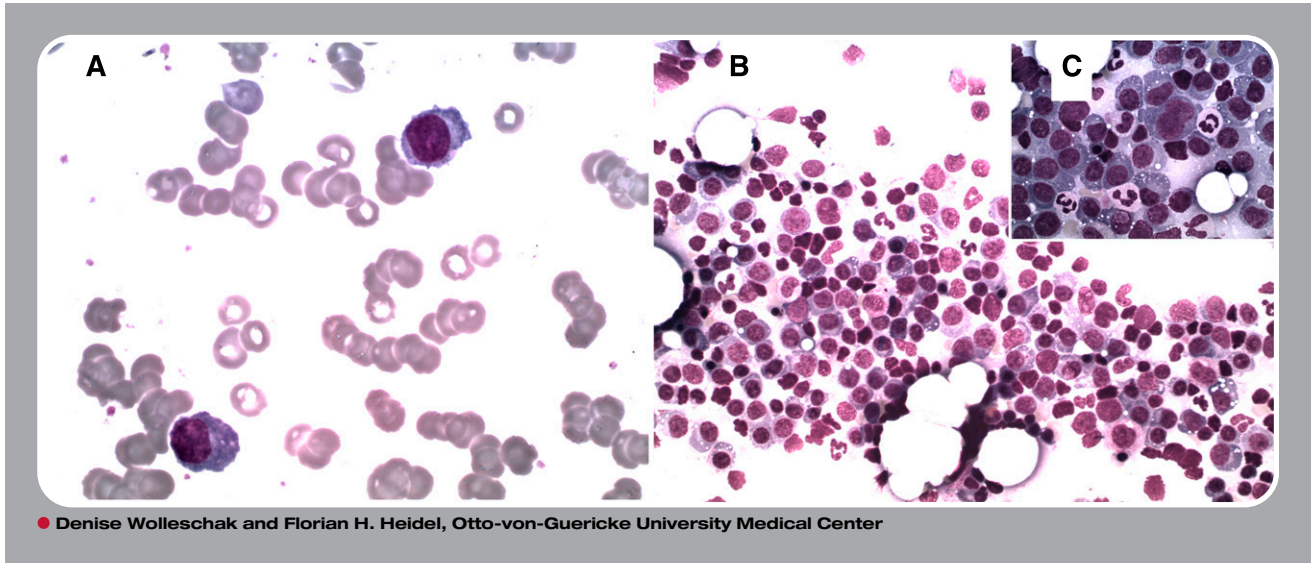


A rare cause of lower back pain



A 51-year-old man who was previously healthy was admitted to the local orthopedics department with lower back pain and weakness of the lower extremities. He presented with leukocytosis of $11 \times 10^9/L$ and his hemoglobin level was 90 g/L. Complete blood count with automated differential reported >20% large unstained cells. The peripheral smear showed rouleau formation and 23% of circulating plasma cells with atypical features such as a higher nuclear-to-cytoplasmic ratio, dispersed nuclear chromatin, and prominent nucleoli (panel A). A bone marrow aspirate smear demonstrated 70% plasma cell infiltration with cells of either mature or plasmablastic morphology and vacuoles in the cytoplasm (panels B-C). Serum immunofixation revealed production of IgA with λ light-chain restriction. The diagnosis was primary plasma cell leukemia (PCL), which is a rare presentation (4/10 000 000 persons per year) of multiple myeloma (MM).

Primary PCL is characterized by leukocytosis with circulating plasma cells that typically exceed 2000/ μ L or 20% of the peripheral white blood cell count. The bone marrow infiltration is usually extensive and consists of plasma cells with atypical and immature morphology. Compared with MM, PCL has a poor prognosis, with a median survival of 8 to 12 months.