Subcutaneous bronchogenic cyst of the chest wall: A case report with brief literature review

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ABSTRACT

Introduction: Bronchogenic cysts are congenital lesions found in the mediastinum, particularly the posterior-superior area. The current study aims to report a rare case of a subcutaneous bronchogenic cyst in the chest wall.

Case report: A 41-year-old patient presented with a swelling of the chest wall. The mass had been present since birth. On examination, there was a large soft, round mass over the sternum subcutaneously. It was a fixed, non-flatulence, non-pulsatile, and non-tender mass.

Discussion: Usually, the condition develops between the fifth and sixteenth weeks of gestation, when the primordial intestine separates into two parts: dorsal, which gives rise to the esophagus, and ventral, which gives rise to the pulmonary bud and tracheobronchial tree. As a result, the cyst is an ectopic lung bud that may or may not be connected to the tracheobronchial tree but lacks mesenchymal tissue.

Conclusion: Although chest wall bronchogenic cysts are uncommon, they should be considered in the differential diagnosis of cystic and soft tissue lesions in adults with chest wall swelling.

1. Introduction

Bronchogenic cysts are congenital lesions caused by defective ventral foregut budding during tracheal and major bronchial structural development [1]. Bronchogenic cysts are rare, with a frequency of 1 in 42,680,000 people. When they occur, the majority are found in the mediastinum, particularly the posterosuperior area [2]. Intrapulmonary cysts account for approximately 15–20% of all bronchogenic cysts and are typically found in the lower lobes. They have been observed not just in infants and children but in adults as well later in life [3]. They can form in ectopic positions anywhere along the foregut developmental pathway [4]. On rare occasions, they have been observed in atypical sites such as the scapula, paravertebral, cervical, retroperitoneal, pericardial, omental, and perianal regions [5]. A chest wall bronchogenic cyst is an extremely rare condition, with just a few occurrences previously recorded in the literature [4–6].

The current study aims to describe a case of a subcutaneous bronchogenic cyst. The report is organized in line with SCARE 2020 criteria and contains a brief review of the literature [7].

2. Case report

2.1. Patient’s information

A 41-year-old male patient presented with a swelling of the chest wall. The mass had been present since birth, but it grew in size later in life till it burst 15 years ago. Three months after it burst, it began to gradually grow in size till the current presentation to the clinic. Dyspnea and chest discomfort are common symptoms. He has a history of asthma and underwent PCI in 2019. He is now taking aspirin, Plavix, and an inhalational short-acting beta-blocker as needed. He is not a smoker.

2.2. Clinical examination

On examination, there was a large (7 cm * 7 cm), soft, round mass over the sternum subcutaneously. It was a slightly mobile, non-
flutulence, non-pulsatile, and non-tender mass. There was no discol-
oration of the overlying skin, and the temperature was normal.

2.3. Diagnosis

The provisional diagnosis was lipoma. No investigation was per-
formed, the patient insisted on the removal of the mass.

2.4. Therapeutic intervention

Under local anesthesia, a transverse incision exposed a cystic lesion
containing milky fluid and adhering to the chest wall. Complete exci-
sion was performed, a corrugate drain was placed, and the wound was closed
in layers. Histopathological examination revealed fibrofatty tissue
transversed by cystic space and lined by cuboidal to pseudostratified
ciliated columnar epithelium (respiratory type) resting on a fibrous
stoma. The overall picture goes with a subcutaneous cyst of the chest
wall.

2.5. Follow up

The patient had a non-eventful postoperative period. After one week,
the stitches and drain were removed without complications.

3. Discussion

Congenital cystic lung lesions, which include congenital cystic ade-
nomatoid malformation, pulmonary sequestration, congenital lobar
emphysema (CLE), and bronchogenic cysts, are an uncommon but
clinically significant set of anomalies [8]. Bronchogenic cysts are non-
malignant congenital anomalies of the primordial ventral foregut. The
cyst wall contains structural characteristics of the airway such as carti-
lage, smooth muscle, mucous glands, and respiratory epithelium [9,10].
They are most commonly found intra-thoracic and are classified as
mediastinal or parenchymal, depending on where they are found. However,
they can also be found intra-abdominal or, in rare cases, cervical [11]. Up to 86% are mediastinal (middle and posterior medi-
astinum), and of these, some may be adjacent to the distal third of the
trachea or close to the main bronchus; thus, they can be subdivided into
pericarinal, paratracheal, para-esophageal, and retro-cardiac, with the
majority being in the right [12]. They’ve been found attached to the
sternum, pericardium, skin, and even the diaphragm [13].

The precise method through which bronchogenic cysts become
subcutaneous is unknown. Proposed mechanisms include the aberrant
tracheobronchial bud being separated from the respiratory tract by
growing skeletal structures and the bronchogenic cyst migrating into the
subcutaneous tissues [14]. They are typically unilocular and contain clear fluid or, less frequently, hemorrhagic secretions or air [15].

Bronchogenic cysts present clinically and radiologically in a variety
of ways, ranging from incidental radiologic findings without symptoms
to giant mass-like structures accompanied by severe symptoms [16,17].
The variability of presentation depends on their size, location, and
compression or invasion of adjacent tissues [5]. According to one study,
71% of children with mediastinal cysts were symptomatic [18]. A cough,
difficulty breathing or swallowing, hemoptysis, and infection are
symptoms of enlarged cysts. Symptoms of cutaneous bronchogenic cysts
include pain, a growing mass, a draining sinus, and, in rare cases,
cellulitis and abscess [5]. Lee et al. reported that more than half of their
cases were symptomatic, with cough being the most common symptom
[17]. Almost always, subcutaneous bronchogenic cysts are asymptom-
atric. The only symptoms reported in the cutaneous bronchogenic
are nasal puffiness and discharge [19]. Shilova et al. reported a case of a
pre-sternal bronchogenic cyst with tender, erythematous swelling [14].
Sirvani et al. reported a 6-year-old male child who presented with a pre-
stenal discharge sinus [20].

Although no modality has proven particularly specific for the
diagnosis of bronchogenic cysts, pre-operative imaging such as ultra-
sonography, CT, and MRI has been employed to better describe these
masses. Plain radiographs have minimal use. A unilocular fluid-filled
cystic tumor is often seen on ultrasonography [21]. CT and MRI pro-
vide excellent details about the size of the cyst and its connection to
adjacent tissues, which is very useful in pre-operative planning [22]. CT
usually shows an encapsulated mass without contrast enhancement and
various levels of fluid attenuation, but MRI shows enhancement on T2-
weighted images owing to mucinous and proteinaceous debris [23].
After a comprehensive clinical assessment, no imaging was deemed
necessary in the current case. A histopathological examination confirms
the definitive diagnosis, which shows a cyst with ciliated pseudostratified
columnar epithelium, indicating a respiratory origin. The his-
topathology distinguishes it from other types of cystic masses, such as
epidermoid cysts, teratomas, lymphangioimias, and dermoid cysts [24].
The same findings were observed in the current case.

The differential diagnosis for bronchogenic cysts is established by
their location in the body. A cystic neck mass might be an abscess, a
thyroglossal duct cyst, or a bronchogenic cyst [25]. A cystic chest wall
mass can be diagnosed as a dermoid cyst or teratoma, and a mediastinal
cyst can be TB, foregut cyst, pericardial cyst, congenital cystic adenoid
malformation, pulmonary sequestration, large B cell lymphoma, or
enterogenous cyst [6].

All bronchogenic cysts should be excised owing to the danger of the
consequences such as enlargement, compression of adjacent tissues,
infection, and cancer (bronchiolavolcar carcinoma, adenocarcinoma,
squamous cell carcinoma, and melanoma) [18,26]. Patients with bron-
chogenic cysts have a 0.7% lifetime risk of developing cancer [18]. This
supports the choice of early excision as soon as possible. The current
case underwent surgical excision as recommended.

In conclusion, although chest wall bronchogenic cysts are uncommon,
they should be considered in the differential diagnosis of cystic and
soft tissue lesions in adults with chest wall swelling. For a definitive
diagnosis and avoiding possible consequences, surgical resection is
suggested.

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None is found.

Ethical approval

Approval is not necessary for case report (till 3 cases in single report)
in our locality.

The family gave consent for the publication of the report.

Consent

Written informed consent was obtained from the patient's family for
publication of this case report and accompanying images. A copy of the
written consent is available for review by the Editor-in-Chief of this
journal on request.

Author contribution

Abdulwhahid M. Salh: major contribution of the idea, literature re-
view, final approval of the manuscript.

Sangar Abubakir A. Mirawdali: Surgeon performing the operation,
final approval of the manuscript.

Fahmi H. Kakamad, Marwan N. Hassan: Writing the manuscript,
literature review, final approval of the manuscript.

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