Scapular Bronchogenic Cyst: A Case Report and Literature Review
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Abstract
Bronchogenic cysts originate from abnormal budding of the tracheal diverticula during the embryological period. Inaccuracy in the process of growing of the ventral foregut will give rise to bronchogenic cyst. Scapular bronchogenic cyst is an extremely rare form of this anomaly. A three years old boy suffered for 2 years with left sided suprascapular cystic lesion which was gradually increasing in size. The swelling was 4 × 3 cm in size and non-tender. The cyst was evaluated by CT scan that showed complex cystic lesion over the left scapular spine. Total excision of the cyst was done and histopathology showed cutaneous bronchogenic cyst. The proposed mechanism for such cutaneous lesion is that the accessory buds from the tracheobronchial tree/primitive foregut migrated from the thorax in an aberrant manner to lie in periscapular positions. The definitive treatment of scapular lesions is total surgical excision. The final diagnosis is based on the histopathological findings in the majority of cases.

Keywords: Congenital; Scapular; Bronchogenic; Cutaneous.

Introduction
Bronchogenic cysts originate from abnormal budding of the tracheal diverticula during the embryological period. During the third week of embryonic development, the dorsal portion of the primitive foregut lengthens to form the oesophagus, and the ventral portion of the primitive gut undergoes differentiation to form the tracheobronchial tree. Insufficiency in the process of growing of the ventral foregut will give rise to bronchogenic cysts. The eventual location of the bronchogenic cysts depends on the stage of development in which the errors occur. Mediastinum is the most common extrapulmonary location. Other rare location like lingual, intra-abdominal and cutaneous regions have also been reported. Cutaneous bronchogenic cysts are the most infrequent lesions that are diagnosed by histopathological examination. However, the scapular bronchogenic cyst is an extremely rare form of this anomaly. Hereby, we describe a boy with a scapular bronchogenic cyst where the diagnosis was made by histopathological examination.

Case report
A three years old boy with no comorbidities was brought to the Pediatric Surgery Department at Sultan Qaboos University Hospital, Sultanate of Oman, with complaints of left sided suprascapular cystic lesion starting 2 years back. It was gradually increasing in size and not associated with pain, fever or weight loss. There was no other swelling elsewhere in the body and no history of trauma. On examination, there was a cystic swelling overlying the left scapular region measuring 4 × 3 cm in size, which was smooth walled, fluctuant, non-tender, and freely mobile. There was a scar overlying the swelling. Systemic examination was normal. He was evaluated by CT scan in another hospital which showed oval complex cystic lesion over the left scapular spine measuring 4 × 3 × 4 cm with calcification as well as the impression of the teratodermoid subcutaneous cyst. Total excision of the cyst was carried out. Pre-operatively, the cyst was adherent to the bone but without any communication with the joint cavity. The post-operative condition of the patient was uneventful. Macroscopically, the specimen consisted of a smooth walled unilocular cyst, 4 × 3.5 × 2 cm with a wall thickness of 0.2 - 0.4 mm and containing yellowish mucoid material. (Fig. 1)

Figure 1: Scapular bronchogenic cyst.
Histopathological examination showed a cyst wall predominantly lined by ciliated pseudostratified columnar epithelium with interspersed goblet cells and partly by stratified squamous epithelium. Smooth muscle cells, adipose tissue, nerve bundles as well as seromucinous glands were present in its wall. The appearances were consistent with the cutaneous bronchogenic cyst. (Figs. 2, 3)

Discussion

Bronchogenic cysts are very rare lesions; the reported incidence being 1 in 42000-68000. Abnormal separations of developing lung tissue result in the formation of these congenital cysts. Therefore, the thoracic cavity is the most common location for these cysts.

Table 1: Scapular bronchogenic cyst reported in the literature (1971-2010).

<table>
<thead>
<tr>
<th>Serial no</th>
<th>Sex</th>
<th>Age at presentation</th>
<th>Symptoms</th>
<th>Histopathology</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>male</td>
<td>4 months</td>
<td>mass</td>
<td>Cpte-smooth muscle</td>
<td>Pul et al.</td>
</tr>
<tr>
<td>2</td>
<td>male</td>
<td>10 years</td>
<td>Mass</td>
<td>Cpte-lymphoid aggregates</td>
<td>Jackson et al.</td>
</tr>
<tr>
<td>3</td>
<td>Male</td>
<td>8 months</td>
<td>Asymptomatic</td>
<td>NS</td>
<td>Fraga et al.</td>
</tr>
<tr>
<td>4</td>
<td>Male</td>
<td>30 months</td>
<td>Asymptomatic</td>
<td>NS</td>
<td>Fraga et al.</td>
</tr>
<tr>
<td>5</td>
<td>Male</td>
<td>01 year</td>
<td>Asymptomatic</td>
<td>NS</td>
<td>Fraga et al.</td>
</tr>
<tr>
<td>6</td>
<td>Male</td>
<td>04 years</td>
<td>Mass</td>
<td>Cpte-goblet cells+smooth muscle</td>
<td>Yu et al.</td>
</tr>
<tr>
<td>7</td>
<td>Male</td>
<td>46 years</td>
<td>Growing mass</td>
<td>Cpte+sebaceous glands+squamous +mucous gland</td>
<td>Tanita et al.</td>
</tr>
<tr>
<td>8</td>
<td>Male</td>
<td>newborn</td>
<td>Growing mass</td>
<td>Epithelium+ malignant melanoma Cpte</td>
<td>Tressner et al.</td>
</tr>
<tr>
<td>9</td>
<td>Male</td>
<td>01 year</td>
<td>Growing mass</td>
<td>Cpte-goblet cells+smooth muscle</td>
<td>Jona</td>
</tr>
<tr>
<td>10</td>
<td>Male</td>
<td>04 years</td>
<td>Mass</td>
<td>Stratified squamous epithelium Alternating with Cpte+goblet cells+sebaceous glands-smooth muscle</td>
<td>Van der Putte et al.</td>
</tr>
<tr>
<td>11</td>
<td>Female</td>
<td>08 years</td>
<td>Asymptomatic</td>
<td>Cpte-goblet cells+mucous glands</td>
<td>Manconi et al.</td>
</tr>
<tr>
<td>12</td>
<td>Female</td>
<td>01 year</td>
<td>Draining sinus</td>
<td>Cpte alternating with stratified</td>
<td>Ozel SK et al.</td>
</tr>
<tr>
<td>13</td>
<td>male</td>
<td>03 years</td>
<td>Mass</td>
<td>Squamous epithelium+mucous glands,Cpte+squamous epithelium</td>
<td>Kundal et al.</td>
</tr>
<tr>
<td>14</td>
<td>Male</td>
<td>03 years</td>
<td>Mass</td>
<td>Cpte+stratified squamous epithelium +goblet cells</td>
<td>Current case</td>
</tr>
</tbody>
</table>

cpte - ciliated pseudostratified columnar epithelium; NS - not specified.
Extra-thoracic cutaneous bronchogenic cysts are observed soon after birth or in early childhood as cystic masses. The usual sites of these lesions are suprasternal notch, pre sternal area, the neck and scapula. Scapular bronchogenic cysts are extremely rare among the extrathoracic variety. The division of developing lung bud, formation of the chest wall and scapula with its musculature are occurrences that imbricate during organogenesis. It is anticipated that accessory buds from the tracheobronchial tree/ primitive foregut migrated from the thorax in an aberrant manner to lie in periscapular positions. Additional proposed mechanisms for such devious cutaneous locations include in situ growth of a respiratory epithelium due to metaplasia of mature pre-existing cutaneous tissue and primary bizarre differentiation (heterotopia) in the developing skin.

Scapular cutaneous bronchogenic cyst presents as a progressively enlarging soft tissue swelling or a discharging sinus. Most of these are present shortly after birth or in early childhood and are more common among the male population. Along with the current case, 69 cutaneous bronchogenic cysts have been reported in the literature, with 14 (20.3%) located in the scapular region, as other locations of cutaneous bronchogenic cysts. Also, among these cases, 11 (78.6%) were diagnosed before the age of 4 years. In 1971, Fraga reported 30 cases of bronchogenic cysts, which is the largest population of cutaneous bronchogenic cysts, 12 out of the 30 patients (40%) were diagnosed below the age of 4 years. The majority of these cases presented with a swelling adjacent to the scapula that drained clear fluid. Remarkably, all the cysts were located subcutaneously. The reason for early detection may be the anatomical location itself as the scapular area is relatively exposed when compared with other parts of the body.

Proper imaging intervention (sinogram, chest X-ray, CT/MRI) will facilitate the evaluation of intrathoracic continuation as well as planning for the operative approach. The definitive treatment of scapular lesions is total surgical excision due to potential risk of infection; and more rarely, a malignant transformation of the epithelium in a scapular bronchogenic cyst. The pathological diagnosis is carried out based on the presentation of one or more tracheobronchial structures in the cyst wall. Frequently, hyaline cartilage, smooth muscle cells, elastic fibers, fibrous tissue, neural cells and seromucous glands can be seen.

Conclusion

This very rare congenital anomaly may present as common superficial skin lesion. Therefore, pediatric surgeons should be aware while dealing with superficial skin lesions in the scapular regions. The final diagnosis is based on the histopathological findings in the majority of cases.

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References