Ectopic thyroid and Hashimoto’s thyroiditis arising from a thyroglossal duct cyst: a case report

Tiroglossal duktus kistinde ektopik tiroidit ve Hashimoto tiroiditi: Olgu sunumu

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A seventy-five-year-old male patient presented with a palpable smooth mass that moved upwards on swallowing, extending from the hyoid bone to the cricoid cartilage. Ultrasonography, scintigraphy, and thyroid hormone measurements showed the mass free from the thyroid gland. Following a diagnosis of infected thyroglossal duct cyst by fine needle aspiration biopsy, the mass was removed by surgery. Histopathologic diagnosis was Hashimoto’s thyroiditis that developed from the ectopic thyroid tissue on the wall of thyroglossal duct cyst.

Key Words: Biopsy, needle; thyroglossal cyst/pathology/surgery; thyroid gland/pathology; thyroiditis, autoimmune/pathology/complications/surgery.

Thyroglossal duct cysts and fistulas may occur at any place from the foramen ecum at the base of the tongue to the pyramidal lobe of the thyroid.[1,2] This developmental abnormality is frequently encountered in childhood during the first five years of life, but may also be detected in adults.[3,4] Cysts presenting with intralingual or suprahroid localizations may be associated with dysphagia, hoarseness, and altered voice.[5,6] Hashimoto’s thyroiditis is an autoimmune disorder frequently seen in females.[7] Although it usually has an insignificant clinical course, symptoms of hypothyroidism may be the initial manifestations in

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**Öz:** A yetmiş beş yaşına erkek hasta, boyun orta hattında, hyoid kemikinden krikoid kartilajda kadar uzanan ve yutunmakla yakarına hareket eden kitle şikayetile başvurdu. Ultrasonografi, sintigrafi, ve tiroit hormon incelemeleri kitlenin tiroit dokusundan bağımsız olduğunu gösterdi. İnce dene kıyasırayon biopsy is ile enfekte thyroglossal duktus kisti tanı kondu ve kitle ameliyatı çıkarıldı. Histopatolojik tani, thyroglossal duktus kisti duvarında ektopik tiroit dokusundan gelişen Hashimoto tiroiditi şeklinde belirlendi.

**Anahtar Sözcükler:** Biyopsi, iğne; tiroglossal duktus kisti/patoloji/cerrahi; tiroit bezi/patoloji; tiroit, otoimmün/patoloji/komplikasyon/cerrahi.

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approximately 20% of patients, of which only 5% may have hyperthyroidism. Most commonly diffuse goiter and lobular goiter may be found on physical examination; on rare occasions, only one nodule is detected.\(^2\)

In this report, we present a case of Hashimoto’s thyroiditis originating from the thyroid tissue on the wall of a thyroglossal duct cyst. Interestingly, the thyroid gland was found normal at its usual location.

**CASE REPORT**

A seventy-five-year-old male patient was referred to our outpatient clinic in January, 2001. He described a small mass in the middle of his neck for the past 10 years, which enlarged with upper respiratory tract infections and returned to its original size following recovery from the infection. However, there had been a progressive enlargement of the mass after the last infection. Physical examination showed a palpable smooth mass (6x5x4 cm in size) of rubber-like consistency, that moved upwards on swallowing. It extended from the hyoid bone to the cricoid cartilage (Fig. 1). Hemoglobin, hematocrit, and the erythrocyte sedimentation rate were measured as 16.2 gr/dl, 46.7%, and 10 mm/h, respectively. There were no clinical or laboratory signs suggesting hypo- or hyperthyroidism. Serum levels of total T4, free T3, free T4, and TSH were within normal ranges. Serum antithyroid microsomal antibody (240 U/ml) and antithyroglobulin antibody (200 U/ml) titers were significantly increased. Ultrasonographic examination revealed a mass in the midline of the neck with cystic and solid structures. The thyroid gland appeared normal. The mass was completely free from the thyroid gland. Scintigraphy with Tc 99m (3mCi) showed a normal thyroid gland, but an increased uptake of the cystic mass suggesting ectopic thyroid tissue. Fine-needle aspiration biopsy was performed. Cytological examination showed leukocytes and necrotic cells which were small and dysplastic. The number of leukocytes was significantly increased and inflammation was prominent. Histopathologic diagnosis was made as thyroglossal duct cyst.

Total extirpation of the mass was planned. Intraoperatively, the mass was dissected from the surrounding tissues. The thyroid gland was intact; the cyst was attached to the corpus of the hyoid bone and had a duct extending to the base of the tongue. The corpus of the hyoid bone was removed and the ductus was dissected. The mass was removed after placing a bag suture (Fig. 2).

Histopathologic examination of the specimen showed a thin capsule formed by a layer of cubic epithelium. Below this capsule were thyroid follicles. There were scattered lymphoid follicles with germinal centers consisting of Hurthle cells (Fig. 3). A diagnosis of Hashimoto’s thyroiditis was made, which developed from the ectopic thyroid tissue on the wall of thyroglossal duct cyst.

Serum antithyroid microsomal antibody and antithyroglobulin antibody titers fell within normal ranges postoperatively.

**DISCUSSION**

Thyroglossal duct cysts are usually found adjoining the hyoid bone (75%) as asymptomatic masses.\(^3\) If the foramen cecum is patent, then fluid or inflam-
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Infection is associated with 50% of these cysts, being the main reason for patient presentation. An infected thyroglossal duct cyst was also considered in the initial diagnosis of our patient with a swollen mass in the midline of the neck, moving upwards on swallowing. Differential diagnosis should include pyramidal lobe of the thyroid gland, thyroid adenoma, aberrant thyroid tissue, branchial cyst, lipoma, dermoid cyst, and hemangioma.

Hashimoto’s thyroiditis is an autoimmune disorder and a diffuse disease of the thyroid gland. It presents as a chronic inflammatory disease of the thyroid in which cytotoxic T-lymphocytes and antithyroid antibodies play a prominent role. Clinical findings are not significant in many patients. In our case, the patient was euthyroid and clinical findings were not remarkable.

The hallmark of the disease is increased antithyroglobulin antibody and antithyroid microsomal antibody levels in the serum, which are detected in approximately 90% to 100% of patients with Hashimoto’s thyroiditis. In our case, a substantial increase in these antibodies were detected.

Ectopic thyroid tissue may be found anywhere along the embryologic thyroglossal duct. Our literature search revealed two cases of Hashimoto’s thyroiditis that developed from the ectopic thyroid tissue of the thyroglossal duct cyst. In our patient, physical examination and scintigraphy of the thyroid gland were normal. However, increased uptake by the cystic mass indicated the presence of an ectopic thyroid tissue of the thyroglossal duct cyst.

Although rare, thyroid tissues on the wall of thyroglossal duct may lead to malignant transformation of thyroglossal duct cysts. Considering the patient’s age and rapid enlargement of the lesion, a diagnosis of thyroglossal duct carcinoma was also included in the differential diagnosis.

In conclusion, in the presence of a thyroglossal duct cyst, symptoms, findings, and laboratory parameters may indicate Hashimoto’s thyroiditis.

REFERENCES

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