Tracheal stenosis due to huge adenomatous goiter

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Abstract
Background: Some thyroid anaplastic carcinoma and lymphoma have been reported to cause tracheal stenosis or choking. Begin thyroid tumors with dyspnea due to tracheal stenosis are exceedingly rare.

Case presentation: We experienced a huge adenomatous goiter (resected specimen weight: 520g). Total thyroidectomy resolved the dyspnea and there was no tracheomalacia. Postoperative CT confirmed a normal tracheal lumen free from stenosis.

Conclusions: We reviewed the CT findings of thyroid benign tumors with dyspnea and found that tumors occupied the space between the posterior wall of trachea and esophagus leading to tracheal stenosis as in our case. Such tracheal findings would actually lead to dyspnea.

Introduction
Some thyroid malignant tumors and thyroid malignant lymphoma have been reported to cause tracheal stenosis [1,2]. Benign thyroid tumors with dyspnea due to tracheal stenosis are very rare [3]. We experienced a case of a huge benign tumor (resected specimen weight: 520g) with dyspnea due to tracheal stenosis and performed total thyroidectomy. This case report presents the clinical findings, imaging examination, operation and postoperative progress. Moreover, we discuss the tracheal stenosis pattern with dyspnea by referring to various studies.

Case presentation
A 54-year-old Japanese woman consulted our hospital with dyspnea at neck flexion and when supine. Therefore, she went to bed in neck retroflection. In the initial findings, both lobes of the thyroid were diffusely enlarged to the lower jaw (Figure 1). In the CT scan, bilateral lobes of the thyroid tumors were diffusely swollen and had grown into the space between the posterior wall of the trachea and esophagus, leading to the tracheal stenosis by compressing the posterior wall of the trachea (Figure 2). On the blood examination, thyroid function, microsome test, and thyroid test were within normal limits. Thyroglobulin (729.6 ng/ml) was high (normal range: 0-33 ng/ml).

We performed total thyroidectomy with preservation of the bilateral recurrent nerve, main blood vessels and thyroid capsule. However, oozing from the entire thyroid surface continued during surgery. The operation time was 6 hours and the bleeding volume was 1450g. The size and weight of the enblock resected specimen was 13 × 11 × 6 cm and 520g (Figure 3). The postoperative histopathology was adenomatous goiter without malignant cells and chronic thyroiditis (Figure 4). The progress after surgery was good, and there was no dyspnea, hemorrhage or vocal cord paralysis. In postoperative CT, the trachea appeared to be free from stenosis and normal (Figure 5). Postoperative thyroglobulin (10.2 ng/ml) was within the normal range.

Discussion
Huge thyroid malignant lymphoma with dyspnea has been reported...
and Nomizu [4] presented a case with choking. Thyroid lymphoma occurs in chronic thyroiditis [5]. Sato [6] experienced a case of chronic thyroiditis without malignant lymphoma that caused narrowing of the air way, for which emergency tracheotomy was performed. Alfonso [7] demonstrated that the incidence of tracheoesophageal compression in thyroid benign diseases with chronic thyroiditis was 67%. In our case, the histopathological diagnosis was an adenomatous goiter without chronic thyroiditis or thyroid dysfunction, and the microsome and thyroid test results were within normal limits in the preoperative blood examination.

Not all huge thyroid benign tumors with tracheal compression could cause dyspnea. We reviewed the CT findings of thyroid benign tumors with dyspnea in an found that they occupied the space between the posterior wall of the trachea and esophagus leading to tracheal stenosis as our case [6,8]. The above tracheal findings would actually lead to dyspnea.

Agarwal [9] described that the incidence of tracheomalacia with huge goiters was high. Some hospitals performed tracheostomy and inserted a T-tube into the trachea lumen after thyroideectomy for huge thyroid benign tumors [8,10]. In our case we observed no clinical findings of tracheomalacia after surgery. Postoperative CT confirmed a normal tracheal lumen without stenosis.

Declarations

Author’s contributions

Takeshi Kusunoki MD: diagnosis and therapy, composition of this case report. Hirotomo Honma MD: therapy, Yoshinobu Kidokoro MD: therapy, Aya Yanai MD: therapy, Ryo Wada MD: Pathological diagnosis, Katsuhisa Ikeda MD: composition of this case report

Consent for publication

Consent for publication on the clinical details, images and histopathological examination was obtained for this case report. A copy of this document is available for review by the Editor of this journal.

Acknowledgements

The authors thank Mr. Brent Bell for checking English of the manuscript.

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