Introduction

Heterotopic thyroid tissue is described to usually lie at the tongue base, while the 10% at the hyoid bone (1). Different localization are also described in cervical regions and in relation with larynx, trachea, oesophagus, mediastinum, diaphragm and the heart.

We report a case of asymptomatic true mediastinal ectopic goitre in a patient with multinodular cervical goitre and previous excision of thyroid nodule.

Case report

A 35-year-old woman is admitted in our department for an asymptomatic multinodular goiter. She underwent thyroid nodule excision four years ago. Hematologic and thyroid function tests were normal. During the preoperative evaluation a chest X-ray showed a shadow in anterior mediastinum on the right (Fig. 1). A chest CT-scan disclosed a non homogeneous 84x51 mm oval mass lying on the ascending aorta (Fig. 2).

Due to the unknown nature of the mediastinal mass, we decided to refer our patient to the Thoracic Surgery Unit. At sternotomy there was no continuity between the thyroid gland and the mediastinal vascularized from the branches of anonymous artery and thoracic vessels. The mass was excised.

Results

The definitive histology revealed an ectopic goiter without findings of malignancy. The postoperative course was uneventful and the patient was discharged on postoperative day 5th.

Thirteen months later, the patient was readmitted to our unit to perform the total thyrodectomy for multinodular goiter.

Discussion and conclusion

According to the criteria of ectopic thyroid tissue, the true intrathoracic goiter is a rare condition. Indeed this definition is often used for both cervical goiter developed into mediastinum and for thyroid tissue not linked to the thyroid. The first condition, known as secondary intrathoracic goiter, is the result of progressive down-growing of a cervical goiter in to the chest behind the sternum (1-3). On the other hand the so-called primary intrathoracic goiter represents the true ectopic thyroid
tissue in the chest as in our case; it differs from the secondary goiter because of the absence of connection with thyroid and on the blood supply from mediastinal vessels (3). Few patients are symptomatic showing cough, wheezing, dyspnea, hemoptysis, recurrent pneumonia, dysphagia, chest pain or superior vena cava syndrome due to the growing of mediastinal mass (1, 3). But most of the cases are asymptomatic (4). Thus the diagnosis is usually incidental, as in our case, during preoperative exams, i.e. chest X-ray, disclosing a tracheal displacement, or compression, calcifications and soft tissue mass.

Further diagnostic tools are needed for the management plane. The scintigraphy can be useful for the differential diagnosis of thymoma and teratoma, but it is not ever diagnostic since the absence of uptake of the $^{131}$I by the mediastinal thyroid tissue (5, 6). Thus, according with the indications to surgical excision for almost all mediastinal masses, a chest CT-scan is required to show the connection with major thoracic vessels and other mediastinal organs (6-9). The resection is advocated like to solve compressive symptoms, and to rule out the diagnosis of malignancy (9, 10).

Although the thoracoscopic excision has been described, the surgical approach is usually a thoracotomy or sternotomy with a mortality rate ranging from 0 to 2% (9-11).

References