CASE REPORT

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Transient thyrotoxicosis in a patient with a functioning nodule;
A possible occurrence of silent thyroiditis

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A 48-year-old woman with diffuse goiter presented with typical symptoms and signs of thyrotoxicosis. Thyroid scanning with I-123 revealed a localized accumulation of the radionuclide in the left lobe which corresponded to a small nodule later detected by ultrasonography, with suppression of the remaining tissues. Owing to the overall reduced radioactivity in the thyroid, she was suspected of having silent thyroiditis causing thyrotoxicosis. Meanwhile, the thyrotoxicosis subsided concurrently with an increase in radioactivity in the extranodular area that had initially been suppressed. The histology of thyroid tissues obtained at the time of operation revealed follicular adenoma or hyperplasia in the area of the localized I-123 uptake and findings similar to those in Hashimoto's thyroiditis in the remaining tissues, supporting our clinical diagnosis of silent thyroiditis together with a functioning nodule.

Key words: silent thyroiditis, functioning thyroid nodule, thyrotoxicosis

INTRODUCTION

Thyroid scanning with Tc-99m or radioiodine is known to be useful for the differential diagnosis of thyroid diseases causing thyrotoxicosis. We have recently seen a thyrotoxic patient with a functioning nodule, in whom the thyrotoxicosis was caused by concomitant silent thyroiditis. This paper presents her scintigraphic findings, which may support the diagnoses.

CASE REPORT

A 48-year-old woman visited Kyoto University Hospital, because she had noticed general fatigue, palpitation and shortness of breath on effort, excessive sweating, and weight loss of 3–4 kg over past 4 months. Physical examination revealed a diffusely enlarged soft goiter, hand tremor, moist skin, tachycardia (120/min) and systolic hypertension (180/80). Neither exophthalmos nor ophthalmopathy was present. She had no fever or tenderness in the anterior neck region.

In vitro thyroid function tests performed on March 10 revealed high serum T4 concentrations of 16.3 μg/dl (normal range, 5.0–11.0), free T4 of 3.7 ng/dl (normal, 0.99–1.92), T3 of 238 ng/dl (normal, 90–170), and free T3 of 9.0 pg/ml (normal, 2.2–5.0). The serum TSH level was undetectably low (<0.03 μU/ml).

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Fig. 1 The clinical course. As thyrotoxicosis subsided, 123I thyroidal uptake increased. RIU: radioactive iodine uptake.
ml; normal, 0.3–3.9). The serum TBG concentration was 19.2 μg/ml (normal, 14.2–30.5). Antibody titers against thyroglobulin and microsomes were positive, both at dilutions of 1:1600 (normal, both 1: <100). Neither TSH binding inhibitor immunoglobulins (TBII) nor thyroid stimulating antibodies (TSAb) were detected in the serum. CRP was negative. No other abnormal findings were obtained.

Ultrasonographic examination revealed diffuse enlargement of the thyroid and a small, round hypoechoic area 2.0 cm in diameter in the lower and anterior part of the left lobe. Thyroid scanning with 1-123 revealed a small area of localized uptake in the lower part of the left lobe, corresponding to the hypoechoic lesion, with reduced radioactivity in the remaining thyroid tissues.

Overproduction of thyroid hormones by the hot nodule seemed unlikely as a cause of thyrotoxicosis, since 1-123 thyroidal uptake was substantially reduced at 2.7% (24 hours after oral administration of 3.7 MBq of I-123; normal, 7–35) (Fig. 1 & 2). Consequently, silent thyroiditis was thought to be responsible for the thyrotoxicosis, and she received 120 mg/day metoprolol (β-antagonist). The clinical course is illustrated in Fig. 1. Prednisolone was used

Fig. 2 Thyroid scanning with 131I performed on March 28 revealed the presence of a hot nodule in the left lobe with suppression of the remaining tissues (a). The overall 131I thyroidal uptake value 24 hours after oral administration of 7.4 MBq of 131I was 2.7% (normal, 7–35). There was a gradual increase in radioactivity during the course of the illness in the extranodular tissues which had initially been suppressed. The overall 131I thyroidal uptake values at 24 hours were 7.6% on April 27 (b) and 10.5% on August 24 (c).
sized follicles with epithelial hyperplasia were observed in the nodule, while follicular degeneration, lymphocytic infiltration and fibrosis were observed in the remaining tissues (Fig. 3). The pathological diagnoses were follicular adenoma or hyperplasia, and Hashimoto's thyroiditis, respectively.

**DISCUSSION**

The patient was suspected of having Graves' disease at her first visit. A low T3/T4 ratio (14.6)\(^2\), negative TBII and TSAb activities, and low thyroidal uptake of I-123, however, suggested that the thyrotoxicosis was of a destruction-induced type. Subacute thyroiditis could be ruled out, considering her lack of fever or anterior neck pain and negative CRP. Although she was diagnosed as having silent thyroiditis and a functioning nodule, a differential diagnosis from toxic nodular goiter or destruction-induced thyrotoxicosis following hemorrhagic infarction in the functioning nodule\(^4\) could not be definitely made. Results of the patient's history, physical examination and other tests excluded the possibility of other thyroid diseases causing thyrotoxicosis, such as hCG-producing tumors, metastatic functioning thyroid cancer, TSH-producing tumor, pituitary resistance to thyroid hormones, excessive intake of thyroid hormones and struma ovari.

The diagnoses of silent thyroiditis and a functioning nodule became more and more convincing during the course of the illness, because serial changes in serum thyroid hormone concentrations, the I-123 thyroidal uptake value and the thyroid scan image seemed to reflect the healing process of silent thyroiditis. The histological diagnosis was adenoma or hyperplasia with Hashimoto's thyroiditis, supporting the clinical diagnoses. There were none of the cystic lesions in the nodule which can be produced following hemorrhagic infarction.

The reported incidence of adenomas in Hashimoto's thyroiditis is as high as 8.3–25.0\%\(^5\). On the other hand, the association of functioning nodules with Hashimoto's thyroiditis, to our knowledge, has never been reported. Although a functioning nodule coincident with silent thyroiditis may be uncommon, the present case report at least has confirmed the usefulness of thyroid scintigraphy for the diagnosis of thyroid diseases causing thyrotoxicosis.

**REFERENCES**


