Malignant fibrous histiocytoma of the mediastinum

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Summary: A case of malignant fibrous histiocytoma of the mediastinum presenting with unusual features of fever and leucocytosis is reported. This is the youngest patient reported in the literature who had this tumour in the mediastinum.

Introduction

Malignant fibrous histiocytoma is a neoplasm with fibroblastic and histiocytic differentiation that arises usually in the deep soft tissues. It has a predilection for the extremities, especially thighs. It occurs very rarely in the mediastinum. This paper reports a case of malignant fibrous histiocytoma of the posterior mediastinum occurring in a young girl who presented with fever.

Case report

A 16 year old girl presented to us with a 6-month history of fever, anorexia and weight loss. The fever was low grade and intermittent initially but became high grade and continuous for one month prior to coming to this hospital.

Physical examination showed an emaciated girl with moderate pallor and a temperature of 38.5°C. The chest examination revealed findings suggestive of a mass lesion in the interscapular and infrascapular areas on the left side.

Investigations showed a haemoglobin of 4.5 g/dl, white blood cell count of 14.9 × 10^9/l (82% polymorphonuclear leucocytes), ESR was 60 mm in the 1st hour, serum iron was 78 μg/dl and total iron binding capacity 540 μg/dl. The peripheral smear suggested an iron deficiency anaemia. The liver function tests were unremarkable. A bone marrow aspirate showed a hypercellular marrow, erythroid hyperplasia with normoblastic marrow. A chest X-ray revealed a large posterior mediastinal mass on the left side without any bony destruction or calcification (Figure 1). Computerized tomography of the chest disclosed a large soft tissue mass in the posterior mediastinum on the left side with a small area of calcification (Figure 2).

Figure 1 Anteroposterior chest radiograph showing a posterior mediastinal mass on the left side.

A percutaneous biopsy of the mass was done which on microscopy showed a mixture of small and large cells, the latter containing an abundant eosinophilic cytoplasm and a vesicular nucleus. These cells had a vague resemblance to ganglion cells. The biopsy was interpreted as ganglioneuroma.

With a pre-operative diagnosis of ganglioneuroma, the patient was operated. At thoracotomy, a large 15 × 10 cm lobulated mass occupying the lower part of the posterior mediastinum and infiltrating the lower

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loge of the left lung was found. The tumour was excised along with the left lower lobe of the lung. The histopathology of this specimen revealed a variable appearance. Some areas showed spindle-shaped fibroblasts arranged in short fascicles and at places showing a stratiform pattern. In other areas, the tumour was markedly pleomorphic and cellular, composed chiefly of large bizarre cells with abundant eosinophilic cytoplasm and vesicular nucleus. Some cells were binucleate or multinucleated. Mitotic figures were seen in large numbers. There was infiltration by a variable number of chronic inflammatory cells chiefly the lymphocytes and plasma cells (Figure 3).

A final diagnosis of malignant fibrous histiocytoma of the posterior mediastinum was made. The fever and leucocytosis subsided after surgery. The patient was advised radiotherapy post-operatively. However, the patient declined it and was lost to follow-up.

**Discussion**

Malignant fibrous histiocytoma was first described in the mid-1960s. Since then, it has been recognized as the most common soft tissue sarcoma in adult life. It presents primarily in middle to late adult life. Out of 200 cases of this tumour reviewed, only 9 were younger than 20 years. It occurs most commonly in the extremities, thighs being the most common site. In two large series comprising 367 patients with this neoplasm, none had a mediastinal tumour.

Malignant fibrous histiocytoma arising in the mediastinum has rarely been reported and to the best of our knowledge, only 8 such cases have been described to date. The previous youngest patient to have this tumour was 33 years of age. One case was asymptomatic and was detected on a routine X-ray, while other patients had variable, nonspecific features in the form of cough, dyspnoea, back pain, superior vena cava obstruction or plural effusion. The present case had a unique presentation with fever and leucocytosis which have not been described in such tumours located in the mediastinum. In a review of 200 cases of this tumour situated elsewhere, only 2 cases had these features. The exact pathogenesis of these features is unclear but these could be due to secretion of lymphokines by the tumour in some patients. Out of 8 mediastinal tumours, 7 were in the posterior mediastinum while one tumour was situated in the anterior mediastinum.

The treatment of choice is surgical excision with or without radiotherapy. Both patients reported by Mills et al. received post-operative radiotherapy. One of them was alive and doing well 56 months after surgery while the second patient was lost to follow-up. Two other cases were alive after 17 months and 15 months after surgical removal alone. The remaining patients died during follow-up.

Malignant fibrous histiocytoma displays a pleomorphic histological picture because of which diagnosis is sometimes difficult with small biopsy specimens. In our case, the initial biopsy was interpreted as ganglioneuroma. In another reported case, the frozen section of the tumour was reported as ganglioneuroma.

The case of malignant fibrous histiocytoma reported here is of interest because of its origin in the mediastinum, its occurrence at a young age and the presence of fever and leucocytosis which remitted after surgery. Despite its rarity, the possibility of a malignant fibrous histiocytoma should be considered in the differential diagnosis of a mediastinal tumour.

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**Figure 2** Computed tomography of the chest showing a soft tissue mass with an area of calcification in the posterior mediastinum.

**Figure 3** Microscopic section showing a markedly pleomorphic tumour composed of small histiocytic cells, large bizarre cells and multinucleated giant cells. A few spindle-shaped cells are also identified (haematoxylin & eosin x 300).


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