A rare case of papillary carcinoma in thyroglossal cyst

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Abstract

Thyroglossal cysts are commonly encountered and may very rarely harbour malignancy. The incidence of carcinoma coexisting in thyroglossal duct cysts is less than 1%, and most common is papillary carcinomas. In the literature to date about 274 cases have been reported.

We present a case of a 40 year woman presenting with swelling in front of the neck since 3 months. On examination there was a small firm, non-tender, midline swelling in the neck. Ultrasound examination and contrast enhanced CT showed heterogeneous soft tissue lesion having solid and cystic components with calcification.

FNAC showed it to be a benign cystic lesion. After surgical resection, histopathology showed fibro collagenous cyst wall showing thyroid follicular cells with few showing squamous metaplasia and a focal area showing papillae lined by cuboidal cells with nuclear overlapping, grooving and clearing of nuclei suggestive of papillary carcinoma of Thyroglossal Cyst.

Keywords: Thyroglossal cyst, Papillary carcinoma, Histopathology.

Introduction

Thyroglossal duct cysts (TGDC) are the most common anomaly in the development of the thyroid gland, which constitutes 70% of midline masses in childhood and in adults about 7% [1]. Only less than 1% of TGDC develop carcinomas [1]. Brentano in 1911 and Uchermann in 1915 are the first to describe a neoplasm in a thyroglossal duct (TGD) remnant [2]. Most cases of Thyroglossal cyst carcinoma diagnosed during the third and fourth decade of life and are rare in children less than 14 years of age. Most patients are asymptomatic[3]

Case Report

A 45 year old female presented with swelling in front of neck of three months duration. It was associated with pain and increase in size since 1 week. The swelling moved with deglutition and protrusion of tongue. On examination, a midline swelling was noticed about 6 cm above the sternal notch, measuring 1x1 cm, with well-defined borders, smooth surface. Overlying skin was pinchable, firm in consistency, moved with deglutition and protrusion of tongue. There was no local rise of temperature. No lymph nodes were palpable.

Investigations

FNAC of swelling: Aspirate yielded 5 cc of straw colored fluid. There were few benign epithelial cells and cyst macrophages seen but no malignant cells were appreciated and was given as Benign Cystic lesion. On repeat USG guided FNAC smears were scanty cellular and reported as Non-diagnostic.

USG Neck: Solid lesion with cystic component was seen. Solid component measuring 18x14 mm with specks of calcification, cystic component measuring 10x8 mm. Both thyroid lobes were normal.
CT Scan of Neck (Plain and Contrast): An ill-defined irregular heterogeneous soft tissue lesion was seen having both solid and cystic component with calcification and enhancement in infrahyoid region indicating Thyroglossal Cyst with Malignant Transformation.

Thyroid Function test was Normal.

Procedure: Surgical resection was performed with Sistrunk Operation.

Histopathology: Gross: Specimen consisted of single piece of grey-white to grey brown tissue measuring 3.5x1.5x1cms consisting of a part of hyoid bone with attached mass. External surface was congested. Cut surface showed grey white well circumscribed lesion with mucoid areas.

Microscopy: Sections studied showed fibro-collagenous cyst wall showing thyroid follicular cells with focal squamous metaplasia. A focal area showed papillae lined by cuboidal cells with nuclear overlapping, nuclear grooving and clearing of nuclei. Tumor was limited within the cyst with no extension to the cyst wall. Cyst wall was seen infiltrated by lymphocytes. Skeletal muscles were seen outside the cyst wall- The histopathological features were consistent with Thyroglossal Cyst with Papillary carcinoma thyroid.
**Case Report**

**Fig- D:** Showing tumor cells arranged in papillary pattern with nuclear overcrowding, optically clear nuclei and nuclear grooving.

**Fig- E:** Showing tumor cells arranged in papillary pattern with nuclear overcrowding, optically clear nuclei and nuclear grooving.

**Follow up:** On six month follow up with USG neck normal thyroid echo texture was seen with no significant abnormality and no evidence of nodal metastasis.

**Discussion**

Thyroglossal duct cysts are epithelial lined cysts. They result from failure of normal developmental obliteration of the thyroglossal duct during 8th to 10th gestational weeks, and can thus occur anywhere along the course of the duct. Sixty percent of TGDC are located between hyoid bone and thyroid cartilage, 13% in substernal region, 24% above the hyoid bone including the submental site and 2% are intra-lingual [3]. The epithelial lining of the cyst varies with location. Those that form near the tongue are lined by stratified squamous epithelium and those away are lined by pseudo stratified ciliated columnar epithelium and variable amounts of thyroid tissue may remain in the cysts.

Thyroglossal duct cyst carcinoma usually presents with clinical features similar to benign thyroglossal duct cyst and it is often difficult to clinically diagnose it preoperatively. Preoperative evaluation is done with Ultrasonography, CT scan, Magnetic resonance imaging and FNAC in all cases. Even after these investigations a diagnosis of thyroglossal duct cyst carcinoma is usually made postoperatively after histopathological examination. This is due to inadequate sampling done with aspiration cytology studies without USG guidance. FNAC has sensitivity of 50–60% in diagnosing carcinomas of thyroglossal duct cyst [4].

Most common histological Pattern is Papillary Carcinoma of Thyroid (75%-80%), but other thyroid tumors such as Follicular, Hurthle cell, mixed Papillary-Follicular carcinomas may also occur. Squamous cell carcinoma arising within thyroglossal duct cyst is even rarer and contributes less than 5% of all malignant lesions involving thyroglossal cysts [5]. There are two theories to explain the thyrogenic origin of TGD adenocarcinomas. Firstly, the denovo theory which is based on the fact that in 62% of cases, ectopic thyroid tissue can be identified histopathologically, and this can be supported by the absence of occurrence of a medullary carcinoma in the TGD as it arises from parafollicular cells [6].

The second theory is the metastatic theory which suggests that thyroglossal cyst carcinoma is metastatic from an occult primary thyroid gland, as papillary carcinoma is multifocal in nature [7]. Although Thyroglossal duct itself could act as a natural conduit for the spread of thyroid carcinoma, the metastatic theory seems less likely. Squamous cell carcinoma is probably the only true carcinoma of the TGD, since the other malignancies actually develop in ectopic thyroid tissue. Thus, squamous cell carcinoma can be considered as the only primary thyroglossal cyst tumour, being rare and having a poor prognosis with a mortality rate of 30-40% [8].

Joseph and Komorowski proposed strict criteria to diagnose primary TDC carcinoma, these are the presence of a thyroglossal remnant, ectopic thyroid nests within the cyst wall and a clinically normal thyroid gland along with TGDC carcinoma [9].

The prognosis for Papillary carcinoma of TGDC is excellent, with occurrence of metastatic lesions in less than 2% of cases [4].
Conclusion

Because of rarity of TGDC, this diagnosis may be missed, drastically affecting the appropriateness of the treatment provided. The lesion could not be detected by FNAC as it was focal but was suspected for malignancy on CT. Regular follow-up to detect any recurrence in the thyroid gland is essential, particularly in cases where thyroid gland is not resected.

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