Thyroid lipomatosis in a 36-year-old patient with rheumatoid arthritis and a kidney transplant

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Summary

Thyroid lipomatosis is a rare disease, as a total of 20 cases have been described in the literature. It is characterized by diffuse infiltration of the stroma by mature adipose tissue and by progressive growth that produces different degrees of compressive symptoms. Our aim is to present the case of a 36-year-old woman who consulted because of dyspnea caused by a multinodular goiter. She underwent surgery with the presumptive diagnosis of a malignant neoplasia, but the pathological examination of the surgical specimen established the diagnosis of thyroid lipomatosis.

Learning points:

• Thyroid lipomatosis is a rare, benign disease characterized by diffuse infiltration of the stroma by mature adipose tissue.

• The pathophysiology of diffuse proliferation of adipose tissue in the thyroid gland is unclear.

• Thyroid lipomatosis is clinically manifested by a progressive enlargement of the thyroid that can involve the airway and/or upper gastrointestinal tract, producing dyspnea, dysphagia, and changes in the voice.

• Given the rapid growth of the lesion, the two main differential diagnoses are anaplastic carcinoma and thyroid lymphoma.

• Imaging studies may suggest a differential diagnosis, but a definitive diagnosis generally requires histopathological confirmation after a thyroidectomy.

Background

Thyroid lipomatosis is a rare disease hitherto described in only 20 patients. It is characterized by diffuse infiltration of mature adipose tissue in the thyroid gland, resulting in its progressive enlargement and potentially to the development of compressive symptoms. While some theories attempt to explain this disease’s pathophysiology – such as through embryological origin of the adipose tissue, adipose metaplasia in response to hypoxia, or senile involution – the etiology of thyroid lipomatosis is not entirely clear. Imaging studies such as ultrasound, CT scan, and MRI may suggest the presence of adipose tissue in the gland, but a definitive diagnosis is confirmed by pathological study of the surgical specimen. In some cases, the diagnosis may be suggested by fine-needle aspiration (FNA) results.

Case presentation

We present a 36-year-old woman who was referred to an endocrinology clinic for dyspnea and goiter.

Her past medical history showed that she had epilepsy since childhood and that she was currently being treated with phenobarbital and lamotrigine. At age 14 years, she was diagnosed with rheumatoid arthritis, for which she had received treatment with glucocorticoids, methotrexate, gold salts, and infliximab. At age
24 years, she was diagnosed with idiopathic bone marrow aplasia, which was treated with cyclosporine, antithymocyte globulin, and multiple blood transfusions. During a hospital admission for bone marrow aplasia, she developed nephrotic syndrome and renal failure. Her kidney biopsy showed secondary amyloidosis. At age 28 years, she started hemodialysis and underwent a splenectomy for splenomegaly of unknown etiology. Two months later, she underwent a kidney transplant with her mother as the donor. Since then, she received immunosuppressive therapy with methylprednisolone, tacrolimus and azathioprine. Her reproductive history showed a first-trimester spontaneous abortion and a preterm delivery due to proteinuria at 34 weeks of gestation. The only relevant family history was her mother with hypothyroidism.

On admission, the patient reported dyspnea while in a supine position for the previous few nights. Her physical examination showed signs of exogenous hypercortisolism and bilateral cubital deviation of the first phalanges of her fingers. Her vital signs were normal. She presented with an enlarged nodular thyroid gland that was approximately 80 g in mass, mobile, and nontender, with a clearly palpable 30 mm nodule in the right lobe and a 15 mm nodule in the left lobe. No lymphadenopathies were evident to palpation.

The notes in her case file showed that the patient had been assessed for a 60 g goiter 2 years before. A thyroid ultrasound showed a 21 mm multiloculated cystic nodule in the right lobe and a hyperechoic, 20 mm solid nodule with cystic areas in the left lobe. No lymph nodes were detected. Her laboratory tests at that time showed anemia and normal thyroid function. A FNA of the right lobe nodule indicated nodular hyperplasia (Bethesda category II) consistent with a benign nodule. Neck X-rays did not show deviation of the trachea. The patient did not return for follow-up.

**Investigation**

A neck computed tomography (CT) scan was performed, showing a diffuse enlargement of the thyroid gland, predominantly of the right lobe. This enlargement invaded the prevertebral space up to the hyoid cartilage, displacing but not compressing the trachea. The thyroid also showed low tomographic density – still higher than that of adipose tissue – with internal blood vessels and septations (Fig. 1A and B). The routine laboratory workup and thyroid function tests were normal (Table 1).

**Treatment**

A total thyroidectomy was performed.

**Table 1** Laboratory results.

| Parameters (NV)     | Results       |
|---------------------|---------------|
| Hematocrit (37–47%) | 36.5%         |
| Hemoglobin (11.5–16 g/dl) | 11.8 g/dl   |
| White blood cells (5000–10 000/mm³) | 7920/mm³ |
| Glycaemia (70–110 mg/dl) | 86 mg/dl    |
| Creatinine (0.5–1.2 ng/dl) | 0.89 ng/dl |
| Sodium (135–145 mmol/l) | 139 mmol/l  |
| Potassium (3.5–5 mmol/l) | 4.6 mmol/l  |
| Total bilirubin (0.1–1.4 ng/dl) | 0.3 ng/dl |
| Alkaline phosphatase (31–100 UI/l) | 139 UI/l |
| Aspartate aminotransferase (AST, 10–42 UI/l) | 14 UI/l |
| Alanine aminotransferase (ALT, 10–40 UI/l) | 9 UI/l |
| Albumin (3.2–5 g/dl) | 4 g/dl       |
| Calcium (8.5–10.5 mg/dl) | 9.4 mg/dl   |
| Phosphorus (2.5–4.5 mg/dl) | 3.1 mg/dl |
| Magnesium (1.7–2.2 mg/dl) | 1.5 mg/dl   |
| Thyrotropin (0.47–4.64 mU/ml) | 0.8 mU/ml |
| Free thyroxine (0.7–1.8 ng/dl) | 1 ng/dl |
| Anti-TPO (0.9 UI/ml) | 0.9 UI/ml   |

NV, normal values; anti-TPO, anti-thyroperoxidase antibody.
Outcome and follow-up

The pathology results were as follows. Macroscopic examination showed that the thyroid gland had a mass of 144 g and measured $98 \times 95 \times 45$ mm (length $\times$ width $\times$ depth). The section surface of the right lobe showed a soft, pale, greyish-yellow tissue with a cystic area of $20 \times 10$ mm. The left lobe showed similar characteristics with a cystic area of $21 \times 20$ mm (Fig. 2A). Microscopic examination revealed that normal thyroid tissue was replaced by mature adipose tissue except for a small remnant of medium-sized normal thyroid follicles containing colloid and lined by a single layer of cuboidal follicular cells (Fig. 2B). A pale, pink, amorphous material was deposited in the extracellular space and surrounded the thyroid follicles, which showed an apple-green birefringence in polarization microscopy with Congo red staining and positive immunostaining for amyloid A (Fig. 2C). The diagnosis of thyroid lipomatosis and amyloidosis was established.

Levothyroxine was started at a dose of 100 $\mu$g/day orally.

Discussion

Thyroid lipomatosis is a rare, benign disease characterized by the enlargement of the thyroid produced by mature adipose tissue diffusely infiltrating the stroma. Only 20 cases have been reported in the literature (Table 2), 18 of which were confirmed by pathological examination. This disease does not show gender predominance (11 males and nine females), and while it affects mostly middle-aged patients (mean age at diagnosis: 44 years), it encompasses patients with a wide range of ages (11–77 years).

In normal thyroids, adipose tissue is distributed under the capsule and along vessels. Fat-containing lesions of the thyroid are seen as lesions containing macroscopic mature fat or lesions rich in microscopic intracellular fat vacuoles, or lesions that are referred to as clear cell or lipid-rich neoplasms (1). Clear cell changes can occur in thyroid neoplasms of various microscopic types and as a consequence of a variety of mechanisms, which include cytoplasmic vesicles (of mitochondrial, reticulum endoplasmic, or Golgi origin) and accumulation of glycogen, lipid, thyroglobulin or mucin (2).

The amount of adipose tissue in the thyroid can be increased in certain disorders, such as Hashimoto’s thyroiditis, amyloid goiter containing adipose infiltration, heterotopic adipocytes, intrathyroidal thymus, and parathyroid lipoma. Some malignant thyroid lesions, such as encapsulated papillary thyroid carcinoma and thyroid liposarcoma or benign lesions, such as thyroid adenolipoma or thyroid lipoma, can also be associated with increased intrathyroidal adipose content. Thyroid lipomas are characterized by the presence of a capsule, which is absent in diffuse thyroid lipomatosis and is the main differentiating feature (3).

The pathophysiology of diffuse proliferation of adipose tissue in the thyroid gland is unclear. Some authors suggest that heterotopic groups of fat cells are included in the thyroid gland during embryogenesis,
| Reference | Age, sex | Thyroid function | Clinical Features | Diagnostic Procedures | Pathological Features |
|-----------|----------|------------------|-------------------|----------------------|----------------------|
| (11)      | 32, M    | NS               | Diffuse goiter for 3.5 years, local compression | NS                  | Thyroid weight 500 g. Diffuse infiltration of fibrofatty tissue, irregular follicles, focal fibrosis |
| (16)      | 11, F    | NS               | Diffuse goiter since birth, multiple lipomas | NS                  | Lobules with fibrous septa, diffuse fat infiltration, small thyroid follicles |
| (17)      | 15, M    | Normal           | Diffuse goiter since birth | NS                  | Thyroid weight 253 g. Diffuse fat infiltration Diffuse fat infiltration |
| (18)      | 58, M    | Hyperthyroidism  | Diffuse goiter for 5 years, local compression. Tuberculosis | NS                  | Thyroid weight 120 g. Diffuse fat infiltration |
| (19)      | 51, F    | Normal           | Diffuse goiter for 3 years | NS                  | Thyroid weight 415 g. Diffuse fat infiltration |
| (20)      | 73, F    | Normal           | Diffuse goiter for 4 years. Colon cancer | NS                  | Thyroid weight 700 g. Thyroid follicles in close proximity to adipose |
| (21)      | 12, M    | Normal           | Right neck mass for 8 years | NS                  | |
| (12)      | 77, F    | Normal           | Cervical mass for 50 years | US: hyperechoic mass. CT: hypo and isodense mass with slight contrast enhancement. MRI: heterogenous mass without regional invasion by the tumor | |
| (4)       | 38, M    | Normal           | Diffuse goiter for 7 years | Tc99-Scintigraphy: bilateral diffuse enlargement | Thyroid weight 465 g. Diffuse infiltration of mature fat between normal follicles. Amyloid (−) |
| (10)      | 67, F    | Normal           | Nodular goiter for 5 years. Morbid obesity, diabetes, CKD | Tc99-Scintigraphy: Cold 2 cm-nodule in the left thyroid | Thyroid weight 41 g. Diffuse infiltration of mature fat, adenomatous nodules, fibrosis |
| (10)      | 59, F    | Hypothyroidism   | Nodular goiter for 6 years. Renal transplantation | Tc99-Scintigraphy: a 1.5-cm hot nodule in the right thyroid. FNA: follicular cell nodule with cystic changes | Thyroid weight 56 g. Diffuse fat infiltration of thyroid including follicular adenomas, papillary thyroid carcinoma |
| (22)      | 64, M    | Normal           | Thyroid enlargement and dyspnea. Psoriasis and CKD | US: heterogeneous echogenicity. Tc99-Scintigraphy: reduced uptake. CT: goiter, slightly compressing the airway. Low density. FNA: fat cells between normal follicles, without any sign of hyperplasia, malignancy, or amyloid deposition | NS |
| (23)      | 45, M    | Normal           | Goiter for 4 years, dyspnea the last six months | CT: compression of the trachea | Thyroid weight 130 g. The interfollicular stroma showed infiltration by mature adipose tissue. Amyloid (−) |
| (9)       | 40, F    | Hyperthyroidism  | Soft multinodular goiter involving both lobes | US: nodules with cystic components with septations. FNA: colloid goiter | Stroma replaced by adult adipose tissue, fibrous septa separating the adipose tissue. Amyloid (−) |
| (7)       | 43, M    | Normal           | Asymmetrically enlarging thyroid gland. CKD in dialysis | US: heterogeneous gland. Tc99-scintigraphy: diffuse uptake and a cold area in the superior part of the left lobe | Thyroid weight 160 g. Sparse thyroid follicles in dense lipoid stroma. Amyloid (+) |
| (8)       | 52, F    | Normal           | Dyspnea in 3 years | US: Increase in echogenicity and attenuation of sound typical of fat. CT: −30 to −40 HU. Mass effect on the hypopharynx | |
| (13)      | 55, M    | Normal           | Goiter enlargement for two years, cough but no respiratory difficulty. Crohn, AA | FNA: droplets of fat. CT: −40 HU, tracheal deviation but no compression. US: hyperechoic | Thyroid weight 148 g. Extensive infiltration of the gland by adipose tissue. Amyloid (+) |
but others believe that a metaplasia from stromal fibroblasts is produced in response to hypoxia or senile involution (4, 5). Lau et al. (6) have recently published a possible relationship between the mutation of the mitochondrial protein succinate dehydrogenase-subunit B and the abnormal differentiation of adipose tissue in thyroid lipomatosis.

Thyroid lipomatosis is clinically apparent due to a progressive enlargement of the thyroid that can involve the airway and/or upper gastrointestinal tract, producing dyspnea, dysphagia, and changes in the voice. Physical examination usually shows a soft, non-tender goiter that is nodular or diffuse (7). In most cases, tests show normal thyroid function, but both hyperthyroidism and hypothyroidism have been described in a few patients.

Given the rapid growth and the tumor size, the two main differential diagnoses, are anaplastic carcinoma and thyroid lymphoma. Imaging studies are helpful to establish the diagnosis. Ultrasound examination shows an enlarged thyroid with a diffuse increase of echogenicity and an echo attenuation typical of adipose tissue. CT scans show an enlarged thyroid with well-defined limits and diffuse infiltration by fat with negative density in the range of adipose tissue (K30 to K40 Hounsfield units). Extension outside the thyroid bed can involve the upper mediastinum and the retropharyngeal compartment, compressing or displacing the surrounding airway and vascular structures (8).

FNA cytology can suggest a diagnosis of thyroid lipomatosis because it usually describes the presence of abundant fat cells among normal follicular cells (7). Although there are no clear guidelines to establish a diagnosis of thyroid lipomatosis with this method, the cytological findings are specific enough to exclude anaplastic carcinoma and thyroid lymphoma (9).

In a pathological examination, the thyroid reveals a section surface with a pale grey, yellowish, or tan color and, in some goiters, a focal cystic degeneration (10). Some of the published cases have reported massively enlarged thyroids with masses of 500–700 g (11, 12). Microscopic examination confirms the diffuse infiltration of mature adipose tissue among normal thyroid follicles. Fibrosis of the stroma and lymphocytic infiltration have also been occasionally described (10).

Our patient showed amyloid deposits, which had been previously reported in two patients with secondary amyloidosis associated with chronic renal failure and hemodialysis in one case, and with Crohn’s disease and kidney transplant in the other (7, 13). Two other reports describe amyloid goiters with diffuse infiltration by adipose tissue (14, 15). The authors attribute the increase in size to...
amyloid deposits – in spite of the infiltration by fat – which underscores the lack of clear criteria to differentiate thyroid lipomatosis with amyloid deposits from amyloid goiters with fat infiltration. In our case, based on the pathology results that showed predominance of fat infiltration over amyloid content, we concluded that the amyloid deposit was secondary to the patient’s systemic amyloidosis but that the increase in thyroid size was due to the infiltration by adipose tissue.

The aim of this paper was to describe a rare thyroid disease in a patient with secondary amyloidosis associated with rheumatoid arthritis and kidney transplantation. Although this association has already been reported, only a small number of cases have been described, and the relationship between these disorders remains unclear. It is important to underscore that the rapid growth and size of these lesions mandate a careful and prompt evaluation to exclude malignant diseases such as anaplastic carcinoma and thyroid lymphoma.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Patient consent
Written informed consent has been obtained from the patient for publication of the case report and accompanying images.

Author contribution statement
S Bell and G A Sosa were responsible for the conception of the work and acquisition of the data, conducting the literature review, and were the authors of the draft and final manuscript. A V Jaen was responsible for the conception of the work, interpretation of the data, critical revision and the draft and final manuscript. M F Russo Picasso was responsible for the acquisition, analysis and review of the pathological samples, and reviewed authors of the draft and final manuscript. A V Jaen was responsible for the acquisition of the data, conducting the literature review, and were the S Bell and G A Sosa were responsible for the conception of the work and thyroid lymphoma.

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