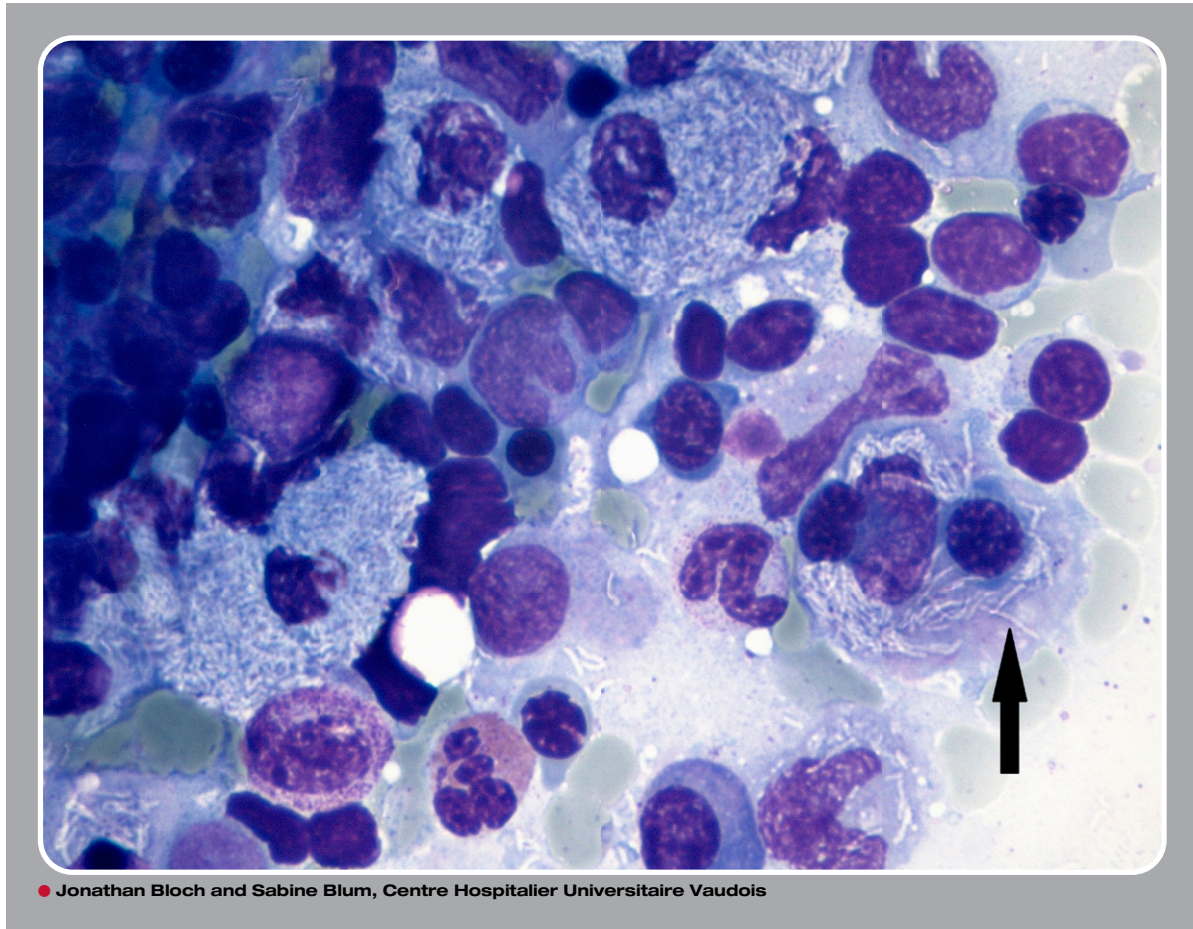


“Atypical” macrophages



A 72-year-old woman presented with a 1-month history of weakness. Medical history included a smoldering myeloma diagnosed 2 years before, an autoimmune pancytopenia treated with prednisone for the past year, and an episode of pulmonary nocardiosis successfully treated with co-trimoxazole 6 months before. Notably, the bronchoalveolar lavage that helped diagnose the nocardiosis also revealed *Mycobacterium avium* that was considered an innocent bystander. On admission, the patient suffered from pancytopenia (hemoglobin, 112 g/L; leukocytes, $2.0 \times 10^9/L$; platelets, $33 \times 10^9/L$; neutrophils, $1.29 \times 10^9/L$) with 6 CD4^+ T cells/ mm^3 . May-Grünwald-Giemsa coloration of the bone marrow aspirate showed an infiltration with numerous pseudo-Gaucher cells containing cell debris and rod-like elements that did not absorb the colorant (arrow). Periodic acid-Schiff staining confirmed that they were mycobacteria (not shown). The patient was started on antimycobacterial antibiotics, but discontinued therapy and died shortly after.

Pseudo-Gaucher cells consist of macrophages whose cytoplasm has a “watered silk” appearance as they are loaded with cellular debris. Pseudo-Gaucher cells have sometimes been described in mycobacterium- and HIV-infected patients. In this case, the patient had a low CD4^+ count due to immune pancytopenia and long-term steroid therapy that allowed the atypical mycobacterial infection to develop.