



Left lower lobe sleeve resection for endobronchial schwannoma

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Abstract: Schwannomas are mesenchymal neoplasms originating from Schwann cells of the nerve sheath. The tumor may appear anywhere in the body, while it has a predilection for extremities, trunk, retroperitoneum, head, and neck. Primary pulmonary schwannoma is extremely rare, accounting for less than 0.2% of all lung tumors. A 42-year-old man was incidentally discovered to have an endobronchial tumor, and we performed the left lower lobe sleeve resection to remove the mass completely. Postoperative histopathological findings confirmed a benign schwannoma. Although endobronchial schwannoma is extremely rare, follow-ups are crucial for monitoring tumor growth and reducing the risk of recurrence.

Keywords: Endobronchial schwannoma; sleeve lobectomy; follow-up

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Introduction

Schwannoma is a mesenchymal neoplasm originating from Schwann cells of the nerve sheath. Primary pulmonary schwannoma is an extremely rare tumor, which accounts for less than 0.2% of all lung tumors. The diagnosis of endobronchial schwannoma is difficult due to the fact that most patients are asymptomatic. The main differential diagnoses include neurofibroma, neurilemmoma and neurogenic sarcoma. Surgery is the preferred treatment of schwannoma. We present a rare case of a schwannoma arising from left main bronchus.

Case presentation

A 42-year-old female presented to our hospital with a 1-month history of hemoptysis. She denied family history and past medical history. Physical examination and routine laboratory analyses were unremarkable. A computed tomography (CT) of chest with contrast revealed a 2.4 cm × 4.3 cm, soft-tissue tumor mass with irregular enhancement located in the left hilar and paraspinal sulcus. The left lower lobe bronchus was embedded in the mass (*Figure 1*). Subsequently, a bronchoscopy was performed

and showed that neoformation protruded into the lower part of left main bronchus and the left lower lobe bronchus was almost completely blocked (*Figure 2A*). With the results presented above, we hypothesized that the tumor may be carcinoid tumour. The endobronchial biopsy showed that a submucosal spindle cell proliferation and a chronic mucosal inflammation accompanied with squamous metaplasia in the bronchial epithelium. It was suspected to be peripheral nerve tumors. However, it does not rule out the possibility of malignant tumors. The patient underwent the left lower lobectomy and end-to-end anastomosis between the left main bronchus and the upper left bronchus after systemic evaluation (*Figure 2B*). Benign nature of the endobronchial tumor was verified via fast frozen section examination. The postoperative pathology study confirmed endobronchial schwannoma (*Figure 3A*), and immunohistochemistry with S-100 protein was positive (*Figure 3B*), but negative for SMA, TTF1, CK7, MART-1 and desmin. The patient underwent an uneventful post-operative course without complications. On the seventh day after surgery, bronchoscopy showed a good end-to-end anastomosis. At one-year follow-up, there was no evidence of tumor recurrence.

Discussion

Schwannomas are mesenchymal neoplasms originating from Schwann cells of the nerve sheath. They may appear anywhere on the body, while they have a predilection for the extremities, trunk, retroperitoneum, head and neck (1). Primary pulmonary schwannoma is an extremely rare tumor, accounting for less than 0.2% of all lung tumors (2). Endobronchial schwannoma was first described by Feldhaus in 1989 (3), and there have been no more than 30 cases reported in the English literature up to now.

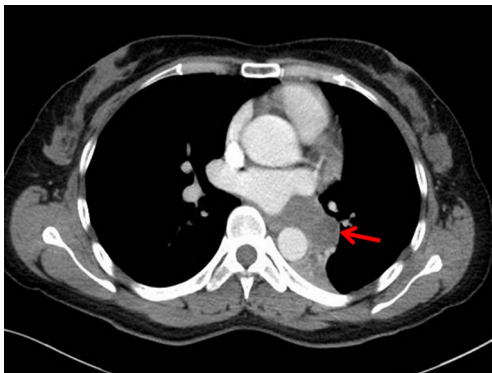


Figure 1 Computed tomography (CT) of chest with contrast revealed a left pulmonary hilar mass (red arrow) with lower lobe atelectasis.

Some endobronchial schwannomas are asymptomatic and discovered incidentally on radiographs. Common signs and symptoms include fever, cough, hemoptysis, dyspnea, and post-obstructive pneumonia in symptomatic patients. It is difficult to diagnose schwannoma based on clinical manifestations. Bronchoscopy is often used to further confirm the location, size, shape, and obstruction of the lesion in the bronchus. The differential diagnosis for endobronchial schwannomas includes neurofibroma, neurilemmoma and neurogenic sarcoma. Pathological biopsy and immunohistochemistry are the gold standard for diagnosis of this tumor.

Treatment options of endobronchial schwannomas are usually determined by the size and location of the tumor. In most cases, surgery is the preferred treatment option (4). Among that, lobectomy is considered the standard treatment for some central pulmonary schwannomas. In our case, although the tumors located in a troublesome position of the lower part of left main bronchus and left lower lobe bronchus, sleeve lobectomy of the left lower lobe not only completely remove the tumor, but also retain the patient's upper lobe. Recently, bronchoscopic treatment has been proved to be a safe and efficient tool for benign tracheobronchial tumors (5). However, there are few reports on bronchoscopic removal of endobronchial schwannoma (6,7).

As schwannomas are benign tumors, there are rarely

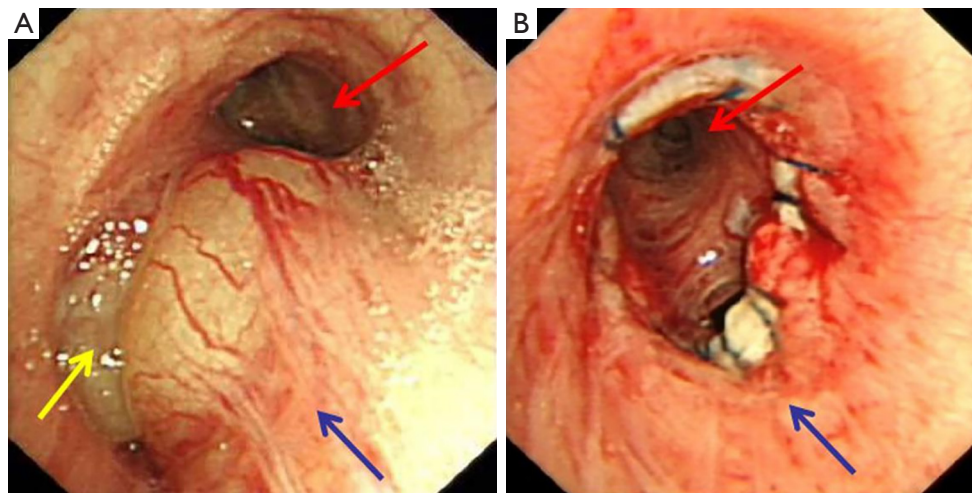


Figure 2 The bronchoscopy images of pre-operation and post-operation. (A) The preoperative bronchoscopy revealed that a neof ormation protruded into the lower part of left main bronchus (blue arrow) and left lower lobe bronchus (yellow arrow) was almost completely obstructed but not the upper left bronchus (red arrow). (B) The postoperative bronchoscopy revealed the end-to-end anastomosis between the left main bronchus (blue arrow) and the upper left bronchus (red arrow) after the left lower lobectomy.

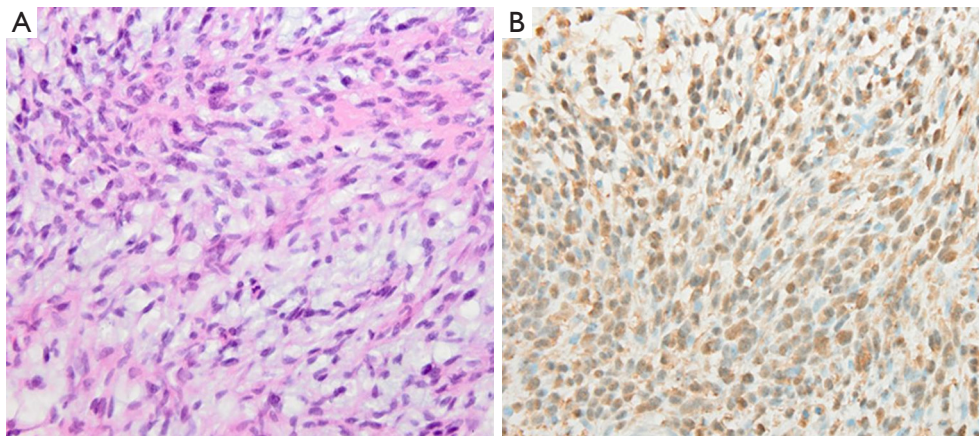


Figure 3 The pathological pictures. (A) The tumor was composed predominantly of uniform spindle cells with ovoid-to-round-shaped nuclei located in the eosinophilic collagen (H&E staining, $\times 200$); (B) immunohistochemistry with S-100 protein was positive ($\times 200$).

reports of schwannoma with malignant transformation, and the prognosis of schwannomas usually is optimistic (8). However, long follow-up is needed for monitoring the tumor growth and reducing the risk of recurrence after resection.

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None.

Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

Informed Consent: Written informed consent was obtained from the patient for publication of this manuscript and any accompanying images.

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