A 15-year-old medium-height male patient was admitted to the thoracic surgery department with a right-sided pneumothorax with persistent air leak. He was a nonsmoker and had not been exposed to pneumotoxic factors. One month earlier, he developed right-sided chest pain, and pneumothorax was diagnosed in a district hospital. Positional therapy was started, but subsequently drainage of the right pleural cavity was implemented, initially passively and then actively. This treatment was ineffective, and the patient was referred to our hospital.

On admission, he was in good condition. Laboratory blood and urine tests were normal. Chest x-ray showed right-sided pneumothorax with a partially collapsed right lung. He underwent video-assisted thoracoscopy, and a small bulla in the right upper lobe was resected. The lung re-expanded, and the chest tube was removed 2 days later. Histological examination of the lung sample revealed an infiltration consisting of large cells with folded nuclei, forming granulomas (FIGURE 1A). These cells were positive for langerin (FIGURE 1B) and CD1a antigen (FIGURE 1C), but negative for BRAFV600E, which led us to establish a diagnosis of Langerhans cell histiocytosis (LCH). High-resolution computed tomography (HRCT) showed a few small nodules at the base of the right upper lobe (FIGURE 1D). Pulmonary function tests and 6-minute walk test were normal (forced expiratory volume in 1 second, 89% pred.; forced vital capacity, 108% pred.; total lung capacity, 88% pred.), while the transfer factor for carbon monoxide was reduced (69% pred.). Computed tomography did not show any osteolytic lesions, and there were no changes on abdominal
ultrasound or brain magnetic resonance imaging. HRCT performed a year later revealed complete regression of the nodules. No progression has been observed during the 5-year follow-up.

Spontaneous pneumothorax is caused by intrinsic factors. Primary spontaneous pneumothorax occurs in patients without a history of pulmonary disease, while secondary spontaneous pneumothorax is observed in patients with underlying diseases. The rupture of a small subpleural cyst is the most common cause of primary spontaneous pneumothorax. The incidence of this condition is 18 to 24/100 000 in men and 6 to 9.8/100 000 in women. The peak incidence is at 15 to 34 years of age. Risk factors include smoking, tall stature, and low body mass. Diseases that can cause secondary pneumothorax include chronic obstructive pulmonary disease, emphysema, lung abscess, tuberculosis, pneumonia, lymphangioleiomyomatosis, lymphoid interstitial pneumonia, Birt–Hogg–Dubé syndrome, alpha-1 antitrypsin deficiency, or pulmonary LCH (PLCH).1-5

PLCH is a rare disease, which usually affects young people and smokers. LCH may be limited to the lungs, or the lungs may be one of the involved organs, both at the time of diagnosis and during follow-up. Pneumothorax is the first sign of PLCH in about 30% of cases. HRCT is important in the diagnosis of PLCH. The radiographic changes range from small intralobular nodules to cystic lesions with sparing of the costophrenic angles.2-4 To our knowledge, this is the first case of PLCH in a nonsmoker with such minimal lesions. The absence of obvious radiographic evidence of lung disease does not exclude the need for lung biopsy in patients undergoing surgical treatment of pneumothorax. It is essential to implement measures aimed at inhibiting the progression of the disease, especially smoking cessation.

**REFERENCES**


**FIGURE 1** Langerhans cell histiocytosis on high-resolution computed tomography scan: D – small nonspecific nodules at the base of the upper lobe of the right lung (arrow)