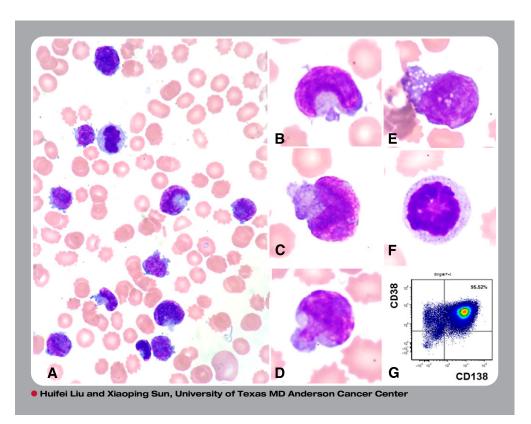


Pleomorphic plasma cell leukemia



T his peripheral blood smear is from a 72-year-old woman, taken several hours before she died of recurrent plasma cell myeloma and newly developed secondary plasma cell leukemia. Her white blood cell count increased from $39.5 \times 10^3/\mu$ L to $233.2 \times 10^3/\mu$ L in 28 hours. The peripheral blood smear shows marked leukocytosis with numerous small- to medium-sized bizarre-looking cells (95% of white blood cells) (panels A-E), which were not seen in her initial bone marrow aspirate. These cells have pleomorphic nuclear shapes such as a fish mouth, a mushroom, or a moon. All cells have scant pale cytoplasm, and some have cytoplasmic vacuoles. Mitotic figures are identified (panels A and F).

Despite the unusual cellular morphology, flow cytometric analysis confirmed these cells to be aberrant plasma cells (positive for CD38, CD138, partial CD56, and κ light chain, and negative for CD45, CD19, CD20, CD28, CD117, and λ light chain) (panel G). Her initial cytogenetics study showed a complex karyotype: $45 \sim 46$,XX,+1,add(1)(p13),der(3)t(3;11)(q29;q13),t(11;14)(q13;q32),+14, del(14)(q24). Fluorescence in situ hybridization reported monosomy 13, t(11;14) IgH/cyclinD1 translocation, CSK1B gene amplification, and TP53 gene deletion. Cytogenetics was not performed during her last admission. Her initial serum lactate dehydrogenase level was 30 489 IU/L (normal, 313-618 IU/L). Her initial and last serum β 2-microglobulin levels were 9.5 and 7.4 mg/L (normal, 0.8-2.3), respectively. This case highlights the broad morphologic spectrum of plasma cell myeloma/leukemia.



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