Introduction

Ectopic thyroid is defined as thyroid tissue not located anterolateral to the second and fourth tracheal cartilages. About 90% of all ectopic thyroid tissues (ETTs) are found along the path of initial embryologic descent of the thyroid gland from the floor of the primitive foregut to its normal pretracheal position. Ten percent have been reported in other anatomical locations, including the mediastinum, trachea, and the heart. ETT is a rare finding in the mediastinum and constitutes only 1% of tumors in this location. We present a case of a 57-year-old female presenting with locally advanced breast cancer and a mediastinal ectopic thyroid.

Case Report

A 57-year-old postmenopausal multiparous female with a history of tuberculous lymphadenitis at 32 years of age, which was treated with a full course of antituberculous treatment, now presented with a lump in the right breast of 6 months. She had no family history of any cancer or thyroid disorder. Fine-needle aspiration cytology from the lump showed ductal carcinoma. Examination revealed a 6 cm × 2 cm hard lump in the upper outer quadrant of the right breast with a movable ipsilateral level 1 lymph node of size 2 cm × 1 cm. General examination including a neck examination was within the normal limits. Digital mammogram showed a normal left breast and BIRDS V lesion of the right breast. Tru-cut biopsy of the lump showed infiltrating duct carcinoma, Grade III and triple-negative. Her hemogram, serum electrolytes, renal and liver functions were normal. Bone scan did not show any evidence of skeletal metastasis. Her chest X-ray showed mediastinal widening. Hence, a computerized tomography scan of the thorax was done, which revealed the enlarged paratracheal nodal mass of size 5.7 cm × 4.4 cm × 2.8 cm in addition to the right breast mass and ipsilateral axillary lymphadenopathy [Figure 1]. Due to the disproportionately large mediastinal lymphadenopathy, in addition to metastasis, possibility of tuberculosis was also considered as the cause for the mediastinal node.

Video-assisted thoracoscopic surgery and frozen section from the mediastinal mass was done which showed normal thyroid parenchyma [Figure 2]. The mass was completely resected. Histopathology examination showed thyroid parenchyma with no significant pathology. Therefore, it was concluded that the mediastinal mass was an intrathoracic thyroid. Her thyroid function test was normal with TSH-0.9 mU/L, T3-105 ng/100 ml, and T4-5.8 ng/100 ml.

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T4-7.5 μg/dL. Thyroid scan was not taken. The clinical stage of the disease was T3N1. She received neoadjuvant chemotherapy with three cycles of 5 fluorouracil, epirubicin, and cyclophosphamide followed by three cycles of docetaxel. Following that, she underwent modified radical mastectomy, and the specimen showed ypT1N0 disease. Following this, she received radiation therapy. At present, she is under follow-up.

Discussion

The thyroid gland originates as a median endodermal thickening on the floor of the primitive hypopharynx about 24 days after fertilization. The primitive gland then descends toward its natural location immediately anterior to the trachea. The developing thyroid gland remains connected to the tongue using a narrow tube, known as the thyroglossal duct. The thyroglossal duct usually involutes at the 6th or 8th week.[1]

Ectopic thyroid is an embryological aberration defined as “thyroid tissue not located anterolateral to the second and fourth tracheal cartilages.” The prevalence of ETT varies between 7% and 10% according to autopsy studies.[4] Ectopic thyroid may develop at various sites from the base of the tongue to any site of the thyroglossal duct due to stoppage of thyroid descent.[1] The cervical midline is the location in 90% of the cases.[2] Less than 1% of all ETT occurs in the mediastinum.[5] Heterotopic thyroid in the mediastinum, pericardium, and heart has probably grown from rudiments of developing thyroid dragged into the chest during the descent of the heart and great vessels with the development of the embryonic neck and the unfolding of the embryo.[5]

Intrathoracic thyroid may be completely asymptomatic, being found incidentally in a chest radiogram performed for other reasons.[3] The symptoms, when produced, are usually related to compression of upper aerodigestive tracts and superior vena cava. Common symptoms are neck mass, shortness of breath, hoarseness of voice, dysphagia, and stridor/wheezing.[6] The diagnostic procedures currently in use for the evaluation of intrathoracic goiter include chest radiography,[7] computed tomography,[8] radionuclide scintigraphy, angiography, and mediastinoscopic biopsy. I-131 scintigraphy can be considered as a diagnostic test only when ETT has to be suspected. Scintigraphy is not always diagnostic because the uptake of I-131 is not always observed in ETT. Patients are usually euthyroid.[9]

Very rarely malignant transformation in ETT can occur.[10] Due to the risks of progressive enlargement, malignant transformation, hemorrhage within the mass, causing respiratory failure and compression of neighboring vital mediastinal organs; these masses should be surgically resected.

Although rare, the possibility of ETT should be considered while evaluating a mediastinal mass. Some of the important differential diagnoses of anterior mediastinal mass include lymphomas, dermoid cysts, and thymic tumors. As in our patient, who has a diagnosis of carcinoma breast, which is otherwise an early-stage lesion, it is extremely important to differentiate it from a metastatic mediastinal lymph node enlargement. Only 2% of patients with breast cancer show mediastinal lymph node metastases.[11] No significant association was found between ectopic thyroid and carcinoma of the breast, after an extensive literature review.

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Conflicts of interest

There are no conflicts of interest.

References