Primary thyroid tuberculosis: a rare etiology of hypothyroidism and anterior cervical mass mimicking carcinoma

Tuberculose tireoidiana primária: rara etiologia de hipotireoidismo e massa cervical anterior mimetizando carcinoma

Bradley Paulino da Silva¹, Erico Gurgel Amorim¹, Elizabeth João Pavin¹, Antonio Santos Martins², Patrícia Sabino de Matos³, Denise Engelbrecht Zantut-Wittmann¹

ABSTRACT
Objective: The involvement of the thyroid by tuberculosis (TB) is rare. Hypothyroidism caused by tissue destruction is an extremely rare report. Our aim was to report a patient with primary thyroid TB emphasizing the importance of diagnosis, despite the rarity of the occurrence. Case report: Women, 62 years old, showing extensive cervical mass since four months, referring lack of appetite, weight loss, dysphagia and dysphonia. Laboratorial investigation revealed primary hypothyroidism. Cervical ultrasound: expansive lesion in left thyroid lobe, involving adjacent muscle. Computed tomography scan: 13 cm diameter cervical mass with central necrosis. Fine needle biopsy: hemorrhagic material. Surgery: total thyroidectomy, left radical neck dissection and protective tracheotomy. The pathological examination showed chronic granulomatous inflammatory process with areas of caseous necrosis and lymph node involvement. The thyroid bacilloscopy was positive. Pulmonary disease was absent. The patient was treated with antituberculosis drugs. Conclusions: Thyroid TB is not frequent, and should be considered as differential diagnosis of hypothyroidism and anterior cervical mass. Arq Bras Endocrinol Metab. 2009;53(4):475-8.

RESUMO

INTRODUCTION
Tuberculosis (TB) has been described in many parts of the body, but the involvement of the thyroid gland is rare. The first description of thyroid TB in the literature was reported by Lebert, in 1862, in a patient with miliary TB. In 1893, Bruns described primary TB...
Thyroid tuberculosis of the thyroid gland with cervical lymph node involvement, without pulmonary disease (1,2,3). The scientific literature reports about 200 cases of TB in the thyroid, most with primary focus outside the thyroid gland (4,5).

The TB bacillus reaches the thyroid through hematogenous and lymphogenous dissemination or directly by cervical lymphadenitis. The clinical presentation is variable, with subacute or chronic thyroid enlargement, but most cases manifest itself as a nodule or caseous abscess (6). The diagnosis can be made by bacilloscopy (Ziehl-Neelsen staining for acid-fast bacilli) and culture of material collected through a fine needle aspiration (FNA) or histopathologic exams (3,6,7).

We described the case of a patient with primary thyroid TB associated with hypothyroidism, cervical lymph node and thyroid mass mimicking thyroid carcinoma, without lung injury.

CASE REPORT

Female patient, 62 years old, presented with a four-month history of progressively growing painless left cervical mass, anorexia, 6 kg weight loss, dysphonia and dysphagia. Past medical history included diabetes mellitus and hypertension in the last eight years, and a primary hypothyroidism in the last three months. Medications included metformin, captopril, NPH insulin, AAS and L-Thyroxin (50 μg/day). She denied family history of thyroid disease, TB or contact with TB patients.

Physical examination showed good general condition with no fever, and normal cardiovascular and pulmonary findings. Head and neck examination revealed a 10 x 4 cm solid and hard left cervical mass in levels III and IV, in continuity with left lobe of the thyroid gland. Laboratory evaluation: TSH = 10.3 IU/mL (VR = 0.41-4.5), free thyroxin (FT₄) = 0.96 ng/dL (VR = 0.9-1.8), negative antithyroperoxidase (TPOAb, VR ≤ 76 IU/mL) and antithyroglobulin antibodies (TgAb, VR ≤ 120 IU/mL). Head and neck endoscopy was normal, except for left vocal cord paralysis. Neck and thyroid ultrasonography revealed enlarged thyroid lobes (more pronounced in the left), and multiple enlarged left cervical lymph nodes with sternocleidomastoid muscle involvement and anterior left submandibular gland displacement. The cervical tomography showed a left neck mass of coalescent nodes, measuring 13 x 4 x 4 cm, involving levels IB, II, III and IV left, in continuity with the sternocleidomastoid muscle and the left lobe of the thyroid and with mediastinal extension. Chest radiography and esophagoscopy were normal.

The FNA, done three times, was non-conclusive, with no evidence of malignant cells.

Despite the negative initial biopsy and frozen section, and with the presumptive diagnosis of malignant thyroid tumor based in the patient’s medical history and in the surgical findings, the patient was submitted to total thyroidectomy and left radical neck dissection. Right and left upper parathyroid glands were identified and preserved as well as both recurrent nerves. However, a protective tracheotomy was felt to be necessary. Surgical findings included a bilateral hard thyroid mass and multiple involved left neck nodes level I thru VI, fixed to sternocleidomastoid muscle and internal jugular vein at levels III and IV.

The pathological examination showed little residual thyroid tissue, with extensive area of white tissue, elastic, joined to the soft tissue around and multiple yellowish nodules. The optical microscopy showed chronic granulomatous inflammatory process with areas of caseous necrosis (Figures 1A and 1B) and calcification, compatible with TB etiology. The cervical lymph nodes at level I were affected and the cytological smear presented acid-fast bacilli. The patient was treated with rifampin, isoniazid and pyrazinamide for nine months. The HIV serology proved negative. There was no clinical recurrence in the following two years.

DISCUSSION

The diagnosis of TB in the thyroid is rarely performed by preoperative clinical suspicion (2). The clinical presentation can mimic cancer, similar to the present report, bleeding or abscess in the thyroid and, in general, the diagnosis is made after biopsy (3,6). In postmortem studies, TB in the thyroid was recognized in 0.003 to 0.1% of the cases. Graham (1932), found 21 cases of thyroid TB among the 20,758 partial thyroidectomy specimens (2,3). However, thyroid foci of TB can happen in 14% of miliary TB (8).

The cause of the rarity of thyroid involvement in TB is unknown, with the following possible explanations: bactericidal thyroid gland colloid action, extremely high thyroid blood flow, increased phagocyte activity in hyperthyroidism, and antituberculous action of the thyroid hormone (6,9). The TB bacillus reaches the thyroid through hematogenous and lymphogenous...
dissemination or directly by cervical lymphadenitis. Microscopically, there is destruction of thyroid tissue by tuberculous granuloma (3,6).

The histopathological aspect of the disease varies depending upon the patient immunity. The disease may present as multiple granulomatous foci throughout the gland or miliary TB, or as a mass with caseous necrosis simulating a goiter with abscess formation, or even as a chronic granulomatous process with fibrosis of the gland. The differential diagnosis of the histological picture should be done with sarcoidosis and subacute (giant cells) thyroiditis (3,8,9). Many cases of primary thyroid TB reported at the beginning of the 20th century had characteristics of giant cell thyroiditis (4), a fact later reassessed in the literature (10). The three criteria for diagnosis of TB in the thyroid, described by Seed, in 1939 included: presence of acid-fast bacilli, necrosis or abscess in the gland and tuberculous focus outside the thyroid (not essential criterion) (2,4,6). The usual result of the involvement of the thyroid is focal caseous TB, presenting as localized swelling, cold abscess, nodule with or without cystic component, multinodular goiter and – more rarely – as acute abscess, possibly confused with carcinoma (9,11). Fibrosis associated with TB is described and adherence to the underlying tissue can cause dysphagia and dyspnea (2,6).

Eventually, the patient may be asymptomatic (9). The clinical presentation may be acute by abscess or thyroiditis, but is often subacute. The duration of symptoms varies from two weeks to one year, with reports of dysphagia, dyspnea and dysphonia due to paralysis of laryngeal nerve. The thyroid affected is increased in volume, hard consistency, adhered to the underlying tissues (2,3). The thyroid function is preserved in the vast majority of cases reported (10) and the description of cases with thyroid hormones abnormalities is extremely rare (9). The description of thyrotoxicosis, generally at the beginning of glandular involvement, is reported as “a clinical syndrome of hyperthyroidism” since the beginning of the last century (2,12). The hypothyroidism is caused by extensive glandular destruction by caseous necrosis (3,6). In recent review of the literature, only one case of hypothyroidism due to thyroid TB was reported (9,13).

Furthermore, the ultrasonographical findings includes a heterogeneous mass or multiple hypoechoic lesions in the thyroid associated with cervical lymph node enlargement; the computed tomography may reveal suggestive image of abscess with increased peripheral density (2,14). As a diagnostic method, the FNA can be effective and confirm the presence of TB in the thyroid, through collecting material for bacilloscopy and culture (7,9). Polymerase chain reaction analysis to detect DNA of Mycobacterium tuberculosis can be done in the cytology material (3,4).

The preoperative diagnosis is rare and it is usually made after surgical resection of the lesion. The treatment of choice is based on surgery and anti-TB drugs administration (8,9). Since the diagnosis is difficult to be performed before surgery and considering the differential diagnosis of cancer, most cases are treated with thyroidectomy (2). The preoperative diagnosis allows
more limited surgical approach, such as abscess drainage (3,15). The use of anti-TB drugs is the basis of thyroid TB treatment and its effectiveness does not differ in relation to other organ sites of involvement of TB (9,15).

In retrospect, the medical history of this patient suggested thyroid TB in the differential diagnosis (multiple negative FNA and negative frozen section), except for the rapid thyroid mass enlargement, multiple left neck nodes and left vocal cord paralysis (all symptoms resembling thyroid carcinoma). However, like the usual literature, the diagnosis was made only with the histopathological examination. Additionally, three months before the complaints of enlargement of the thyroid, the patient had developed primary hypothyroidism, presented with undetectable serum levels of antithyroperoxidase and antithyroglobulin antibodies, without thyroid lymphocytic infiltration in the cytological and histopathological specimens, facts that point out to non-autoimmune cause of gland dysfunction. Thus, the primary hypothyroidism is due to destruction of the thyroid parenchyma by TB infectious process, which is rarely reported in the literature. Clinical, laboratorial and radiological approaches followed the steps described in the literature, and the treatment succeeded by thyroidectomy and administration of anti-TB drugs.

In conclusion, TB in the thyroid is not frequent, but should be considered as differential diagnosis of anterior cervical mass with acute or subacute growth. The diagnosis can be achieved through cytology by FNA, but generally it is reached after surgical resection of the lesion and pathological examination. The treatment is based on surgery and anti-TB drug administration. Additionally, hypothyroidism associated with primary thyroid TB is rare, but it could be seen in suggestive cases.

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REFERENCES