

Pleuropericardial Cyst

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Abstract

Introduction: Cystic lesions within the pericardial space are a rare entity and comprise 7% of the mediastinal masses and 33% of mediastinal cysts. The reported incidence of pleuropericardial cyst is 1 in 1,00,000 population and most cases are detected incidentally. Most cases are congenital and asymptomatic but life threatening complications may occur in the course of disease. *Case Report:* A 60 year old male pt. came to our hospital with chief complains of chronic cough, chest pain, dyspnoea, palpitation, a feeling of retrosternal pressure and right shoulder discomfort radiating to right arm since 2 months. Chest X-ray showed e/o isolated cystic shadow in right mediastinum. Patient was advised surgery and underwent Anterior Thoracotomy and excision of cyst. *Discussion:* Patients with pleuropericardial cysts are usually asymptomatic (up to 60-75% cases). Symptoms usually appear when the cyst compresses on a nearby structure. Common symptoms include chronic cough, chest pain, dyspnoea and a feeling of retrosternal pressure. *Conclusion:* Pleuropericardial cyst occurs rarely, is clinically silent in majority of cases but can cause life threatening complications. Therefore it is prudent to keep such anomalies in mind when evaluating patients from third and fourth decade of life either during routine health check-up or when such patients present with chest complaints.

Keywords: Pleuropericardial cyst; mediastinum; chronic cough; chest pain; dyspnoea; retrosternal pressure.

Introduction

- Cystic lesions within the pericardial space are a rare entity and comprise 7% of the mediastinal masses and 33% of mediastinal cysts [1,2].
- The reported incidence of pleuropericardial cyst is 1 in 1,00,000 population and most cases are detected incidentally [3-6].
- They are usually found in third or fourth decade of life and male and the females are affected equally [5].
- Most cases are congenital and asymptomatic but life threatening complications may occur in the course of disease. They are found in the literature under various names: le kyste pleuropericardique (Jeaubert de Beaujeu, 1945; Roche, 1954), pleural cyst, pericardial cyst, pericardialcoelomic cyst (Lambert, 1940), springwater cyst (Greenfield, Steinberg, and Touroff, 1943), mesothelial cyst (Churchill and Mallory, 1937), and thin-walled cyst [7].

Case Report

- A 60 year old male pt. came to our hospital with chief complains of chronic cough, chest pain, dyspnoea, palpitation, a feeling of retrosternal pressure and right shoulder discomfort radiating to right arm since 2 months.
- There was no past history of diabetes mellitus, hypertension, bronchial asthma, tuberculosis or Ischaemic heart disease.

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- On examination, vitals were normal.
- All routine blood investigations were within normal limits.
- Chest X-ray showed e/o isolated cystic shadow in right mediastinum.
- HRCT Thorax (Plain) revealed a well-circumscribed thin walled cystic lesion in right anterior mediastinum extending along the right cardiac border and IVC upto the level of aortic arch measuring approximately 11.5×7.7×9.4 cms in craniocaudal (CC), anteroposterior (AP) and Transverse (Tr) dimensions respectively. It extended posteriorly upto right interlobar artery division. Anteriorly the mass was limited by the right anterior chest wall extending from the 1st anterior rib upto 6th anterior rib. Inferiorly it was limited by right hemidiaphragm.
- Impression- Likely to be 1) Pleuropericardial cyst or 2) Lymphangioma.
- Patient was advised surgery and underwent Anterior Thoracotomy and excision of cyst. 5cm wide incision was taken over Right inframammary line. Flaps were raised. Pectoralis major and pectoralis minor were retracted. Space was created through 4th Intercostal space (ICS). Large cystic cavity involving pleura and pericardium was seen. Fluid was drained and whole cyst was excised. Closure was done in layers after putting Intercostal drain (ICD). The procedure was uneventful.
- Cyst wall specimen was sent for histopathological examination for confirmation. Section showed fibrofatty tissue with thin fibro-collagen cyst wall lined focally by flattened to cuboidal epithelium. Large congested vessel also seen in adjacent fibrofatty tissue.
- Impression - Simple cyst consistent with Pleuropericardial cyst.



Fig. 1: 5 cm wide incision taken over Right inframammary line

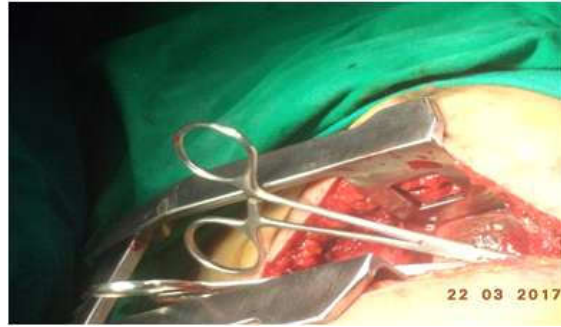


Fig. 2: Large Pleuropericardial Cyst involving pleura and pericardium



Fig. 3: Fluid drained and whole cyst was excised



Fig. 4: Closure done in layers



Fig. 5: Pre-op Chest X-ray showing isolated cystic shadow in Right mediastinum



Fig. 6: HRCT Thorax(Plain) showing a well circumscribed thin walled cystic lesion in Right anterior mediastinum



Fig. 7: Post-op Chest X-ray

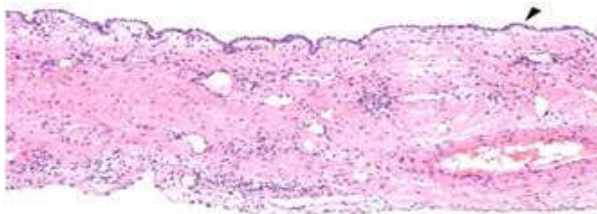


Fig. 8: HPE section showing fibrofatty tissue with thin fibrocollagen cyst wall lined focally by flattened to cuboidal epithelium

Discussion

- Patients with pleuropericardial cysts are usually asymptomatic (up to 60-75% cases). Symptoms usually appear when the cyst compresses on a nearby structure. Common symptoms include chronic cough, chest pain, dyspnoea and a feeling of retrosternal pressure.
- Pleuropericardial cysts are usually benign in nature but complications may arise eventually in the form of compression, inflammation, haemorrhage or rupture.
- Pleuropericardial cyst may appear as an incidental finding in chest X-ray in an asymptomatic patient.
- CT scan is the diagnostic modality of choice in all cases and diffusion weighted cardiac MRI for cases with diagnostic confusion. Echocardiography provides a narrow window and lesions at unusual locations may be missed with this technique.
- Management protocol is similar to that of mediastinal mass. Large cysts should be resected surgically.
- An algorithmic approach should be followed for management depending on size, shape and compressibility of the mass, patient symptoms, surgical fitness and patient preference.

Conclusion

- Pleuropericardial cyst occurs rarely, is clinically silent in majority of cases but can cause life threatening complications.
- Therefore it is prudent to keep such anomalies in mind when evaluating patients from third and fourth decade of life either during routine health check-up or when such patients present with chest complaints.
- A high index of suspicion and a subsequent thorough work-up is necessary to reach a diagnosis.

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Ethical Approval: Not required.

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