Pulmonary involvement with multiple myeloma occurs infrequently and may be difficult to distinguish from more common primary lung tumours, metastatic disease, or other pleural and parenchymal abnormalities. A patient who developed acute respiratory distress syndrome (ARDS) was subsequently found to have multiple myeloma with involvement of lung parenchyma by neoplastic plasma cells. Only one other report of ARDS in association with multiple myeloma was found, and there are no previous reports where the appearance of ARDS antedated a diagnosis of multiple myeloma. In patients with ARDS, parenchymal involvement from multiple myeloma should be included in the differential diagnosis.

Ultrasound guided thoracentesis yielded 200 ml of bloody exudative effusion. Cytology was negative for malignant cells. Thoracoscopic biopsy revealed diffuse and nodular infiltration of the lung parenchyma by neoplastic plasma cells (fig 2). The 24 hour urine contained 918 mg protein, and immunoelectrophoresis revealed a monoclonal band in the gamma region identified as IgG kappa with excess kappa light chains. Serum immunoelectrophoresis similarly showed hypergammaglobulinaemia with a monoclonal IgG kappa band.

Treatment continued with piperacillin/tazobactam, azithromycin, and vancomycin for pneumonia, and methylprednisolone for myeloma. She was judged to be a poor candidate for more aggressive chemotherapy owing to progressive multisystem organ failure. In keeping with the patient’s advance directive and after consultation with her family, supportive care was withdrawn. Permission for bone marrow biopsy or necropsy was withheld.

**DISCUSSION**

This patient was diagnosed with multiple myeloma on the basis of a monoclonal serum protein spike, the presence of urinary Bence Jones proteins, and infiltration of the lungs by neoplastic plasma cells. She developed acute respiratory distress syndrome (ARDS) secondary to this nodular involvement of the lungs by the myeloma. Multiple myeloma presenting as ARDS has not been reported previously. Absence of leucocytosis on admission, failure to respond radiographically or clinically to antibiotics, the absence of positive cultures for microorganisms, and pathological findings without evidence of infectious pneumonia all argue against infection as a major factor contributing to the development of ARDS.

Multiple myeloma may involve the thorax in a variety of ways, but pulmonary parenchymal involvement is uncommon. Thoracic manifestations include skeletal abnormalities.
Multiple myeloma may produce varied patterns on chest radiographs including multiple masses mimicking solid tumour metastasis, diffuse interstitial disease from alveolar septal amyloidosis, or consolidation. Our patient presented with a chest radiographic pattern that was consistent with the diagnosis of ARDS, as defined by the American-European Consensus Conference on ARDS. However, both the ARDS Clinical Trials Network (in terms of inclusion criteria) and a recent review of "imitators" of ARDS highlight the pathophysiological heterogeneity that may underlie such a broad clinical definition. Since our patient had neither prior history nor clinical evidence of multiple myeloma, this pathological diagnosis was not made until an open lung biopsy was obtained. A greater clinical suspicion could conceivably have resulted in bone marrow biopsy, thereby obviating the need for lung biopsy. We suggest that multiple myeloma with direct plasma cell invasion of pulmonary parenchyma be added to the list of pathological mechanisms that can produce the clinical syndrome of ARDS.

**REFERENCES**


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Figure 2 Photonicrograph of thoracoscopic biopsy specimen. Stain: haematoxylin/eosin; original magnification ×10.