Malignant Pleural Mesothelioma Detected by Endobronchial Ultrasound-Guided Transbronchial Needle Aspiration

**Abstract**

A 63-year-old man was admitted to our clinic with left hilar lymphadenopathy on his chest radiography. Fluorodeoxyglucose positron-emission tomography showed multiple regions of high metabolic activity on the left pleura and left interlobar lymph nodes. Histopathological examination of the lymph node material obtained by endobronchial ultrasound-guided transbronchial needle aspiration (EBUS-TBNA) revealed malignant mesothelioma. This case report demonstrates the usefulness of EBUS-TBNA in the diagnosis of metastatic mesothelioma.

**Key words:** Endobronchial ultrasonography (EBUS), mediastinal lymph node, mesothelioma.

Malignant mesothelioma is a rare and fatal neoplasm, which is strongly associated with asbestos exposure. Malignant pleural mesothelioma (MPM) arising from the parietal pleura is more frequent (65 to 70%), as inhalation is the typical route of asbestos pathogenicity (1). Various techniques have been used for the diagnosis of MPM, including thoracentesis with pleural fluid cytology and closed pleural biopsy; however these techniques may not obtain enough tissue to confirm the diagnosis. Video-assisted thoracoscopic surgery (VATS) is recommended for a definitive diagnosis; however, VATS is an invasive and expensive technique (2). A minimally invasive procedure, endobronchial ultrasound-guided transbronchial needle aspiration (EBUS-TBNA), provides sampling of mediastinal and hilar lymph nodes, as well as centrally located lung lesions. In the literature, there is only one study in which the value of the EBUS-TBNA was investigated in the diagnosis of MPM (3). Herein, we present a case with MPM diagnosed using EBUS-TBNA.

**Özet**

Posteroanterior akciğer grafinde sol hiler lenfadenopati saptanan 63 yaşında erkek hasta kliniğimize başvurdu. PET-CT'de sola lenf bezlerinde ve plevralda artmış florodeoksiglukoz tutulumu saptandı. Endobronşyal ultrason transbronşyal iğne aspirasyonu (EBUS-TBİA) ile elde histopatolojik incelmede malign mezeotelyoma saptandı. Bu olgu ile EBUS-TBİA'nın metastatik malign mezeotelyoma olgularında kullanımı faydali olduğu gösterildi.

**Anahtar Sözcükler:** Endobronşyal ultrason (EBUS), mediastinal lenfadenopati, mezeotelyoma.
CASE
A 63-year-old male patient with a 102-pack per year smoking history was admitted to the outpatient clinic for the evaluation of a left hilar lymphadenopathy. He did not have a history of asbestos exposure; however, he had a history of left pleural effusion from 10 years previously. No malignant cells were found after a closed pleural biopsy, followed by computed tomography (CT) once per year.

No abnormalities were found at his physical examination. Laboratory tests showed high CRP levels (9.40mg/dL, normal level: 0-0.8 mg/dL), and the Hb level was 11.3g/dL (normal level: 13.5-18g/dl). Chest X-ray revealed a left hilar fullness, which was an enlarged hilar lymph node on the contrast-enhanced CT scan. There was pleural thickening, but no pleural effusion (Figure 1a).

Subsequently, EBUS-TBNA was performed to obtain a tissue specimen for diagnosis (Figure 1b). The cytopathological examination showed a spindle cell malignancy with vimentin. Calretinin and CD31 expression by immunohistochemistry was compatible with malignant mesothelioma (Figure 2a and b). The EBUS-TBNA findings were confirmed with a tru-cut biopsy (Figure 2c).

DISCUSSION
Mesothelioma is the most common primary malignant tumor of the pleura. It is a very rare disease, originating from the mesothelial cells lining the pleura. Extrapleural nodal metastasis is unusual in mesothelioma, while hilar and mediastinal lymph node involvement occurs in less than 50% of the patients (1).

The pathogenesis of this disease remains unclear, as the information about the role of asbestos exposure in the development of malignant mesothelioma can not be defined for the majority of cases. In such cases, dust exposure, chemicals, genetic factors, and viral infections have been reported (4). There was no asbestos exposure in our case, either; however, our case was a chronic smoker. The rarity of the disease, the absence of asbestos exposure, and the unusual presentation prompted us to report this case.

Radiological modalities are currently available for the detection of nodal metastases in patients with MPM. Computed tomography (CT) has a sensitivity of 60% in detecting nodal involvement. In recent years, the role of FDG-PET has been increasing in the evaluation of MPM (4). However, imaging studies can not replace pathological examinations, and invasive procedures may be needed for a definitive diagnosis (4). For instance, video-assisted thoracoscopic biopsy has a high sensitivity for diagnosis; however, it is a more invasive procedure which usually requires general anesthesia. In addition, EBUS-TBNA is a proven alternative to surgery for sampling mediastinal and hilar lesions. It is recognized as an accurate and minimally invasive procedure for the diagnosis of hilar and mediastinal lymph nodes (3).

As MPM is an aggressive disease, early diagnosis must be achieved for successful treatment. Despite the advances in our understanding of the pathobiology of MPM, the definitive diagnosis of MPM still requires histopathological tissue examinations, usually via direct thoracoscopic biopsy, combined with a panel of immunohistochemical markers (4). Furthermore, MPM is a very heterogeneous malignancy, often resulting in misdiagnosis with other conditions. Mesothelioma is classified into three major histological types: epithelioid, sarcomatoid and mixed. The best prognosis is in the epithelioid type, which is the most common (50 to 60 %). Clinical and radiographic evaluations alone may be often insufficient for the diagnosis, while thoracoscopy is the most of definitive method (4).
In the literature, there are few cases about malign pleural mesothelioma diagnosed by EBUS-TBNA. In one case, EBUS-TBNA was applied to subcarinal lymph node and MPM was diagnosed (5). In our case, histological specimens were obtained from the left interlobar lymph node by EBUS-TBNA, which is a minimally invasive procedure. Therefore, we showed that EBUS-TBNA is an excellent and safe tool for obtaining adequate histological nodal tissue for examination to achieve a rapid and definitive pathological diagnosis.

CONFLICTS OF INTEREST
None declared.

AUTHOR CONTRIBUTIONS

REFERENCES