Pulmonary sequestration mimicking lung cancer – A case report

Plučni sekvestar koji imitira karcinom pluća

Tatjana N. Adžić Vukičević*,†, Dragan V. Radovanovič†‡, Bojana D. Aćimović†, Marko P. Popović‡

*Faculty of Medicine, University of Belgrade, Belgrade, Serbia; †Clinic for Pulmonology; ‡Clinic for Thoracic Surgery, Clinical Center of Serbia, Belgrade, Serbia

Abstract

Introduction. Pulmonary sequestration is a rare congenital anomaly and most intralobar sequestrations were located in lower lobes. Case report. We reported an unusual 28-year-old female patient with intralobar pulmonary sequestration on the left lower lobe, successfully treated with lobectomy. Computed tomography (CT) of the chest with intravenous contrast revealed multiple clustered cystic lesions in the left lower lobe with aberrant artery from descending aorta. Additional aortography showed an aberrant artery (3 mm in diameter) arising from the abdominal aorta and flowing into the lesion. Conclusion. Standard therapy regimen for pulmonary sequestration includes surgery. CT scan of thorax with intravenous contrast and aortography represent the gold standard for its diagnosis. Tumor-like shadows seen on the chest radiography or CT scans should not be always suspected on malignant lesions.

Key words: bronchopulmonary sequestration; diagnostic techniques and procedures; diagnosis, differential; thoracic surgical procedures.

Correspondence to: Tatjana Adžić Vukičević, Clinic for Pulmonology, Clinical Center of Serbia, Višegradska 26, 11 000 Belgrade, Serbia. Phone: +381 11 366 33 07, E-mail: adžić_tatjana@yahoo.com

Pulmonary sequestration is a rare congenital malformation of lung characterized by abnormal pulmonary parenchyma without tracheobronchial airway connection and with blood supply from a systemic artery. Definitive treatment includes surgical excision, usually lobectomy, followed by division of the anomalous artery via standard thoracotomy.

Case report

A 28-year-old female was hospitalized due to chest pain and high temperature. Chest radiography showed a tumor-like shadow in the left lower lobe (Figure 1). Laboratory test findings showed elevated sedimentation rate – 94 mm/h, C-reactive protein (CRP) – 198 mg/mL, blood cell count – 17.3 × 10³/mm³ and D-dimer – 4.39 mg/L fibrinogen equivalent units (FEU). Electrocardiogram (ECG), heart ultrasound, lung functions tests and blood gas analysis were normal. A bronchoscopy finding was normal. The computed tomography (CT) scan of the chest with intravenous contrast revealed multiple clustered cystic lesions in the left lower lobe with the aberrant artery from the descending aorta (Figure 2). Aortography showed the aberrant artery (3 mm in diameter) arising from the abdominal aorta and flowing into the lesion (Figure 3). Left lower lobectomy was done. The intraoperative finding showed the aberrant artery coming...
Fig. 1 – Chest radiography: tumor-like shadow in the left lower lobe.

Fig. 2 – a) Thorax computed tomography (CT) scan: multicystic mass in the left lower lobe; b) Thorax CT scan: aberrant artery arises from the descendenting thoracic aorta.

Fig. 3 – Aortography showing the aberrant artery.

Discussion

Pulmonary sequestration is a rare congenital lesion of the lung parenchyma of unknown etiology without normal connection with the tracheobronchial tree and with the blood supply directly from the descending aorta. It is a rare developmental abnormality which account for 0.15–6.4% of all congenital lung anomalies. There are two morphological types: extralobar (25% of all cases), which have their own pleura, and intralobar (75% of all cases), which are surrounded by visceral pleura. Patients were presented in younger age with symptoms of recurrent bronchopulmonary infections and if they are recognized they should be operated on. If

the symptoms of patients with pulmonary sequestration are not serious, intervention should be prolonged for several years. In the cases of repeated episodes of pneumonia, after physical investigation, diagnostic methods should include usual laboratory analysis, chest radiography and thorax CT scan with intravenous contrast. In our patient thorax CT scan with intravenous contrast revealed the aberrant artery from the descending aorta. Examination was completed with digital subtraction arteriography. Pulmonary sequestration is often misdiagnosed as congenital pulmonary cysts of bronchiectasis complicated with infection, benign lung tumor, diaphragmatic hernia and lung cancer. Surgical resection is a definitive treatment in patients with pulmonary sequestration. Standard posterolateral thoracotomy has been used for a years. Because of its benign etiology partial lung resection for pulmonary sequestration should be more appropriate than lobectomy. Video assisted thoracoscopic surgery (VATS) is a better alternative to standard thoracotomy for pulmonary sequestration because of minimal surgical trauma, postoperative pain and duration of hospitalization. In the presented patient VATS was not done because of deficient technical equipment. Potential barriers to VATS include intrapleural adhesion due to the inflammatory process and dissection and dividing of the feeding artery, but rarely reported in case series. There are a few reports on successful management of asymptomatic pulmonary sequestration with angiographic embolization.

**Conclusion**

In a young patient tumor-like shadow on chest radiography should be suspected on a benign lesion. Detailed radiological investigations, including thorax CT scan, followed with arteriography, are mandatory.
REFERENCES


