shape are absent. Short-term follow up is useful to document regression or resolution. In addition, a prompt clinical response to anti-inflammatory therapy is highly diagnostic.

In the acute phase, US can detect a hypoechoic ill-defined mass, usually tender. The adjacent thyroid tissue is heterogeneous in echotexture. In the subacute phase, the hypoechoic area increases in size to involve the ipsilateral thyroid lobe and sometimes extends to the contralateral lobe. And during recovery phase, thyroid appearances returns to normal or atrophy may develop [62]. Typically, enlarged and activated lymph nodes can be found on US as well.

Color Doppler imaging

No increase in vascularity occurs at CDI. However, these findings are nonspecific because they may also be seen in CLT, multinodular goiter, and Graves’ diseases, and the clinical findings may assist in the differential diagnosis [59,62].

On CDI, the hypoechoic lesions show reduced or absent vascularity, whereas the remaining thyroid displays normal vascularity. Peak systolic velocity of the afferent arteries is typically normal [63].

The sonographic changes described above are no longer demonstrable after complete remission of the disease.

Elastography

Stiff areas are seen in the thyroid gland in SGT and these stiff areas can be stiffer comparing with chronic thyroiditis, resembling malignant nodules on elastography (fig 11) [64,65]. Xie et al found that real-time strain US elastography does not provide conclusive information in the diagnosis and differential diagnosis of SGT due to its inability to distinguish between this pathology and thyroid cancer [64].

Yang et al found that the SR (mean±standard deviation) (calculated by using the sternocleidomastoid muscle on the same side of the thyroid as the reference tissue) of patients with hyperthyroidism, Hashimoto’s thyroiditis, and SGT were 2.30±1.08, 7.04±7.74, and 24.09±13.56, respectively. The SR of the control group was 1.76±0.54. SR values ranked in ascending order were control group, hyperthyroidism group, Hashimoto’s thyroiditis group, SGT group. There were statistically significant (p<0.05) differences in thyroid hardness between groups with different diffuse thyroid diseases [29].

Thyroid tissue stiffness was higher in SGT at baseline (214.26±32.5 kPa) in comparison with values recorded at a 4-week follow-up (45.92±17.4 kPa) and at 10 weeks following diagnosis and treatment initiation (21.65±5.3 kPa, p<0.0001). Baseline thyroid stiffness in SGT was higher than in CAT (36.15±18.7 kPa, p<0.0001) and healthy control subjects (16.18±5.4 kPa, p<0.0001). In the remission of SGT, thyroid stiffness was lower than that found in CAT (p=0.006), and higher comparing with healthy control subjects (p=0.0008) [66].

Subacute inflammatory pseudonodules were found to have greater stiffness comparing with nodules found in chronic thyroiditis, and in this condition, elastography could be unable to detect thyroid cancer [66].

Contrast enhanced ultrasound

To date, no specific papers, to the best of our knowledge, have been published focusing on subacute granulomatous thyroiditis.

Tips and tricks

SGT may be diagnosed conclusively by B mode ultrasound in 90% of cases if the clinical picture is
typical. Rarely, other thyroid diseases with hypoechoic echo texture must be considered, and CDI contributes with important additional information towards the differential diagnosis. Complementary US guided FNA, demonstrating giant cells and histiocytes, confirms the diagnosis and may be helpful in unclear cases. Scintigraphy, as in other cases of thyroid inflammation, does not contribute any additional information and should not be performed [38].

d) Riedel struma

Riedel’s struma (fibrosing thyroiditis) is an extremely rare, local manifestation of a systemic form of fibrosclerosis. Most patients with Riedel’s thyroiditis (RT) are between 30 and 50 years old at the time of diagnosis [67]. It is a rare inflammatory disease that results in fibrosis of the thyroid gland and invasion to the surrounding structures of the neck [68]. Although the etiology of RT is unclear, the most probable cause is an autoimmune process [69]. The condition is associated with extensive fibrosis in the gland that spreads to the neighboring tissues and is associated with the presence of inflammatory cell infiltrates. It is characterized by an increasingly massive fibrous alteration of the thyroid and the surrounding tissues, resulting in indolent hard swelling of the neck [70]. RT is more common in females, with an operative incidence of 0.06% reported. The disease is frequently associated with fibrosis of the mediastinum and retroperitoneum, similar to IgG4-mediated disease such as Ormond’s disease [71].

Conventional B-mode ultrasound

US shows an enlarged, hypoechoic gland or with a coarsened echotexture with fibrous septations resulting in pseudonodular appearance (fig 12a) [72]. Perithyroid extension is present [73-75].

Color Doppler imaging

Previously published CDI features of RT include a decrease in vascular flow [76], or only slight vascularization of the mass (fig 12 b) [73]. The US vascular en-casement and improvement with corticosteroid treatment clearly appear to be specific to RT [75].

Elastography

Elastography showed heterogeneity in the stiffness values of the thyroid parenchyma varying between 21 kPa and 281 kPa. USE revealed heterogeneity in the stiffness values of the thyroid parenchyma. It should be noted that the hypoechoic areas corresponded to the highest stiffness values. These areas could correspond to a high degree of fibrosis. Longer follow-up times are necessary to evaluate the extent to which corticosteroid treatment can alter such stiffness values [75].

Contrast enhanced ultrasound

To date, no specific papers, to the best of our knowledge have been published focusing on RT.

Tips and tricks

Currently, there are no well-established cytomorphologic criteria published in the literature in RT [77].

e) Post partum thyroiditis

Post-partum thyroiditis (PPT) is a special form of autoimmune thyroiditis, which occurs in the first year after parturition. PPT is characterized by transient hyperthyroidism followed by transient hypothyroidism during the first six postpartum months. It is seen in 5-9% of women and recurred in 75% of cases in subsequent pregnancies, often being not recognized because the symptoms of hyperthyroidism (1-3 months post partum) and of slight hypothyroidism (4-10 months post partum) are misinterpreted as puerperal depression or as a [78]. The disease is self-limiting, and thyroid function commonly returns to normal after one year; permanent hypothyroidism is rare. Diagnosis is based upon clinical manifestations and thyroid function tests (TSH and free T4). Assessment of anti-TPO antibodies may also help in the diagnosis and thyrotropin receptor antibodies (TRAB) should be negative [79].

Conventional B-mode ultrasound

Typically the gland appears hypoechoic either diffuse or with multifocal areas of low echogenicity scattered throughout both lobes of the thyroid resulting in a heterogeneous appearance [80].

Color Doppler imaging

In comparison to Graves’ disease, there is no ‘vascular inferno’. In older studies, blood flow is reported to be mostly reduced and increased vascularity is rare. Peak systolic velocity is normal [38]. However, depending on US-devices used, an increased blood flow can be detected very similar to that seen in the acute phase of Hashimoto’s disease.

Elastography

To date, no specific papers, to the best of our knowledge have been published focusing on postpartum thyroiditis.
**Contrast enhanced ultrasound**

To date, no specific papers, to the best of our knowledge have been published focusing on postpartum thyroiditis.

**Tips and tricks**

Scintigraphy does not have any role in diagnosis or follow-up of patients with PPT, especially since the postpartum patients might be breastfeeding her child, which is a contraindication to radioiodine administration. Neither is biopsy recommended in these patients, unless a concomitant focal lesion is seen.

**f) Silent thyroiditis**

Silent thyroiditis is a rare painless disease. Slight hypothyroidism is clinically apparent, seldom followed by hypothyroidism. Anti-TPO antibodies are often low and only temporarily present. The disease shows spontaneous remission within a few months [81].

**Conventional B-mode ultrasound**

Sonographic findings equal post-partum thyroiditis [82].

**Color Doppler imaging**

No specific information has been listed in the literature in regards to CDI in silent thyroiditis. From personal experience it can be reported that an increased blood flow can be detected very similar to that seen in the acute phase of Hashimoto’s or postpartum thyroiditis.

**Elastography**

To date, no specific papers, to the best of our knowledge have been published focusing on silent thyroiditis.

**Contrast enhanced ultrasound**

To date, no specific papers, to the best of our knowledge have been published focusing on silent thyroiditis.

**Tips and tricks**

Because silent thyroiditis might show US features very similar to other forms of hyperthyroidism, a radioscans can be helpful in differentiating silent thyroiditis from TRAB negative Graves’ disease or a diffuse autonomy. Scintigraphy during the hyperthyroid phase of thyroiditis would show low values for radioiodine uptake, usually less than 1% compared with high values seen in Graves’ disease. This can help in differentiating between the two diseases [83].

**g) Varia: Amiodaron and interferon induced thyroid disease**

Amiodarone is a potent iodine-rich drug; however, side effects on the thyroid and other organs may counterbalance its beneficial effects on the heart. It can cause isolated abnormalities of thyroid function tests and overt thyroid dysfunction, either hypothyroidism (amiodarone-induced hypothyroidism [AIH]) or thyrotoxicosis (amiodarone-induced thyrotoxicosis [AIT] [84]). It may develop early on or after many years of amiodarone treatment, sometimes following drug withdrawal due to prolonged tissue storage of iodinated amiodarone metabolites [85].

AIH occurs more frequently than AIT in iodine-sufficient areas. In contrast to AIT, AIH is more frequent in females with a female to male ratio of 1.5:1 [86]. AIH patients are also older than AIT with circulating thyroid autoantibodies seen in almost 53% of AIH patients with underlying thyroid abnormalities especially Hashimoto’s thyroiditis [87].

Males are more frequently affected by AIT (M/F ratio 3:1). Incidence has been reported to range between 1% and 23% [88]. 15–20% of patients on amiodarone treatment develop either AIT or hypothyroidism. Two types of AIT are present; type 1 is a form of iodine-induced hyperthyroidism that develops in abnormal glands (multinodular goiter or latent Graves’ disease). The thyrotoxicosis in this type is caused by excessive thyroid hormone synthesis and is treated with anti-thyroid drugs. The more common type 2 AIT occurs in patients without underlying thyroid disease. Thyrotoxicosis in these patients is caused by amiodarone-induced destructive thyroiditis; the disease is treated with steroids [89].

In addition, thyroid disease is a common side effect of interferon-based antiviral therapy for chronic hepatitis C, which may lead to dose reduction or discontinuation of therapy. The two entities observed are destructive forms of thyroiditis as well as INF-induced Graves’ disease [90].

**Conventional B-mode ultrasound**

In patients with AIT type 1, US shows a diffuse or nodular goiter (fig 13), while in patients with AIT type 2, ultrasound shows a normal or small diffuse goiter. US-findings of IFN-induced thyroid disease are similar to the entities observed without IFN.

**Color Doppler imaging**

CDI has been shown to be a useful tool in differentiating type 1 AIT from type 2. Using a classification that subdivides CDI features into four patterns (0–III), some

---

**Fig 13.** A 69 year old male patient with known cardiac disease being treated with Amiodarone. Patient had initially hyperthyroidism and hence was sent for a thyroid US which showed an echogenic appearing thyroid lobe with no nodules.
authors have reported that CDI showed mild to markedly increased flow in the thyroid in patients with AIT type 1, whereas flow within the thyroid is markedly decreased or absent in patients with AIT type 2 [91-93]. Moreover, the analysis of nodular and extranodular blood flow was useful in the definition of the underlying thyroid diseases in type 1 AIT, being able to differentiate toxic multinodular goiter and toxic adenoma from Graves’ disease [91]. This differentiation appeared to be of clinical relevance regarding therapeutic choice and outcome [94,95]. Separate evaluation of parenchymal blood flow from that of nodules may prove beneficial in the diagnosis of underlying thyroid diseases in patients with type 1 AIT [91].

In IFN-induced thyroid disease, a reduction in echogeneity suggestive for a destructive process of the thyroid gland was observed on US even before changes in thyroid function of antibody status could be measured. Risk factors for the development of thyroid dysfunction were age, female gender, pre-treatment thyroid volume, pre-existing thyroglobulin/thyroid peroxidase antibodies and viral load. Power-Doppler ultrasound could reliably differentiate between destructive thyroiditis and Graves’ disease [90].

**Elastography**

To date, no specific papers, to the best of our knowledge have been published focusing on Amiodarone induced thyrotoxicosis

**Contrast enhanced ultrasound**

To date, no specific papers, to the best of our knowledge have been published focusing on Amiodarone induced thyrotoxicosis

**Tips and tricks**

Scintigraphy can differentiate between the two forms of AIT with positive scans seen in AIT I and negligible uptake seen in AIT II [96].

**h) Acute thyroiditis**

Acute thyroiditis is a very rare disease, predominantly occurring in immuno-suppressed patients. It may develop locally or hematogenically in sepsis. The classical symptoms of a florid inflammation with fever, sore throat, painful swelling, reddening of the skin and lymph node enlargement can be found [38]. Acute suppurative thyroiditis is uncommon due to the excellent lymphatic drainage, encapsulation, and high iodine content in the gland. It usually affects children and young adults with congenital fourth branchial pouch sinus tracts [27].

**Conventional B-mode ultrasound**

Conventional US scan depicts ill-defined thyroid borders, the echo texture is inhomogeneous; hypoechoic and anechoic areas represent colliquation. In most cases, lymph nodes with inflammatory changes may be demonstrated [38]. US findings are nonspecific and the gland may appear enlarged and hypoechoic due to inflammation. Focal fluid collection with bright echoes in it from gas can suggest an abscess [48].

**Color Doppler imaging**

May show normal or increased blood flow.

**Elastography**

To date, no specific papers, to the best of our knowledge have been published focusing on acute thyroiditis

**Contrast enhanced ultrasound**

To date, no specific papers, to the best of our knowledge have been published focusing on acute thyroiditis

**Tips and tricks**

FNA confirms the diagnosis by identifying the causative pathogen. If clinical and sonographic findings are inconclusive, other forms of thyroiditis or malignant tumor might have to be excluded by cytological examination. Acute thyroiditis may usually be cured by antibiotic therapy. In advanced stages with abscess formation, local drainage or even surgery might become necessary [38,97].

**Conclusion**

A wide spectrum of DTD’s affect the thyroid gland and US is not generally required for the diagnosis of DTD. In some cases, like Hashimoto’s thyroiditis, US helps exclude focal thyroid disease in these patients and in assessing the size of the thyroid gland. US contrast and elastography is in its infancy and larger studies are needed to evaluate the utility of these new techniques in diffuse thyroid diseases. We also refer to the recently published World Federation in Ultrasound and Medicine (WFUMB) thyroid elastography guidelines [98].

**Conflict of interest:** none

**References**

2. Haugen BR, Alexander EK, Bible KC, et al. 2015 American Thyroid Association Management Guidelines for Adult Patients with Thyroid Nodules and Differentiated Thyroid Cancer: The American Thyroid Association Guidelines Task Force on Thyroid Nodules and Differentiated Thyroid Cancer. Thyroid 2016;26:1-133.
3. Thyroid/Parathyroid Ultrasound. American College of Radiology Practice Guideline. In: http://www.acr.org/Sec-
42. Vanderpump MP, Tunbridge WM. Epidemiology and prevention of clinical and subclinical hypothyroidism. Thyroid 2002;12:839-847.
79.  