# Giant Cystic Chondroid Hamartoma

# Özgür Katrancıoğlu<sup>1</sup>, Yücel Akkaş<sup>1</sup>, Hakan İmamoğlu<sup>2</sup>, Sema Polat<sup>3</sup>

<sup>1</sup>Sivas Numune Hastanesi, Göğüs Cerrahisi Kliniği, Sivas

# **Abstract**

Pulmonary hamartoma is the most common benign neoplasm of the lung. Its size is usually smaller than 4 cm, and a pulmonary hamartoma larger than 10 cm is rare. Pulmonary hamartoma with air-filled cystic areas is also a rare form. Since it is a rare condition, in this article, we reported a case of giant pulmonary cystic hamartoma in a 49-year-old patient admitted with cough and breathing difficulty.

Keywords: Giant, cystic, chondroid, pulmonary hamartoma

### Özet

Pulmoner hamartom, akciğerin en sık görülen iyi huylu tümörüdür. Boyutu genellikle 4 cm'den küçüktür ve 10 cm'den büyük pulmoner hamartom nadirdir. Hava dolu kistik alanlar içeren pulmoner hamartom nadir görülen bir formdur. Bu yazıda; nadir görülmesi nedeniyle, öksürük ve nefes darlığı şikayeti ile başvuran 49 yaşındaki dev pulmoner kistik hamartom olgusu sunulmuştur.

Anahtar Kelimeler: Dev, kistik, kondroid, pulmoner hamartom

#### INTRODUCTION

A hamartoma is a benign neoplasm that consists of an abnormal quantity, mixture, and arrangement of the tissues that are normally found in the organ. Pulmonary hamartoma (PH) is the most common neoplasm in the lung; however, its cystic form is rare (1). Tumor size is usually smaller than 4 cm. Pulmonary hamartomas range in size from 1 cm to 8 cm in diameter in different series (2). A pulmonary hamartoma of larger than 10 cm in size is very rare (3). We report a case of giant cystic PH, which is a rare type of PH.

#### **CASE PRESENTATION**

A 49-year-old female was admitted to our hospital with cough and dyspnea for 1 year. On physical examination, breath sounds were absent in the right lower lung field. Chest radiography demonstrated a huge cystic mass in the right hemithorax and computed tomography of the chest revealed a large cystic and solid mass in the right lower lobe of the lung (Figure 1). The tumor was removed by wedge resection through a right lateral thoracotomy under the impression of ruptured hydatic cyst.

On gross examination, a well-circumscribed, lobulated, large, solid, and cystic mass was found. Its size was 14x10.5x8.5 cm (Figure 2). Microscopically, the solid component was composed of cartilaginous and adipose tissue. The cystic and cleft-like areas were lined by ciliated columnar epithelium, and there were also papillary projections through the cleft-like areas (Figure 3, 4). The patient was diagnosed as giant cystic chondroid PH. No signs of recurrence were detected for 1 year after the operation.

#### DISCUSSION

Pulmonary hamartoma is the most common benign tumor of the lung, with an incidence of 0.025%-0.32% according to varied series (4). Giant cystic chondroid hamartoma occurs generally in adults and rarely in children (5). Pneumothorax and malignant degeneration can occur with the tumor as a complication (6, 7). In this case, no complications were observed. Congenital abnormalities and benign and malignant tumors can be associated with PH (8). No associations were found in this case. Most PHs are parenchymal (peripheral) tumors, and only 10% of PHs are endobronchial tumors (4). Most PHs have a large

Received Date / Alındığı Tarih: 13.02.2012 Accepted Date / Kabul Tarihi: 07.07.2012

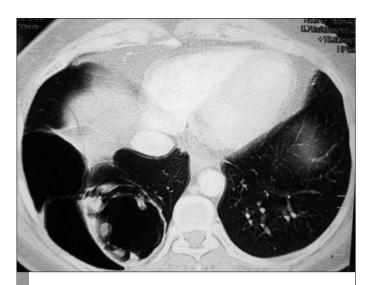
Address for correspondence / Yazışma Adresi Yücel Akkaş, Sivas Numune Hastanesi, Göğüs Cerrahisi Kliniği, Sivas, Türkiye E-mail / E-posta: y.akkas@yahoo.com

© Copyright 2014 Turkish Respiratory Society (TRS) Eurasian J Pulmonol 2014 DOI: 10.5152/ejp.2014.53765

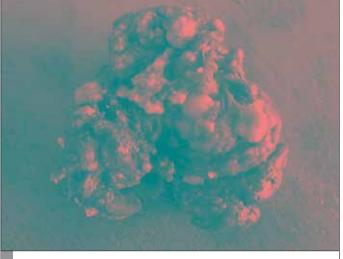
• Available online at www.eurasianjpulmonol.com

<sup>&</sup>lt;sup>2</sup>Sivas Numune Hastanesi, Radyoloji Kliniği, Sivas

<sup>&</sup>lt;sup>3</sup>Sivas Numune Hastanesi, Patoloji Kliniği, Sivas



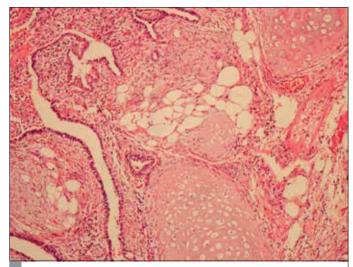
**Figure 1.** A large cystic and solid mass in the right lower lobe of the lung with the chest computed tomography



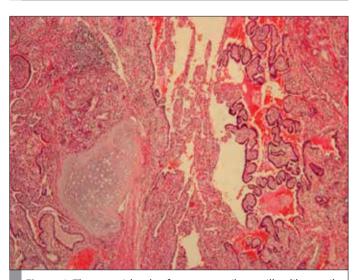
**Figure 2.** A well-circumscribed, lobulated, large solid, and cystic mass is shown.

cartilaginous component and are called chondromatous hamartoma. Other subtypes are leiomyomatous, lymphangiomyomatous, adenofibromatous, and fibroleiomyomatous hamartomas. Their size may begin to increase slowly at a rate of 3.2±2.6 mm per year (9). Pulmonary hamartomas bigger than 10 cm and cystic type PHs are extremely rare (3). Cysts are air-filled cystic areas, and why they occur in hamartomas is unknown. There are several hypotheses to explain this development. One of them is a check-valve mechanism owing to partial obstruction of the bronchioles by the mass. The route of air entry into the tumor might result in gradual expansion of small epithelial-lined tubules resembling bronchioles (10). But, if this hypothesis is true, there must be an airway connection to cystic areas. In our study, this connection was not detected. The other hypothesis is that cleft-like spaces may become cysts during the growth of the tumor (1).

Giant cystic chondroid PH is an unusual presentation of PH. This form is rare but may put in the differential diagnosis of large cystic pulmonary tumors and tumor-like conditions.



**Figure 3.** The cysts and cleft-like spaces are lined by ciliated columnar epithelium, mature adipose tissue, and mature cartilage (HEX100).



**Figure 4.** There are islands of mature cartilage, villus-like papillary projections, and cystic space (HEx40).

Etik Komite Onayı: N/A.

Hasta Onamı: Yazılı hasta onamı bu olguya katılan hastadan alınmıştır.

Hakem Değerlendirmesi: Dış bağımsız.

Yazar Katkıları: Fikir - Y.A.; Tasarım - Ö.K.; Denetleme - Y.A.; Kaynaklar - H.İ.; Malzemeler - H.İ.; Veri toplanması ve/veya işlemesi - Y.A.; Analiz ve/veya yorum - H.İ.; Literatür taraması - Y.A.; Yazıyı yazan - Y.A., S.P.; Eleştirel İnceleme - H.İ.

Çıkar Çatışması: Yazarlar çıkar çatışması bildirmemişlerdir.

**Finansal Destek:** Yazarlar bu çalışma için finansal destek almadıklarını beyan etmişlerdir.

## **REFERENCES**

- Miura K, Hori T, Yoshizawa K, Hamaguchi N, Morita J. Cystic pulmonary hamartoma. Ann Thorac Surg 1990; 49: 828-9.
- Salminen US. Pulmonary hamartoma. A clinical study of 77 cases in a 21year period and review of literature. Eur J Cardiothorac Surg 1990; 4: 15-8.
- Lee SY, Park HJ, Lee CS, Lee KR. Giant pulmonary hamartoma. Eur J Cardiothorac Surg 2002; 22: 1006. [CrossRef]
- 4. Cosío BG, Villena V, Echave-Sustaeta J, de Miguel E, Alfaro J, Hernandez L, et al. Endobronchial hamartoma. Chest 2002; 122: 202-5. [CrossRef]

- Ozbudak IH, Dertsiz L, Bassorgun C, Ozbilim G. Giant cystic chondroid hamartoma of the lung. J Ped Surg 2008; 43: 1909-11. [CrossRef]
- Stella F, Davoli F, Brandolini J, Dolci G, Sellitri F, Fiorentino M, et al. A rare case of giant cystic chondroid hamartoma of the lung presenting with left side pneumothorax. Minerva Chir 2009; 64: 117-9.
- 7. Basile A, Gregoris A, Antoci B, Romanelli M. Malignant change in a benign pulmonary hamartoma. Thorax 1989; 44: 232-3.
- Gabrail NY, Zara BY. Pulmonary hamartoma syndrome. Chest 1990; 97: 962-5. [CrossRef]
- 9. Hansen CP, Holtveg H, Francis D, Rasch L, Bertelsen S. Pulmonary hamartoma. J Thorac Cardiovasc Surg 1992; 104: 674-8. [CrossRef]
- Doppman J, Wilson G. Cystic pulmonary hamartoma. Br J Radiol. 1965;
  38: 629-31. [CrossRef]