Dual ectopic thyroid associated with thyroid hemiagenesis

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Summary
We report a case of a 15-year-old girl with a midline neck mass that was first noted 2 or 3 years previously. She had been treated with levothyroxine (L-T4) for congenital hypothyroidism until 11 years of age. Ultrasonography revealed an atrophic right thyroid (1.0 × 1.6 × 2.6 cm in size) and a mass (2.3 × 1.0 × 3.5 cm in size) in the upper part of the neck. No left lobe of the thyroid was detected. On further evaluation, Tc-99m pertechnetate thyroid scintigraphy and CT showed ectopic thyroid tissue in the lingual region and infrahyoid region. Thus, she was diagnosed as having dual ectopic thyroid and thyroid hemiagenesis. The atrophic right thyroid was thought be non-functional. Treatment with L-T4 was started to reduce the size of the dual ectopic thyroid tissue. This may be the first reported case of dual ectopic thyroid associated with hemiagenesis detected only by ultrasonography.

Learning points:
- Ultrasonography can confirm the presence or absence of orthotopic thyroid tissue in patients with ectopic thyroid.
- The cause of congenital hypothyroidism should be examined.
- Clinical manifestation of ectopic thyroid may appear when the treatment with L-T4 is discontinued.
- Annual follow-up is needed in all children when their thyroid hormone replacement is stopped.

Background
Ectopic thyroid tissue is a rare congenital anomaly characterized by the presence of thyroid tissue in regions other than its normal position, usually from the base of the tongue to the final pre-tracheal position. The prevalence of ectopic thyroid is about 1 per 100,000–300,000 individuals (1). It is very uncommon for ectopic thyroid tissue to be present simultaneously in two regions (2, 3). Thyroid hemiagenesis is also a rare congenital anomaly in which one lobe of the thyroid gland is not developed. It is generally diagnosed incidentally in association with abnormalities of the remaining opposite thyroid lobe, such as hyperthyroidism, hypothyroidism and adenoma. Korpal-Szczyrska et al. (4) detected thyroid hemiagenesis in two out of 4,004 unselected school children (aged 7–15 years) by ultrasonography (i.e. about 1 per 2,000).

Here, we report the case of a patient with dual ectopic thyroid associated with thyroid hemiagenesis.

Case presentation
A 15-year-old girl was referred to our hospital for further evaluation of her 2- or 3-year history of anterior neck swelling. She presented with occasional feelings of discomfort and pressure in the neck and a choking sensation. Her medical history demonstrated hypothyroidism that was detected by neonatal screening (thyroid-stimulating hormone (TSH) level: 78.5 μU/mL; free T4 level: 1.05 ng/dL) and treated with levothyroxine (L-T4) up to 11 years of age (the first dose of L-T4 was 30 μg/day and the final dose of L-T4 was 5 μg/day).
A morphological examination was not performed when her hypothyroidism was first detected. Treatment with L-T4 was discontinued because a euthyroid status was achieved with a small dose of L-T4. She had no history of neck surgery or irradiation. Her family history revealed no other family members with thyroid disease.

**Investigation**

Her height was 165.5 cm and her body weight was 59.9 kg. Her physical examination revealed a soft nontender mass (2×2 cm in size) in the upper part of the anterior neck, but the thyroid gland was not palpable in the normal position. The serum levels of free T4, TSH and thyroglobulin (Tg) were 1.01 ng/dL (normal range: 0.70–1.80 ng/dL), 2.622 μU/mL (normal range: 0.40–4.30 μU/mL) and 24.2 ng/mL (normal range: ≤33.7 ng/mL), respectively. Anti-Tg, anti-peroxidase and anti-TSH receptor antibodies (TRAb) were negative. The serum Tg and TRab were measured because they were useful to distinguish between the various causes of neonatal hypothyroidism. Ultrasonography demonstrated a heterogeneous mass (2.3×1.0×3.5 cm in size) in the upper part of the neck (Fig. 1) and an atrophic right thyroid lobe (1.0×1.6×2.6 cm in size), but no left thyroid lobe was detected (Fig. 2). She was diagnosed as having thyroid hemiagenesis on the basis of these findings. Tc-99m pertechnetate thyroid scintigraphy (Fig. 3) showed uptake in the upper part of the neck and the center of the oral cavity, without uptake in the normal position of the thyroid or other positions. Neck CT (Fig. 4) revealed two hyperdense masses at the base of the tongue and in the pre-tracheal region below the hyoid bone, but not in the normal position. She was additionally diagnosed as having dual ectopic thyroid on the basis of these CT findings. Fine-needle aspiration biopsy of the pre-tracheal mass did not show malignant cells.

**Discussion**

Tc-99m pertechnetate thyroid scintigraphy and CT demonstrated dual ectopic thyroid (lingual and infrahyoid regions), but thyroid tissue in its normal position was not detected by these two procedures. In 2014, Meng et al. (2) studied 42 patients with dual ectopic thyroid including their own six patients and reported that one lesion is in the lingual (sublingual) region and the other is in the subhyoid (infrahyoid) region in about half of the patients. In 2015, Gu et al. (3) identified five patients with dual ectopic thyroid among 42 patients with ectopic thyroid. Among these five patients, four had ectopic thyroid tissue in the lingual and anterior neck regions, one in the lingual...
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region and in the floor of the mouth. Thus, the association of ectopic thyroid tissue with regions above and below the hyoid bone may not be unusual in dual ectopic thyroid. In about 90% of patients, ectopic thyroid is found at the base of the tongue (lingual thyroid) (1). Normal thyroid tissue is absent in 75% of patients with lingual thyroid (1). In dual ectopic thyroid, a normally located thyroid gland is extremely rare. In 1966, Kuehn et al. (5) reported the first case of dual ectopic thyroid with normal pretracheal thyroid. After this report, orthotopic thyroid glands in dual ectopic thyroid have been demonstrated by scintigraphy or ultrasonography. Karakoc-Aydiner et al. (6) reported that repeated ultrasonography showed the disappearance of previously detected orthotopic atrophic thyroid tissue in six patients with ectopic thyroid. Thus, the atrophic right lobe in our patient may disappear in the future because of the non-functional features.

Ectopic thyroid gland typically becomes evident during adolescence or pregnancy since the requirement for thyroid hormones increases during this period. About 55% (23/42) of the patients with dual ectopic thyroid are also reported to seek medical consultation when they are between 11 and 28 years of age (2). The treatment with L-T4 was stopped in our patient at 11 years of age. This stopping might have increased the size of dual ectopic thyroid mass gradually resulting in the symptoms.

A few reports on patients examined by thyroid scintigraphy demonstrated the association of thyroid hemiagenesis with single (7, 8) or dual ectopic thyroid (9). In these reports (7, 8, 9), orthotopic thyroid is clearly demonstrated, whereas ectopic thyroid is slightly visible by scintigraphy. On the other hand, orthotopic thyroid (hemithyroid) was not demonstrated by scintigraphy or CT in our patient, but was demonstrated by ultrasonography. Silva et al. (10) reported that ultrasonography demonstrated thyroid tissue in 6 out of 18 patients scintigraphically diagnosed as having thyroid agenesis. The right thyroid lobe was atrophic and non-functional and had a low iodine concentration. These features may explain why the right thyroid lobe was not visible on CT. Therefore, not only thyroid scan or CT, but also ultrasonography is required to confirm the presence or absence of orthotopic thyroid tissue in patients with ectopic thyroid. This may be the first reported case of dual ectopic thyroid associated with hemiagenesis detected only by ultrasonography.

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 for publication of her clinical details and/or clinical images was obtained from the parent of the patient.

Author contribution statement
S Nakamura wrote the majority of the paper. T Masuda and M Ishimori wrote and contributed as well. Every author played a significant role in drafting this case report.

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