Malignant fibrous histiocytoma (MFH) is the most common soft tissue sarcoma in adults, most frequently involving the extremities, retroperitoneum, and trunk, but it is rare as a primary tumor of the lung [1–3]. Chest roentgenography, chest computerized tomography (CT), and magnetic resonance imaging (MRI) disclose a well-defined mass that is not very lobulated [3–5]. Early diagnosis is very important because of the poor prognosis. Bronchoscopy and percutaneous needle aspiration are of limited value in differential diagnosis, and thoracotomy is warranted for definitive diagnosis [2,6]. Surgery is the primary mode of therapy [7].

We report a case of primary MFH of the lung illustrating the clinical, radiologic, histologic, immunohistochemical, and ultrastructural features, and the difficulty in treating such cases.

**Case Presentation**

An 86-year-old male was admitted to our hospital in February 2002 due to insidious onset of exertional dyspnea and poor appetite lasting for 1 month. He had previously been quite well and had no history of smoking. Physical examination revealed pale conjunctivae and decreased breath sounds over the left basal lung field. Laboratory examinations showed mild anemia, hyponatremia, hypoalbuminemia, and a high C-reactive protein concentration. Tumor markers, including carcinoembryonic antigen, tissue plasmin antigen, and squamous cell carcinoma antigen, were within normal limits. Chest roentgenography demonstrated a 9 × 15 cm, pleural-based opacity in the left lower lobe (Figure 1). Chest CT revealed a well-
defined mass with heterogeneous density in the left lower lung field. Local extension to the left chest wall and right pleural effusion were noted (Figure 2). Gallium whole-body scan revealed intensively increased radioactivity in the anterolateral aspect of the left lower chest with a photopenic area in the adjacent middle portion and gradually increasing contrast throughout the serial 6-, 24- and 48-hour images.

Thoracoscopic lung biopsy yielded eight tissue fragments, measuring up to $1.8 \times 1.1 \times 0.8$ cm. Grossly, they were reddish, soft and elastic, or whitish, soft and jelly-like. A storiform pattern of spindle cells and multinucleated giant cancer cells with atypical mitotic figures were seen on microscopy (Figures 3 and 4). Hemorrhage and necrosis were noted. Immunohistochemical studies were negative for cytokeratin, CK5/6, and calretinin, but positive for vimentin. Therefore, MFH was diagnosed. Because the patient was extremely old in age and had poor performance status (Karnofsky performance status = 30), he received supportive care. He died in April 2002 from respiratory failure.

**DISCUSSION**

MFH is the most common (10%) soft tissue sarcoma in adults and mostly occurs in males. It mainly arises from soft tissue in the extremities (68%), the retroperitoneum (16%), and lung (7.5%) [1]. Primary MFH comprises 0.04% of all pulmonary malignant tumors [2]. There must be no evidence of another primary site on careful physical examination and radiologic evaluation before MFH can be considered a primary rather than a metastatic lesion. The mean age of patients with primary lung MFH is 55 years (range, 10–80 years) [7,8]. There are no predisposing factors. The most common complaints are cough, dyspnea, fatigue, hemoptyis, weakness, headache, fever, chills, weight loss, and chest pain. Patients may be asymptomatic at presentation [7,9–11]. Our patient was older than usual (86 years old), and his initial presentation was exertional dyspnea and poor appetite.

Typically, lung MFH appears as a large, solitary, well-circumscribed, non-cavitating, non-calcified, peripheral lung mass with a slight preponderance in the middle and lower lobes. Approximately 20% of patients have associated pleural effusion [3,4]. CT and MRI can provide useful information to demonstrate the nature and extent of tumor invasion [5]. On CT, lung MFH appears to be of soft-tissue density, sometimes with central areas of low attenuation. The tumor displaces rather than invades local structures. In our case, imaging showed a huge, well-defined tumor in the left lower lobe with soft-tissue density.

Figure 1. Chest roentgenography shows a 9 × 15 cm, pleural-based opacity in the left lower lobe.

Figure 2. Chest computerized tomography reveals a well-defined mass with heterogeneous density in the left lower lung field. Local extension to the left chest wall and right pleural effusion are evident.
and no mediastinal or hilar lymph node involvement. The tumor had local extension to the left chest wall and right pleural effusion was also noted.

Immunohistochemical examination permits the line of differentiation to be established in a number of primary or metastatic sarcomas of the lung. Mesenchymal neoplasms are positive for vimentin and negative for cytokeratin. Both leiomyosarcoma and rhabdomyosarcoma contain myoglobin. S-100 protein is a marker for neurogenic sarcoma. MFH tumor cells show positive reactions for vimentin, α₁-antitrypsin and α₁-antichymotrypsin, but are negative for cytokeratin, desmin, myoglobin, smooth muscle actin, and S-100 protein [8]. Enzinger and Weiss described four histologic types of soft-tissue MFH: storiform pleomorphic; myxoid; giant cell; and inflammatory [12]. Most primary lung tumors are of the storiform variety. No definite relationship between survival and histology has been identified because of the predominance of the storiform subtype. The most significant prognostic factors are the presence of metastases, local recurrence, and more than 15% necrosis [13]. In our case, histology revealed a storiform pattern in spindle cells and multinucleated giant cells. Hemorrhage and necrosis were noted. Immunohistochemical studies were negative for cytokeratin, CK5/6, and calretinin, but positive for vimentin.

Bronchoscopy and percutaneous needle aspiration are of limited value in differential diagnosis, and thoracotomy is warranted for definitive diagnosis [2,6]. Mediastinoscopy is not useful because MFH usually disseminates hematogenously and does not commonly invade lymphatics [2,12]. The brain is a common site of metastasis [9,14–17].

Complete tumor resection is the treatment of choice, with high rates of local and distant recurrence reported. The roles of radiation and chemotherapy have not been clearly defined [7]. Systemic chemotherapy has been used largely for metastatic disease, but has yielded poor results [7,15,18,19]. In our case, the patient only received supportive care due to his extreme old age and poor performance status.

In conclusion, MFH of the lung is a rare clinical entity with variable outcome. Aggressive surgery is the mainstay of treatment. The role of adjuvant radiation and chemotherapy remains undefined.

**References**

Primary malignant fibrous histiocytoma of the lung