Cervical Thymic Cysts—Rare Abnormality: A Report of Two Cases

ABSTRACT
Cystic Lesions in the neck have a long differential diagnosis. Thymic cysts are considered rare in the cervical region, whereas they are more common in the superior mediastinal region. They may become more important diagnosis especially in the younger age group. In this article we will consider two cases, one male and other female with similar features and both being diagnosed as having cervical thymic cysts.

Keywords: Aberrant development of the thymus, Cervical thymic cysts, Cystic lesions in the neck, Ectopic thymic tissue.

CASE REPORTS

Case 1
Patient is 21 years male, complaining of a painless swelling in midline neck since birth. The swelling gradually and progressively increased. No complains of difficulty in swallowing, breathing or change of voice. On examination, single well defined nontender, noninflamed swelling, in suprasternal region, not moving on swallowing or deglutition, firm in consistency, no cough impulse or pulsations, immobile, nonreducible, noncompressible (Fig. 1).

INVESTIGATIONS

- Fine needle aspiration cytology: Benign cystic lesion
- Ultrasonography of neck: Cystic lesion in suprasternal region, 4 × 3 cm, both thyroid lobes normal.

Fig. 1: Male patient showing midline neck lesion
Cervical Thymic Cysts—Rare Abnormality: A Report of Two Cases

Computed tomography (CT) of neck: Well defined cystic peripheral enhancing lesion measuring 4.2 × 3.9 cm in midline, suprasternal region, lesion appears separate from surrounding structures, both lobes of thyroid gland and remaining structures in the neck are normal (Fig. 2).

Case 2

A female patient, 21 years old, complained of single painless swelling in midline neck since birth. Clinical examination and investigations were similar to the previous case, except the size, which was smaller (Fig. 3).

Computed tomography scan suggested single unilocular midline swelling slightly to right, measuring 3 × 3 cm with substernal extension, appears free from other structures (Fig. 4).

After obtaining informed consent from patient and preanaesthetic fitness, excision of the cyst was done under general anesthesia. Case 1 on operation showed midline swelling, in suprasternal region without any attachment to thyroid gland. Case 2 on operation showed swelling toward right, in suprasternal region, extending in the substernal region, with fibrous attachment to the thyroid gland. No major complications were encountered during and after the operation. Specimens were send for histopathological examination. Patients were discharged on 3rd postoperative day, followed-up on 7th day for removal of sutures. Repeat follow-up at 3 months showed no recurrence and adequate wound healing.

Histopathological report on both occasions showed squamous, nonkeratinized epithelium which was replaced by an inflammatory infiltrate containing numerous cholesterol clefts. The presence of thymic tissue in the walls of the cyst along with the pathognomonic presence of Hassall’s corpuscles identified the thymic origin of the cyst.

DISCUSSION

Thymus gland derives its origin from the third and, in some instances, the fourth pharyngeal pouch; it develops bilaterally early in fetal life and descends down the neck during the sixth to eighth week of gestational life, where the two primordia fuse to form the gland, which then reaches its definitive position in the mediastinum, behind the sternum. Clinically, in most cases, ectopic thymic tissue presents as a unilateral, asymptomatic neck mass, commonly in the left side of the neck. Thymus gland reaches its relative maximum size in children aged 2 to 4 years, attaining its final size at puberty when it weighs 30 to 40 gm. The gland then involutes and is replaced by fibro fatty tissue.

The pathogenesis of the ectopic thymus has not been fully clarified yet. Several theories have been proposed, including the following:

- Complete or partial failure of the unilateral gland to migrate to its normal position. The presence of parathyroid gland in the mass supports the idea of
nondescending tissue.\textsuperscript{11,12} Also, the fact that aberrant thymus is usually found in the normal pathway that the gland follows for its descent to the thorax is in favor of this theory.

- Sequestration of accessory cervical foci along the normal pathway of descent. In that case, there would be normal thymic tissue in the mediastinum, as most of it would have migrated.\textsuperscript{13}
- Ectopic thymic gland, in the form of masses located in the pharynx, the trachea or the base of the skull.\textsuperscript{14}
- Failure, after descent of the majority of the gland, ofrostal fragments to involute, leading to separate accessory lobes or cords.

There are two theories about the pathogenesis of a thymic cyst as follows:

- Relates it to acquired progressive cystic degeneration of Hassall’s corpuscles, of unknown etiology.
- Cystic changes in persistent unincorporated remnants of the thyropharyngeal duct.

According to Speer, it may be due to embryological remnants in the neck, sequestration products in pathological involution, neoplastic process, degeneration of Hassall’s bodies, mesenchymal elements.\textsuperscript{15}

These lesions usually lie medial to the sternocleidomastoid muscle, anterior to the carotid sheath and lateral to the thyroid gland. Preoperative CT and chest X-ray should be obtained to confirm the presence of normal or abnormal thymic tissue before surgery for suspected CTC. Intact complete surgical removal of the CTC is the treatment of choice, bearing in mind that it may be adherent to the surrounding structures, such as the vagus nerve, the internal jugular vein, and carotid artery as well as the phrenic, hypoglossal and recurrent laryngeal nerves. About 50\% of CTCs may have a fibrous connection to normal thymic tissue or active thymic cells, and removal of the duct may leave the patient athymic. This is not a problem in an adult, but a child may develop immunodeficiency problems.

REFERENCES