A Case of Recurrent Plasmacytoma with Endobronchial Involvement

Endobronşiyal Tutulum ile Birlikte Rekürren Plazmositom Olgusu

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Abstract
Extramedullary plasmacytomas and/or endobronchial plasmacytomas are rare conditions. We present this case due to its rarity. The patient who was diagnosed with extramedullary plasmacytoma of the right vocal cord ten years ago and nasopharynx two years ago was referred to our unit with hemoptysis. A suspicious lesion was detected in the right intermediate bronchus in the thoracic computed tomography (CT) scan and bronchoscopy was performed. A wide-based polypoid lesion in the distal part of the right intermediate bronchus was observed. Examination of the biopsy samples from the endobronchial polypoid lesion showed the presence of plasmacytoma. Extramedullary plasmacytomas are those which involve any soft tissue other than the bone marrow with the upper respiratory tract, being the most frequent site of extramedullary involvement. There are only six cases of endobronchial plasmacytoma in the English literature.

Key words: Plasmacytoma, extramedullary plasmacytoma, endobronchial plasmacytoma.

Pulmonary involvement may be the initial sign of multiple myeloma or may occur due to a primary pulmonary plasmacytoma (PPP). In contrast to the poor prognosis of multiple myeloma with pulmonary involvement, PPPs are likely to have a better prognosis. In a patient series, the lower airway involvement was found to occur in 4.7% of the patients with multiple myeloma (1). Primary endobronchial plasmacytoma (PEP) is a much less frequent condition with only six cases reported in the English literature.

Of these cases, two were treated with Yttrium-Aluminum-Garnet (YAG) laser, three with surgical resection, and one with rigid bronchoscopic debulking using the argon plasma coagulation (APC) (2-7).

We present this rare PEP case due to the obscurity surrounding its definition, and absence of published diagnostic and therapeutic algorithms.

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CASE

A 62-year-old male patient was diagnosed with an extramedullary plasmacytoma after histopathological examination of the biopsy specimen obtained from the right vocal cord 10 years ago. This was followed by local radiotherapy. Two years ago, he had another new-onset nasopharyngeal lesion, which proved to be a plasmacytoma after histopathological examination. Subsequent scans and bone marrow biopsy showed no involvement at other sites, shifting the focus of diagnosis from multiple myeloma to a plasmacytoma. Thus, he was re-administered radiotherapy.

The patient presented to our unit with hemoptysis. Thoracic computed tomography (CT) showed a suspicious lesion in the right main bronchus and bronchoscopy was performed (Figure 1). The bronchoscopy showed a wide-based polypoid lesion in the distal right main bronchus (Figure 2), as well as a mucosal irregularity at the upper third of the tracheal mucosa. The histopathological examination of the biopsy samples confirmed the diagnosis of amyloidosis in the tracheal lesion and a plasmacytoma in the endobronchial polypoid lesion.

Figure 1: A suspected nodular lesion in the right main bronchus

A positron emission tomography (PET) showed no involvement in other sites or at the site of the endobronchial lesion. Bone marrow biopsies did not show any systemic involvement and the protein electrophoresis demonstrated monoclonal immunoglobulin lambda-band. Urinary immune-fixation electrophoresis showed a monoclonal band of lambda-light chain. A diagnosis of primary endobronchial pulmonary plasmacytoma was established on the basis of these findings. An endobronchial intervention was planned (YAG-laser). However, the patient refused further work-up and is currently under follow-up without any treatment.

DISCUSSION

Extramedullary plasmacytomas are plasma cell tumors which tend to arise from the mucosal lymphoid tissue of the upper airways. They may involve any soft tissue other than the bone marrow with the upper respiratory tract being the most frequent site of involvement (80%) (8). While the upper airway involvement is more common in men, no sex differences have been reported for pulmonary involvement (8). The most frequent radiological appearance are pulmonary nodules and mass lesions in the surrounding hilus. Also, lobar consolidation and diffuse infiltration have been reported (1), as well as multiple pulmonary nodules (9) or cystic lung lesions (10). In our patient, we detected a suspicious pulmonary nodule and it was not clearly diagnostic. Therefore, the final diagnosis was able to be made only after bronchoscopy. An endobronchial involvement of plasmacytoma is not very well-known and often includes the mucosal involvement. In addition to rare polypoid tumors, it may also mimic a bronchogenic carcinoma. In our case, polypoid lesions were detected.

Also, some cases were diagnosed after thoracotomy performed for cystic lesions with a pre-diagnosis of aspergillosis (8, 10), as well as those diagnosed with video-assisted thoracoscopic surgery (11). In these patients, the histopathological diagnosis was based on immunohistochemical methods (Ig kappa light chain monoclonality). After establishing a diagnosis of plasmacytoma with immunohistochemical analysis following surgical resection of the tumor, a diagnosis of multiple myeloma should also be performed to exclude multiple myeloma (8).
our case, the diagnosis was based on bronchoscopic biopsy and PET scanning and bone marrow biopsy were performed to rule out multiple myeloma.

No clearly-defined treatment protocols have been developed for this condition, yet. Some cases undergoing surgical treatment also receive radiotherapy, while others receive both chemotherapy and radiotherapy. On the other hand, some authors have reported good long-term survival and excellent results with surgery (10). The majority of the patients are treated with surgical resection alone (8). Radiotherapy is generally recommended for the plasmacytomas of the head and neck. Although there is no clear recommendation on the use of radiotherapy in the treatment of endobronchial plasmacytomas, some authors have suggested that endoscopic resection and laser may represent inadequate therapeutic options, necessitating the use of radiotherapy (2). In case of a primary endobronchial plasmacytoma, YAG-laser, surgical resection, and rigid bronchoscopic debulking with argon plasma coagulation (APC) can be used (2-7). In our case, laser-YAG treatment was offered. However, the patient did not give consent for the treatment and we were unable to perform laser-YAG treatment. Currently, he has been under our follow-up without treatment for two years. The most important prognostic factor in extramedullary plasmacytomas is the transformation into multiple myeloma. There is a lack of information on its prognosis, since no survival differences between different therapeutic modalities are able to be shown (8).

Multiple or recurrent plasmacytomas may occur in 5% of the cases (1). In patients with multiple or frequent recurrences, systemic chemotherapy and autologous stem cell transplantation represent an important therapeutic option. In our case, an upper respiratory tract plasmacytoma was present in multiple locations and subsequently a pulmonary lesion was also detected.

In conclusion, a possibility of recurrence in patients who were previously diagnosed with an extramedullary plasmacytoma at another site should be kept in mind. Although an endobronchial plasmacytoma is a rare entity, it should be considered in the differential diagnosis of endobronchial neoplastic lesions. Treatment protocols and survival have not been clearly defined, yet.

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CONFLICTS OF INTEREST
None declared.

AUTHOR CONTRIBUTIONS

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