Extramedullary endobronchial plasmacytoma

Sebastián Fernández-Bussy, Gonzalo Labarca, Erik Folch, Adnan Majid

Division of Interventional Pulmonology, Clinica Alemana-Universidad del Desarrollo, Santiago, Chile
Clinica Alemana-Universidad del Desarrollo, Santiago, Chile
Division of Thoracic Surgery and Interventional Pulmonology, Beth Israel Deaconess Medical Center, Harvard Medical School, Santiago, Chile

Correspondence to
Dr. Sebastián Fernández-Bussy, sfernandezbussy@alemana.cl

DESCRIPTION
A 58-year-old man with a history of multiple myeloma was admitted to the hospital with progressive dyspnoea and hypoxemia. Chest examination revealed decreased breath sounds bilaterally. Blood examination revealed anaemia. Chest x-ray and chest CT scan revealed multiple pulmonary nodules, segmental atelectasis and central bronchial narrowing. Flexible bronchoscopy showed nodular lesions with extensive mucosa and submucosal infiltration, which extended from the proximal bronchus intermedius to the right lower lobe bronchus. A large nodular lesion was also noted at the posterior membrane of the proximal left mainstem bronchus (figure 1) and in the right bronchus intermedius (figure 2). Biopsies were taken with flexible forceps.

Endobronchial biopsies showed extensive tumour infiltration with plasmacytoid cells, ovoid irregular eccentric nuclei, prominent nucleoli, eosinophilic cytoplasm with a paranucleus pale area. High mitosis rate and frequent apoptotic figures were seen (figure 3). Immunohistochemistry was positive for CD-138 and MUM-1. Only k-light immunoglobulin chains were noted. This results suggest that the tumor was plasma cell origin with light chain restriction. Extramedullary plasmacytoma (EMP) is a plasma cell neoplasm that arises outside the bone. EMP is rare and is observed in only 5% of all plasmacytomas. Endobronchial plasmacytoma is a rare manifestation from EMP.

Figure 1 A large nodular lesion was noted at the posterior membrane of the proximal left mainstem bronchus that extends distally with mucosal erythema.

Figure 2 Extensive mucosa and submucosal infiltration at the distal-end of the bronchus intermedius with complete narrowing of the middle lobe and severe narrowing of the right lower lobe.

Figure 3 Left: ×40 hematoxylin and Eosin staining of the bronchial partition; presence of a lymphoproliferative infiltration. Right: ×100 hematoxylin and Eosin staining showing atypical plasma cells with a high mitosis rate.

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Learning points

▸ Extramedullary plasmacytoma (EMP) is an uncommon condition associated with haematological disease when neoplastic plasma cells arise outside the bone.
▸ Lower airways compromise is an unusual presentation of EMP; few cases have been reported.
▸ Histopathological study is required for diagnosis; bronchoscopy is an effective tool to visualise and obtain a biopsy sample.

Competing interests  None.
Patient consent  Obtained.

REFERENCES