A case of endobronchial leiomyoma treated by sleeve resection of the right upper lobe bronchus

Bolesnik sa endobronhijalnim lejomiomom lećen sleeve resekcijom bronha za gornji režanj desnog pluća

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Abstract

Introduction. Bronchial leiomyoma is extremely rare. Most reported have been resected by either lobectomy or pneumonectomy. We presented a case treated by sleeve bronchoplasty without pulmonary resection. Case report. The presented case, 39-year-old male, had been admitted to our hospital complaining of hemoptysis. Chest X-ray showed no abnormality in either lung field, but computed tomography scan found the tumor in the upper right bronchus. The diagnosis was made by histological and immunohistochemical examination of the specimens obtained during bronchoscopy. Conclusion. The presented patient was treated by thoracotomy and sleeve resection of the right upper lobe bronchus with the removal of all the tumor.

Key words: bronchial neoplasms; leiomyoma; diagnosis; thoracic surgical procedures; treatment outcome.

Introduction

Primary pulmonary leiomyomas are extremely uncommon both in adults and children, constituting approximately 2% of benign lung tumors 1. They are thought to arise from the smooth muscle of the bronchus 2. The affected patients usually have respiratory symptoms due to partial or complete airway obstruction which deteriorate persisting asthma 1, 2 or be complicated with bronchiectasis and recurrent pulmonary infection 1,3,4.

Case report

A male, 39-years old smoker, was admitted to hospital due to months’ long polymorphic problems in the form of productive cough with haemoptysis, feeling of languor, fatigue and haemiparesthesia (feeling of numbness experienced on the left side of his body). Chest radiograph was normal (Figure 1), while computed tomography (CT) showed intraluminal, round, clearly circumscribed lesion in the upper lobe bronchus, 9.5 × 10 × 12.3 mm in diameter, which was post contrast homogeneously coloured without mediastinal lymphadenomegaly (Figure 2). During bronchoscopic examination, a cystic, moderately vascularized tumor change was noticed on the bronchus carina for the upper lobe bronchus, which was of smooth surface, white color, and softer consistency. The surrounding was moderately hyperemic without signs of mucosa infiltration. After bronchoscopy, a histopathological examination of the tumor change led to a leiomyoma diagnosis. Histologically, tumor tissue was built of...
intertwined bundles of spindle medium sized cells, abundant pale eosinophilic cytoplasm and elongated nuclei, yet edges that were blunt as in cigarettes. There was hyaline binder among the cells. Mitosis was not present. It was used enlargement x10, and HE staining (Figure 3). Tumor cells were immunoreactive for desmin and alpha-SMA (Figure 4) and immunonegative for CD117, CD34, MyoD1, S-100 protein and vimentin. Due to neurological problems in the form of paraesthesia of the left part of the body, a CT examination of endocranium had been done which detected no pathological changes. After adequate preoperative preparation, the patient was operated under general anesthesia, when right anterolateral thoracotomy was performed. After cutting the initial portion of the right upper lung lobe bronchus, endoluminally was registered a smooth soft tumor, of soft consistency, whose narrow peducle went from lower lip of bronchus carina for the upper lobe bronchus towards the intermedial bronchus.

Circular resection of the initial part of bronchus for the upper lobe was done with tumor, wherein the section surface of the same with no tumor, as well as resectional surface of the lower lip carina from intermedial bronchus were ex tempore histopathologically examined. After that anastomosis was performed by separate sutures. Post-operative care past with no complications. The control chest radiogram was satisfying. The control laboratory findings were within normal ranges. The definitive histopathological finding matched Hialeah leimyoma (Leiomyoma hyalineum).

Discussion

Leiomyomas of the lung are extremely rare and account for less than 2% of all benign tumors of the lung. These neoplasms can occur in parenchymal, endotracheal or endobronchial locations. Generally it seems that pulmonary leiomyomas affect females more than males with a ratio of approximately 1.5 : 1. Dyspnoea, cough and haemoptysis are the most common symptoms in those patients. On the chest X-ray pneumonitis is usually seen because of infection, which is the result of stasis of secretion. CT scan help us to locate the tumor, but bronchoscopy is a much more helpful diagnostic procedure to define the location of the tumor, and to get histopathological diagnosis, which help us in planning the operation. When the tumor is pedunculated and small, bronchoscopic resection is also useful. Bronchial leiomyomas are thought to derive from smooth muscle layer of bronchi, bronchioles, or blood vessels.

Conclusion

Leiomyomas of the respiratory system are essentially treated with surgical or bronchoscopic resection, depending on the location of the tumor. In some cases those tumors can be treated as conservative as possible, since the tumor is benign. In the presented case we performed a “sleeve” resection of the upper lobe bronchus, and after that anastomosis with the intermediate bronchus. In some other circumstances, with a secondary parenchymal destruction, leiomyomas of the bronchus may be treated with anatomic resection, like segmentectomy, lobectomy and in rare situation with pneumonecetomy.

REFERENCES


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