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Surgical correction of postpneumonectomy-like syndrome in a patient with a tuberculosis-destroyed lung

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Postpneumonectomy syndrome is a rare condition that is characterized by dyspnea resulting from an extreme mediastinal shift and bronchial compression of the residual lung after right or left pneumonectomy.¹⁻³ Severe fibrosis of the lung such as is seen in a lung destroyed by tuberculosis (TB) can cause similar clinical features in the absence of pneumonectomy.⁴

We report here a couple of unique cases of postpneumonectomy syndrome without pneumonectomy in which treatment via pneumonectomy and mediastinal repositioning with tissue expanders was successful.

Clinical Summaries

PATIENT 1. A 47-year-old woman had had severe progressive dyspnea for 6 months. She had had pulmonary TB 20 years earlier, which was completely cured by administration of anti-TB drugs. On physical examination, breath sounds were decreased in the left lung field. A simple chest radiograph and computed tomogram (Figure 1, A) revealed that the left lung was almost completely destroyed and the right main bronchus was compressed by the right main pulmonary artery and the vertebral body.

The operation was performed via a left posterolateral thoracotomy. The severe pleural adhesion was dissected, and then left pneumonectomy was performed followed by insertion of two types of tissue expanders into the pleural cavity. One expander was ellipsoid (350 mL) and the other was crescent shaped (250 mL). After insertion of the tissue expanders, normal saline that included antibiotics

was injected into the tissue expanders, and the thoracotomy wound was closed.

Postoperatively, she recovered without any problem. A plain chest radiograph showed that the mediastinum had returned to a normal position (Figure 1, B). In addition, the expiratory wheezing and dyspnea disappeared.

The patient was discharged from the hospital on the 13th postoperative day. At present, 4 years after the operation, she is being observed at the outpatient department and is without dyspnea.

PATIENT 2. A 50-year-old woman had had severe progressive exertional dyspnea for the previous 6 months. She had had pulmonary TB 30 years earlier, which was completely cured by anti-TB medication. Six months before her current admission, the right lung totally collapsed and she had paroxysmal severe exertional dyspnea, which led her to visit our hospital's emergency room. On physical examination, there was no audible breath sound in the right side of the chest. Chest computed tomography revealed a totally collapsed right lung, extreme deviation of the mediastinal structure, and a narrowed left main bronchus, which was compressed between the left main pulmonary artery and the descending aorta (Figure 2, A). We performed a right pneumonectomy via a posterolateral thoracotomy. There was no pleural adhesion. Two types of tissue expanders were inserted, with 320 mL of normal saline being infused into the ellipsoid one and 450 mL into the crescent-shaped one.

Postoperatively, the mediastinum returned to the normal position (Figure 2, B). The patient was discharged on the seventh postoperative day. Five months after the operation, she remains free of expiratory stridor and dyspnea on exertion.

Discussion

Postpneumonectomy syndrome is a rare complication that is characterized by tracheal stenosis and severe dyspnea; these symptoms are caused by mediastinal shifting after pneumonectomy. In the cases we presented, the right or left main bronchus was compressed between the main pulmonary artery and vertebra/aorta not by surgical pneumonectomy, but by auto-pneumonectomy that was due to TB. These 2 cases show that surgical pneumonectomy is not a prerequisite for postpneumonectomy syndrome because this malady occurs as a result of stenosis of the bronchus after vigorous movement and rotation of the mediastinum.

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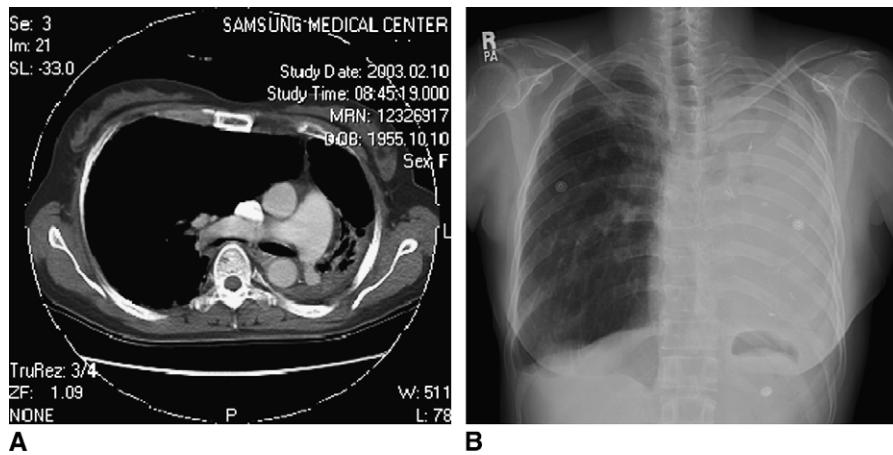


Figure 1. A, Preoperative chest computed tomogram shows narrowing of the right main bronchus compressed between the right main pulmonary artery and thoracic vertebral body. B, Postoperative chest radiograph shows the mediastinum in its normal position.

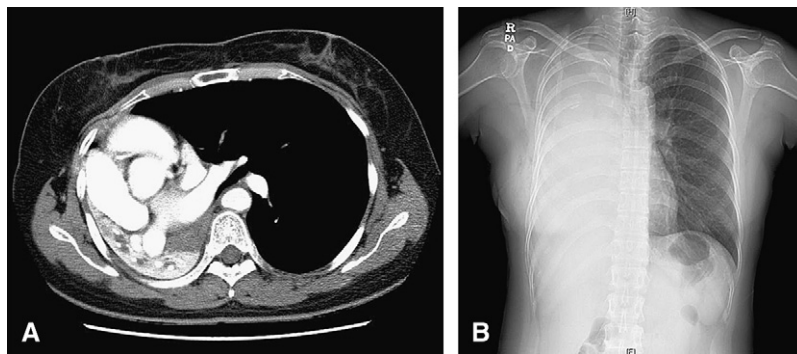


Figure 2. A, Preoperative chest computed tomogram shows narrowing of the left main bronchus compressed between the left main pulmonary artery and descending aorta. B, Postoperative chest radiograph shows the mediastinum in its normal position.

The diagnostic approach for our cases was the same as that for postpneumonectomy syndrome. On a plain chest radiograph, the mediastinum and bronchi are shifted posterolaterally and the opposite lung is herniated toward the lesion. Computed tomography of the chest can reveal that the heart and the great vessels are rotated toward the lesion and the bronchi are compressed by the vertebra and the great vessels.

Many methods have been tried to treat this syndrome. The most favorable method is insertion of tissue expanders in the hemithorax to shift the mediastinum to the midline and alleviate the obstruction of the trachea. Pneumonectomy must precede repositioning of the mediastinum by insertion of one or more tissue expanders in the case of a TB-destroyed lung without surgical pneumonectomy.

In conclusion, postpneumonectomy syndrome must be considered in the differential diagnosis when patients exhibiting dyspnea have a TB-destroyed lung or a reduced lung volume that is caused by severe pulmonary fibrosis. Once this syndrome is diagnosed, me-

diastinal repositioning by insertion of one or more tissue expanders can be expected to lead to successful treatment.

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