Diffuse Lipomatosis of Thyroid Masquerading as Nodular Goitre

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ABSTRACT

Diffuse thyroid lipomatosis is an exceedingly rare histopathological condition of thyroid classically characterized by diffuse swelling and fatty infiltration within the thyroid stroma. We report a case of 40 year old female who presented with a midline swelling in neck since 7 years with a recent history of compressive symptoms. On evaluation, goitre of thyroid gland was revealed on computed tomography. She underwent subtotal thyroidectomy and the final diagnosis of diffuse thyroid lipomatosis was confirmed based on histopathology.

Keywords: thyrolipoma; histopathology; goiter; neck swelling; lipomatosis; thyroid.

INTRODUCTION

Thyroid lipomatosis is defined as a progressive enlargement of thyroid gland by diffuse infiltration of mature adipose tissue intermixed with follicles and lack of encapsulation. [1] It is an extremely rare condition and the first ever case was reported by Dhayagude in 1942. [2] We report a case of thyrolipomatosis in a 40 year old patient presenting with respiratory distress.

CASE REPORT

A 40 year old female patient presented with a history of diffuse midline swelling for the past 7 yrs. With recent increase in the size of swelling, she developed dysphagia, dyspnoea and was admitted in emergency with respiratory distress since 2 days. There were no complains of fever, palpitation, tachycardia or weight loss. Thyroid function tests were within normal limits. Plain radiographs of the chest were unremarkable. Computed tomography scan revealed goitrous enlargement of thyroid gland (Figure-1a and 1b).

Fine needle aspiration cytology (FNAC) was performed in which 10 ml colloid admixed with blood was obtained and was inconclusive. A clinical diagnosis of colloid goitre with tracheal compression was made and subtotal thyroidectomy was performed.
performed. We received thyroidectomy specimen weighing 550gm, comprising of right and left lobe of thyroid measuring 12x7x5 cm and 10x6x6 cm respectively with attached isthmus. The external surface was encapsulated, nodular and the cut surface was tan brown with microcystic change (Figure-2).

A diagnosis of diffuse lipomatosis of thyroid gland was rendered based on these findings.

**DISCUSSION**

Adipose tissue can be seen in parathyroid, thymus, salivary gland, breast or pancreas. However, its presence in thyroid gland is unusual. \[3\] Few adipocytes may be found near the capsule, in the perivascular location or in the connective tissue septa. \[3,4\]

Clinically, it mostly presents as a diffuse enlargement in the anterior cervical region. A large proportion of patients present with hyperthyroidism while others are euthyroid. In our case, the condition of the patient was euthyroid.

Several theories have been proposed to explain the pathophysiology of this condition:

1) Trites et al proposed that thyroid arises from primitive foregut and the presence of these lesions suggest a disturbance in development of primitive foregut. \[5\]

2) Schroder et al suggested that adipose tissue may be a result of metaplasia of stromal fibroblasts due to tissue hypoxia or senile involution. \[6\]

3) Chesky et al and other authors have attributed the lesions to the simultaneous inclusion of fat with striated muscle in thyroid gland during embryogenesis before the development of thyroid capsule. \[7\]

4) Lau et al suggested a possible relationship between diffuse lipomatosis and loss of expression of protein succinate dehydrogenase – B subunit in the follicular or adipose cells. \[8\]

The differential diagnosis of presence of mature adipose tissue in the thyroid include heterotopic nests of adipocytes, diffuse lipomatosis, adenolipoma, amyloid goitre, lymphocytic thyroiditis, intrathyroidthymic or parathyroid lipoma, encapsulated papillary carcinoma and liposarcoma. \[3\]

Thyrolipoma is a well circumscribed, encapsulated nodule composed of thyroid follicles admixed with mature adipose tissue.
while lipomatosis of thyroid is a diffuse process. Similarly, its diffuse nature differentiates it from heterotrophic nests of adipocytes located mainly in the subcapsular areas. Amyloid goitre which can also contain fat cells can be distinguished based on special stain and polarising microscopy. Fat infiltration can be seen in lymphocytic thyroiditis. Parathyroid lipoma should also be considered as a differential diagnosis wherein lobulated adipose tissue is present in the stroma with follicular formations of parathyroid chief cells. 

Our case highlights the importance of evaluation of patients with adipose components in the thyroid in order to exclude other conditions that present as diffuse enlargement of the gland, particularly neoplastic.

CONCLUSION
Diffuse thyroid lipomatosis is a rare entity of unknown etiology and this case point towards significance of including it in the differential diagnosis of a patient presenting with goitre.

Conflict of interest
The authors declare that there is no conflict of interest regarding publication of this paper.

REFERENCES


