

Figure 3. Bronchoscopic biopsy: Hodgkin lymphoma, CD30+.

Keywords: Hodgkin lymphoma, Cavitory lung lesions, Tuberculosis

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Light Chain Myeloma with Highly Atypical Plasma Cells and Extensive Auer Rod-Like Inclusions

Yüksek Atipik Plazma Hücreleri ve Yaygın Auer Cisimciği Benzeri İnklüzyonları Olan Hafif Zincir Myeloma

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To the Editor,

A 73-year-old woman with a history of chronic kidney disease presented with fever (39.8 °C), dyspnea, and fatigue. Complete blood count showed moderate normocytic anemia with hemoglobin of 10.0 g/dL (normal range: 12.0-16.0), mild leukocytosis of $10.8 \times 10^9/L$ (normal range: 4.0-9.0), and thrombocytopenia of $10^2 \times 10^9/L$ (normal range: 150-400). Serum protein electrophoresis showed mild hypogammaglobulinemia of 6.7 g/L (normal range: 7.0-16.0). Serum immunofixation electrophoresis demonstrated monoclonal κ -type light chains without heavy chain correlates (IgG, IgM, IgA, IgD, IgE). Moreover, a serum-free light chain assay measured

Anahtar Sözcükler: Hodgkin lenfoma, Kaviter akciğer lezyonları, Tüberküloz

Informed Consent: It was received.

Conflict of Interest: The authors of this paper have no conflicts of interest, including specific financial interests, relationships, and/or affiliations relevant to the subject matter or materials included.

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a high κ -type light chain level of 2060.0 mg/L (normal range: 3.3-19.4) with a κ/λ ratio of 48.5 (normal range: 0.3-1.7).

A bone marrow aspirate smear showed 40% plasma cells, many of which appeared as binuclear plasmablastic cells with nucleoli ("owl-eyed" plasma cells), bright cytoplasm, and bundles of numerous Auer rod-like cytoplasmic inclusions (Figures 1A and 1B). This unique morphology is remarkable. While the current literature describes Auer rod-like inclusions in single cases of different forms of myeloma [1,2,3,4,5], this is, to the best of our knowledge, the first report on the concomitant appearance with enlarged highly atypical "owl-eyed" plasma cells in a patient suffering from κ -type light chain myeloma.

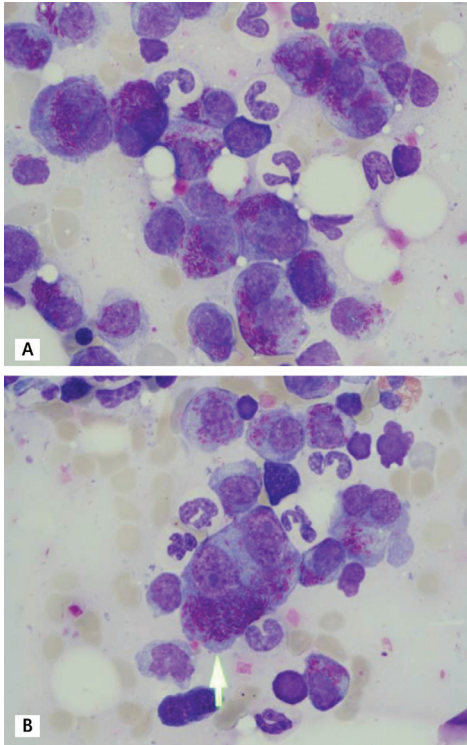


Figure 1. Bone marrow aspirate smear of a 73-year-old patient with κ -type light chain myeloma (A and B). The arrow marks a binuclear plasmablastic cell containing numerous Auer rod-like inclusions.

However, the prognostic value of this unusual plasma cell phenotype remains unclear.

Keywords: Light chain myeloma, Plasma cells, Bone marrow aspirate

Anahtar Sözcükler: Hafif zincir myeloma, Plazma hücreleri, Kemik iliği aspirasyonu

Informed Consent: Was obtained from the patient.

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Aggressive Clinicopathological Course of Myeloma with t(3;16) (q21;q22) Cytogenetic Abnormality

t(3;16)(q21;q22) Anomalili Myeloma Olgusunda Agresif Klinikopatolojik Seyir

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To the Editor,

Multiple myeloma (MM) is a heterogeneous disease and patients present with a wide variety of cytogenetic anomalies reflecting the nature of the disease [1]. The aim of this letter is to report a rare karyotypic abnormality with an aggressive clinical course of MM.

A 56-year-old male patient was admitted to the neurosurgery clinic with dorsal shoulder pain and inability to walk in April 2011. He underwent thoracic and lumbar spinal magnetic resonance imaging. Laminectomy was performed on the patient upon detecting masses at the levels of the first and seventh thoracic vertebrae. The patient was referred to our center when he was determined to have "lymphoma" based on the first evaluation