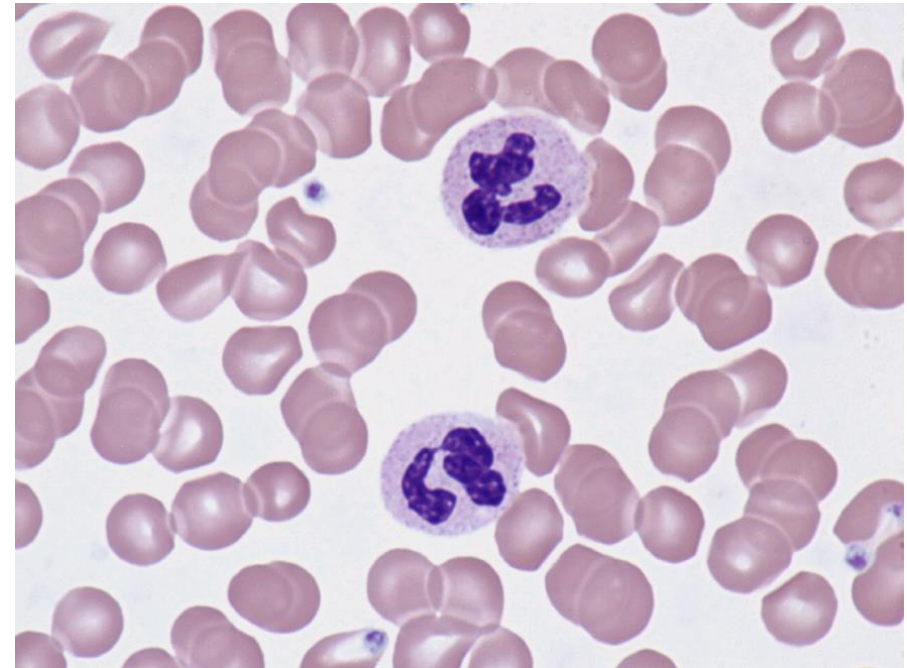


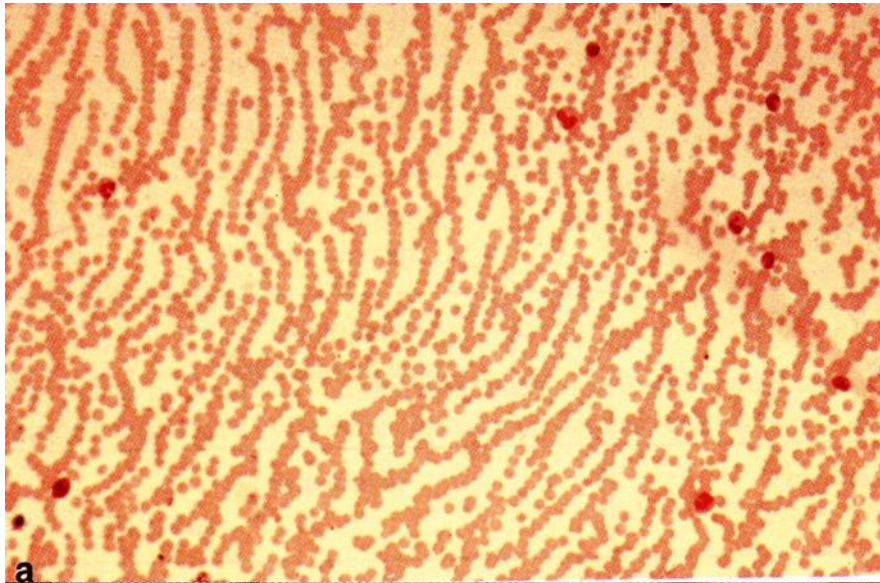
Tecniche di laboratorio di ematologia

Prof. Gian Matteo Rigolin
Ematologia

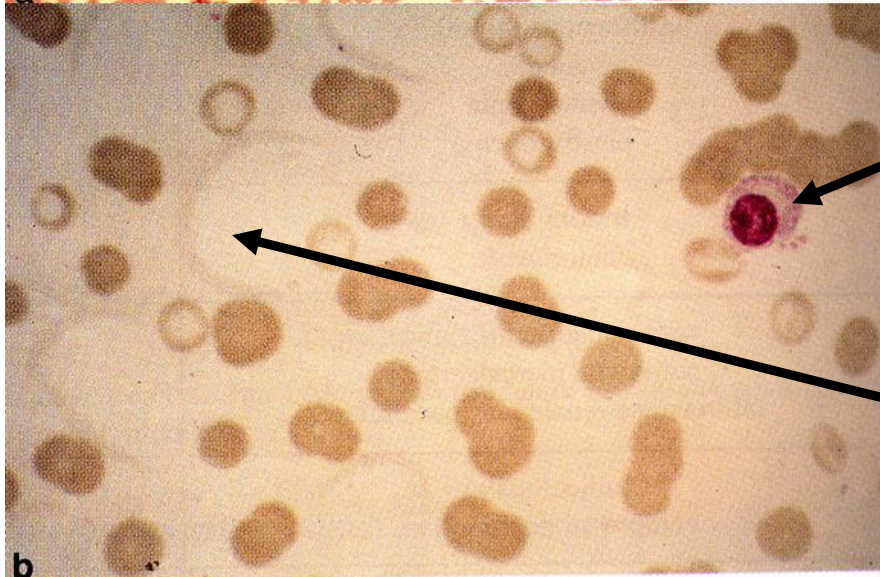


QUADRI MORFOLOGICI

MIELOMA MULTIPLO: sangue periferico



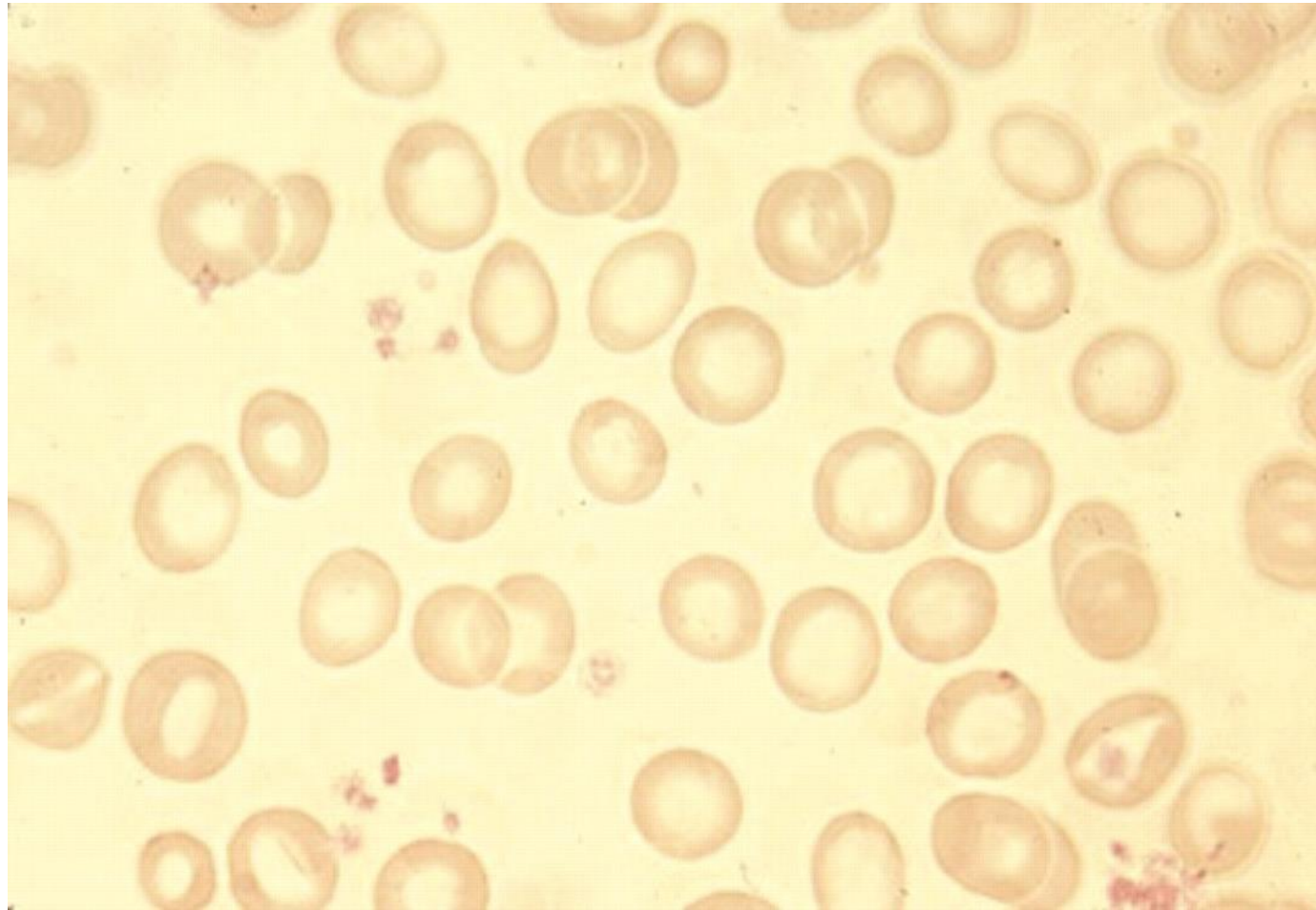
Impilamento delle emazie ("rouleaux")



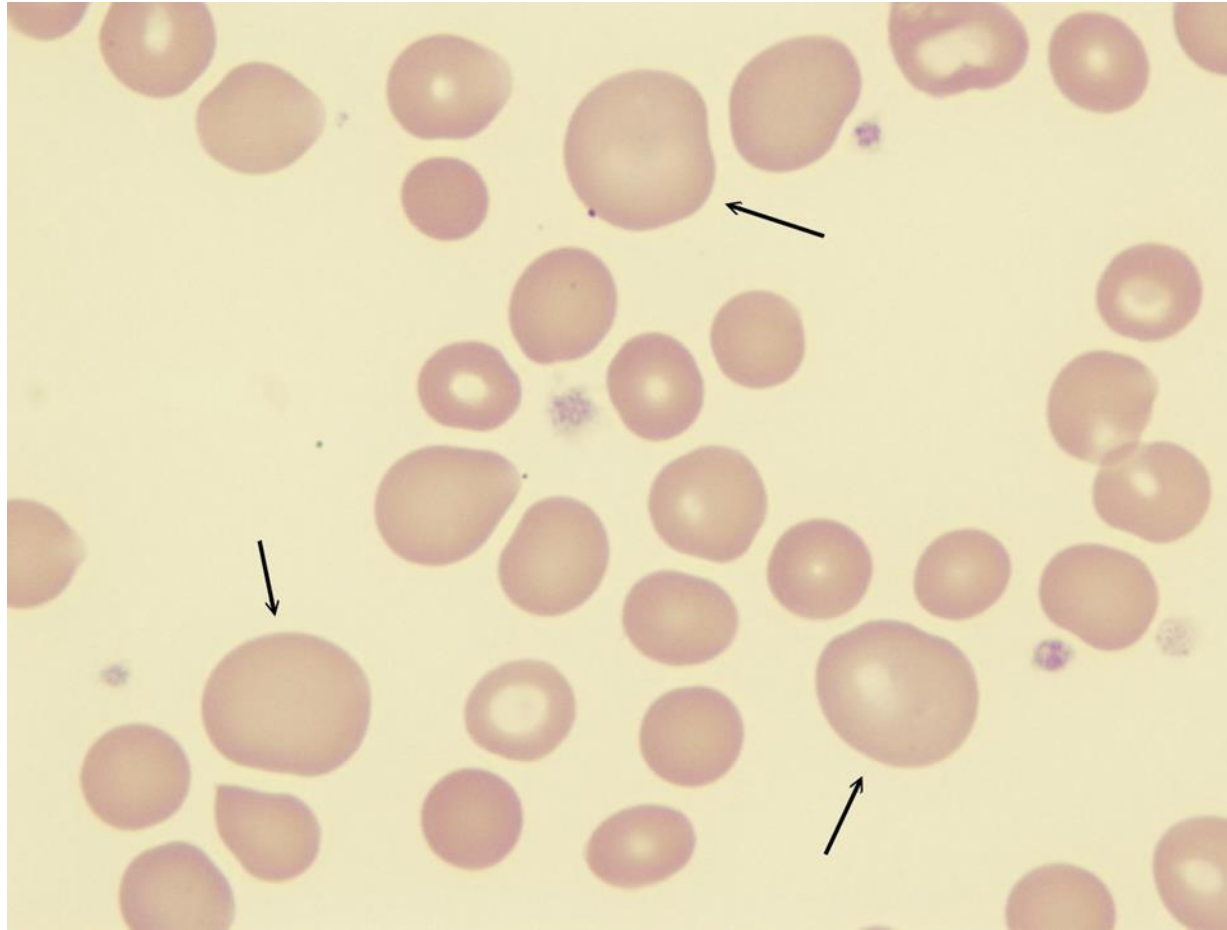
Plasmacellula circolante

Archi di precipitazione

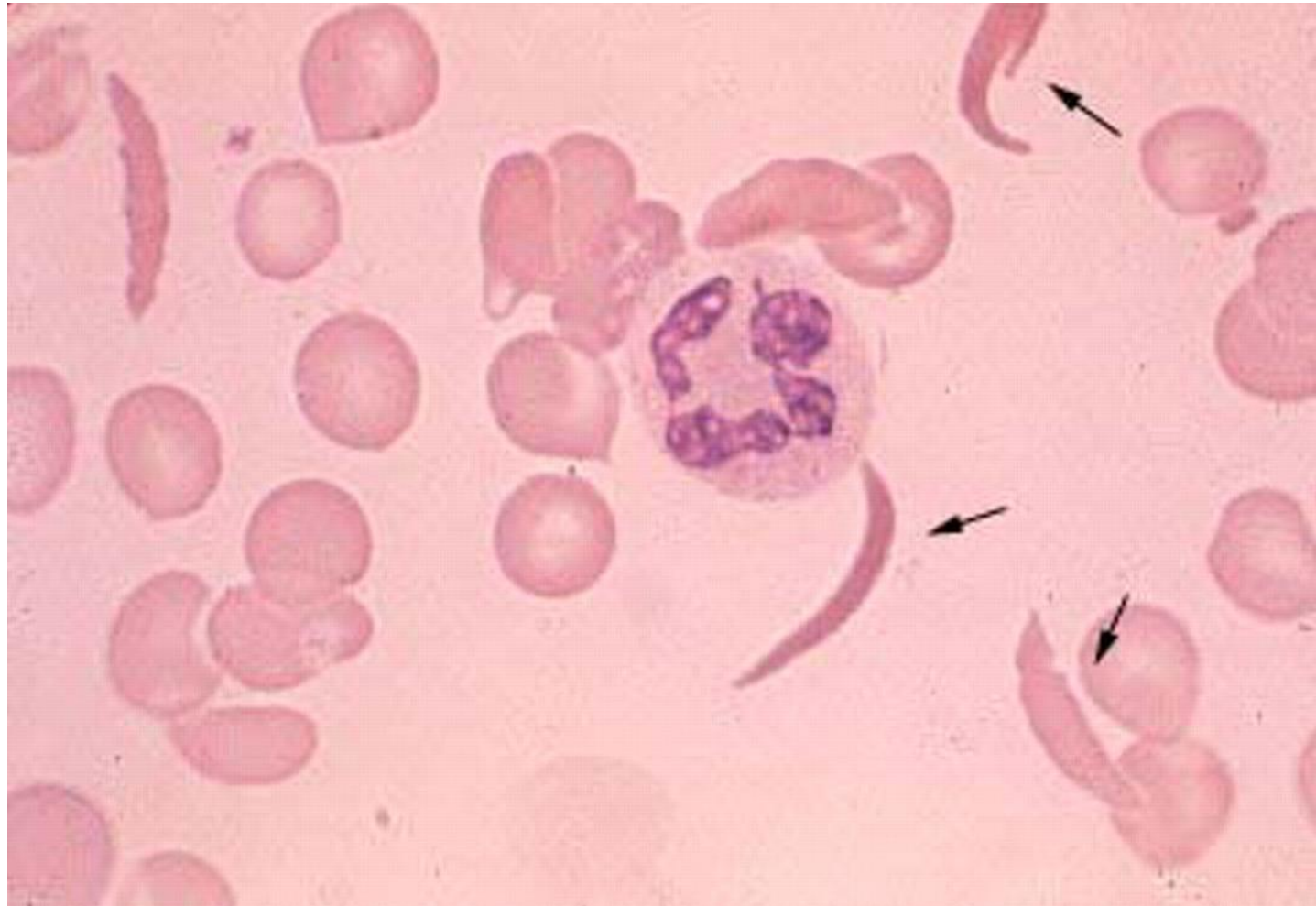
Anemia sideropenica: microciti



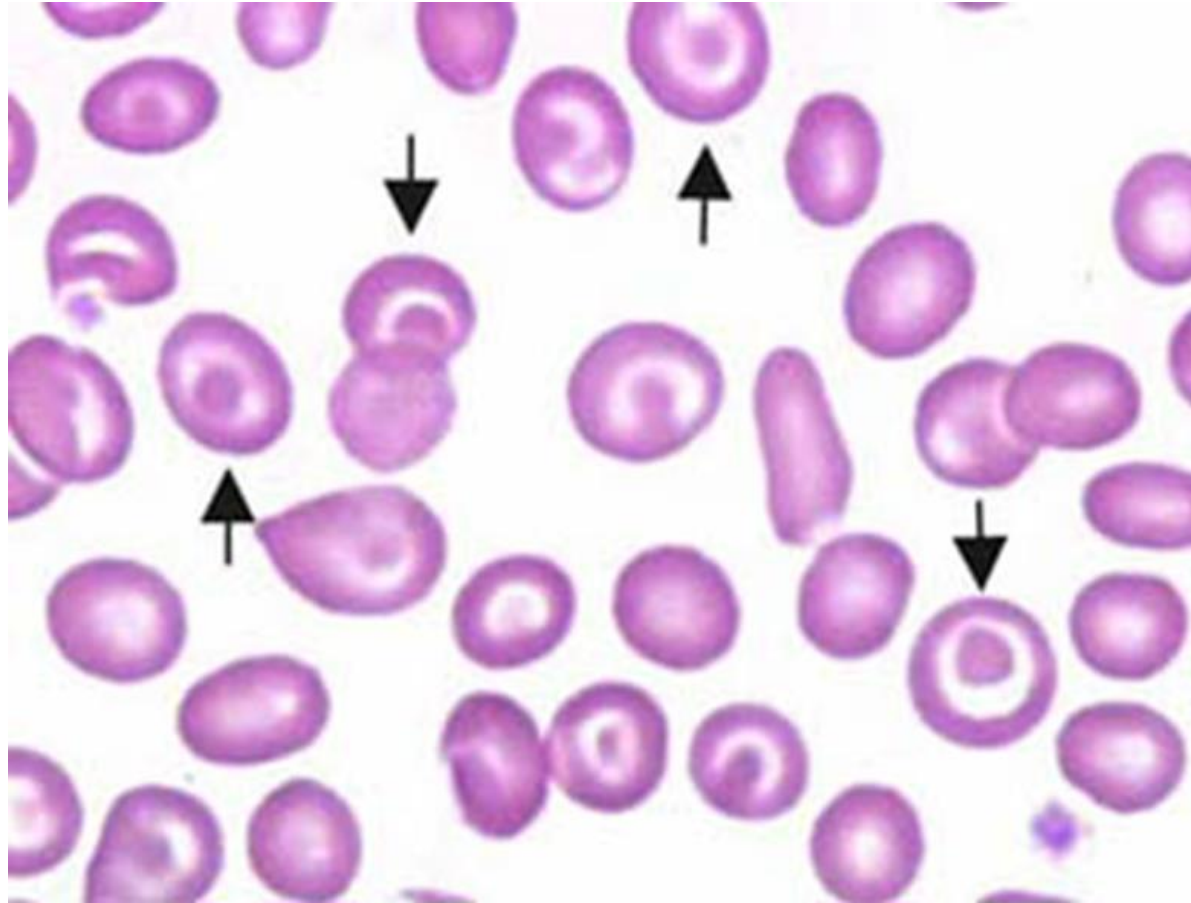
Macrocyti



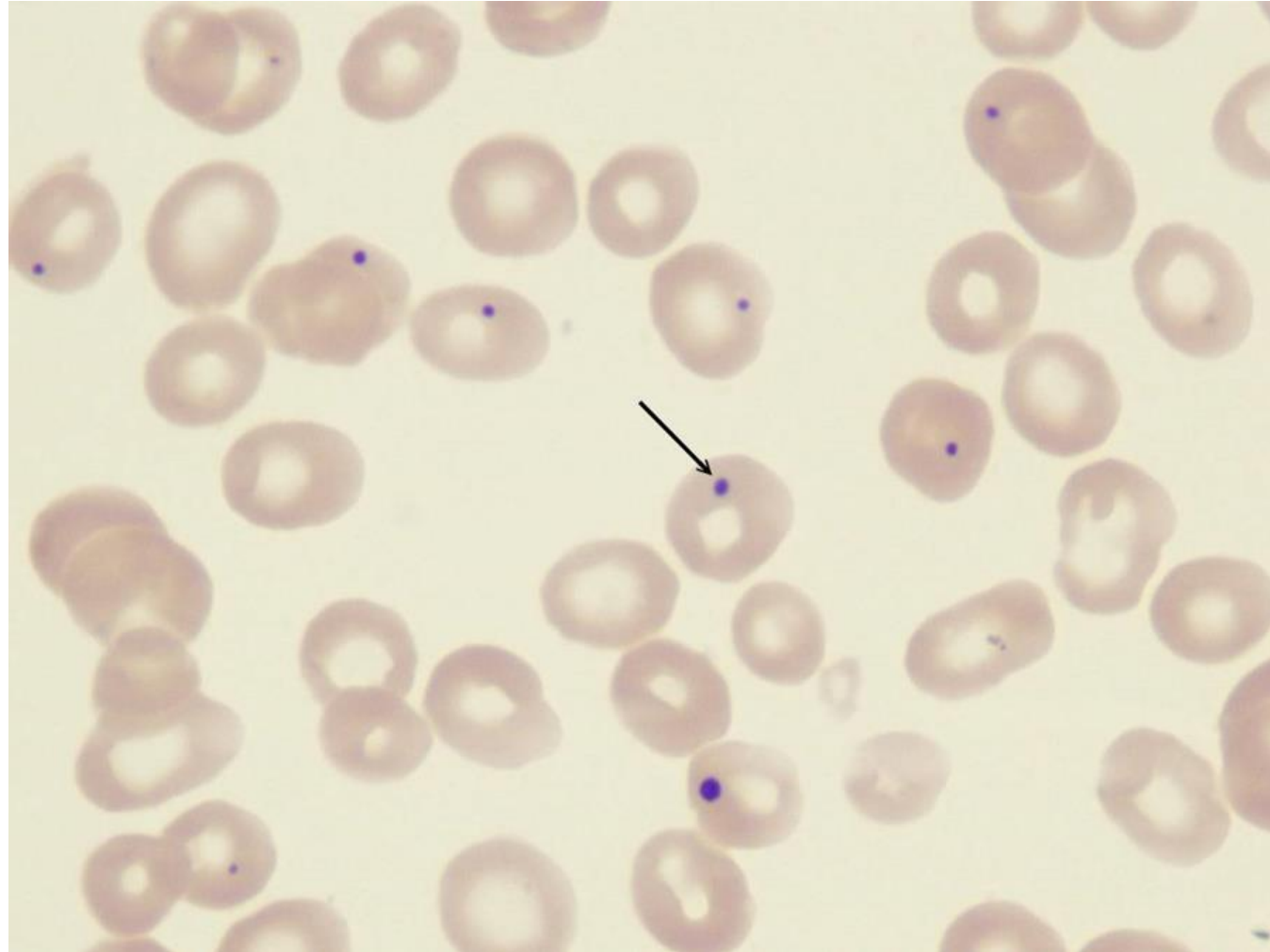
Anemia a cellule falciformi



Target cells



Corpi di Howell-Jolly



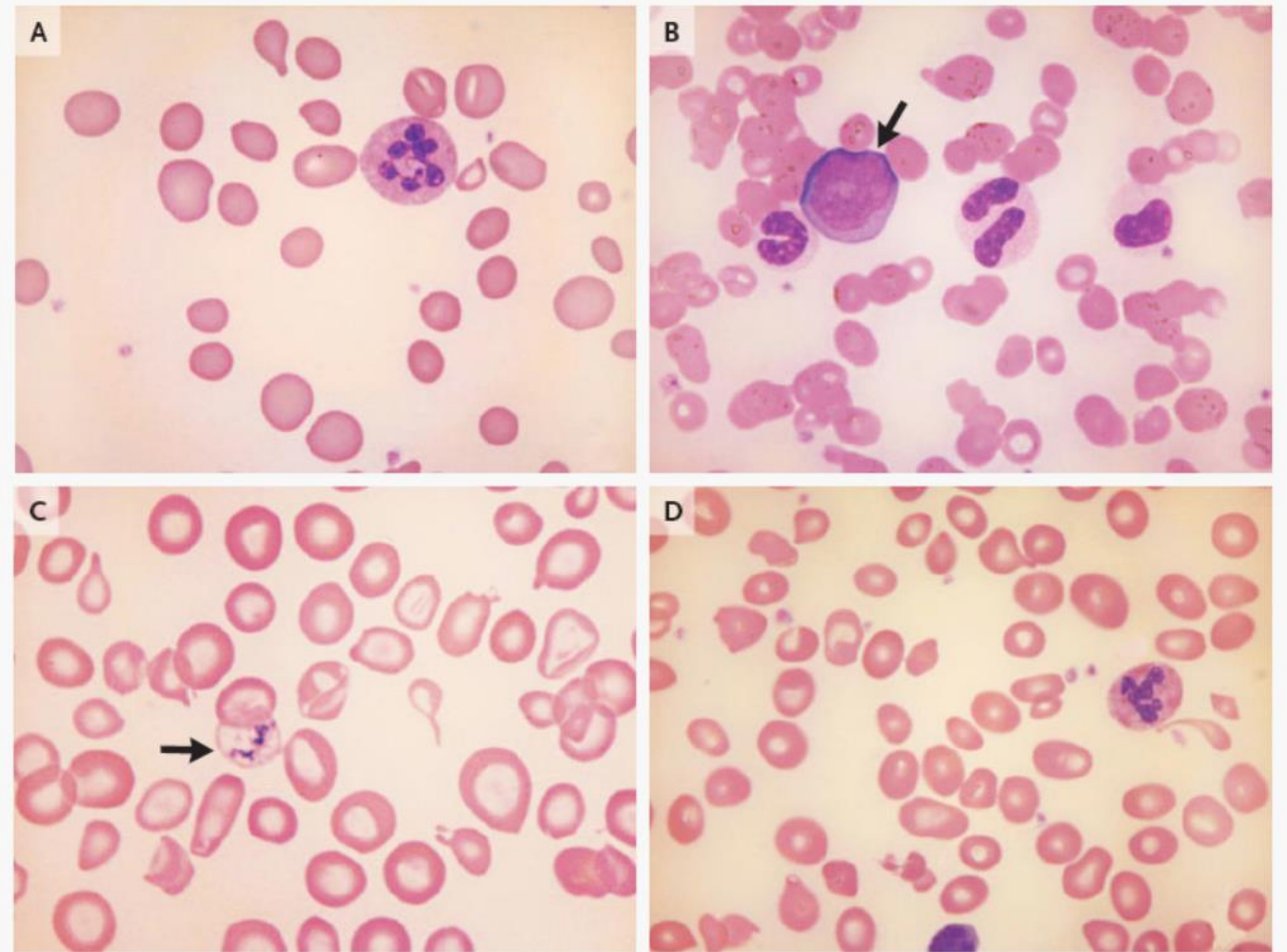


Figure 3. Red-Cell Changes in Various Types of Macrocytic Anemia.

Pernicious anemia is shown in the blood smear in Panel A, with anisocytosis, macrocytosis, and a hypersegmented neutrophil. Panel B shows myelodysplastic syndrome, with a blast cell (arrow) and two neutrophils that have hypolobulated nuclei, one of which is binucleated and the other hypogranular. Panel C shows myelodysplastic syndrome with anisocytosis, poikilocytosis, macrocytes, stomatocytes, and an erythrocyte with prominent Pappenheimer bodies (arrow); the smear is also dimorphic, showing both well-hemoglobinized macrocytes and hypochromic microcytes. Panel D depicts type 1 congenital dyserythropoietic anemia, with anisocytosis, poikilocytosis, and some macrocytes. All specimens were stained with May–Grünwald–Giemsa stain.

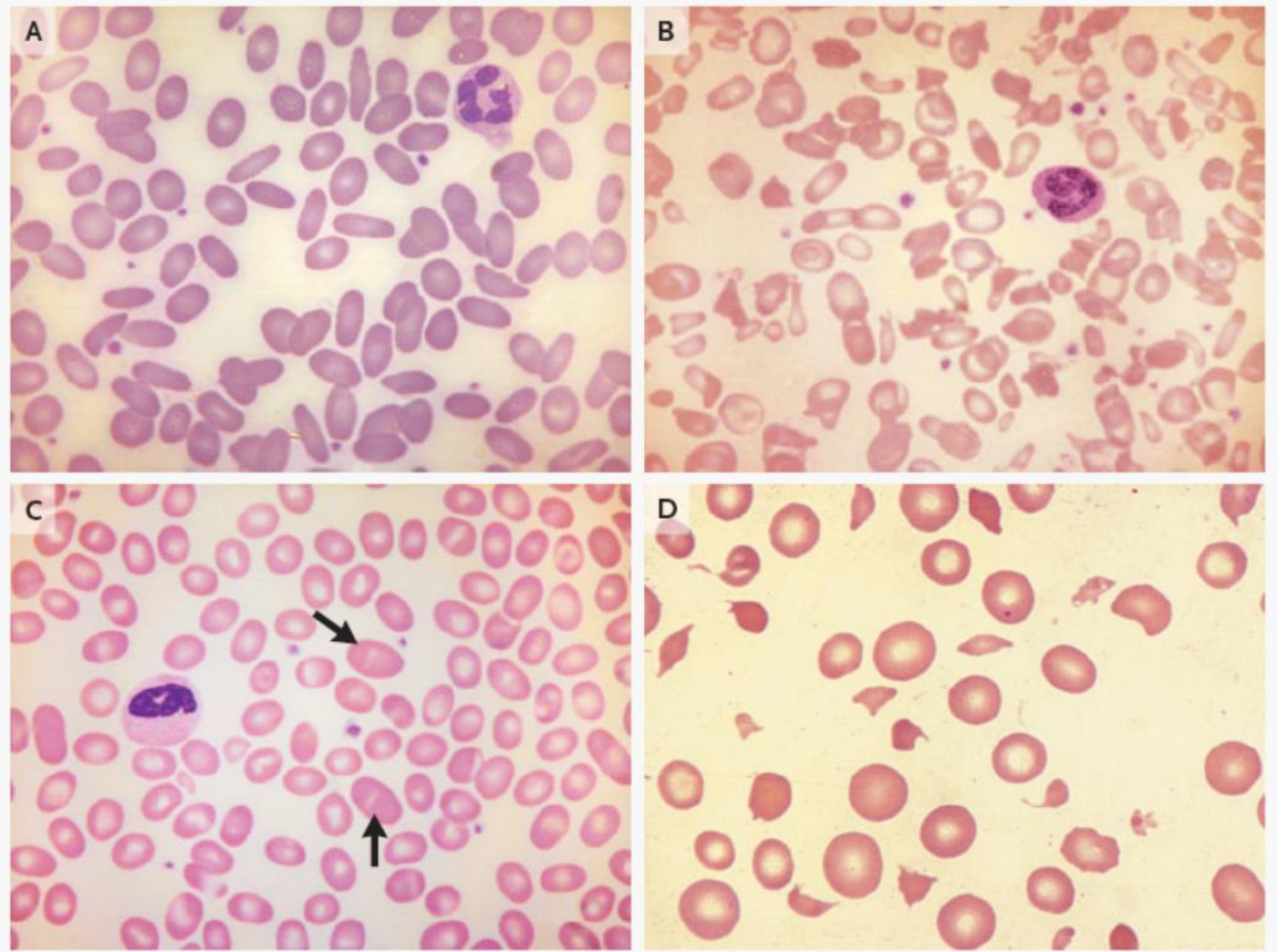


Figure 1. Hemolytic Anemias, Characterized by Different Types of Poikilocytes.

In Panel A, the blood smear shows hereditary elliptocytosis, with numerous elliptocytes and smaller numbers of ovalocytes. Panel B shows hereditary pyropoikilocytosis; there is striking poikilocytosis, with elliptocytes, ovalocytes, and fragments. In Panel C, Southeast Asian ovalocytosis shows moderate poikilocytosis, with the poikilocytes including several macro-ovalocytes (arrows). Panel D shows microangiopathic hemolytic anemia resulting from cyclosporine therapy, with numerous red-cell fragments. All specimens were stained with May-Grünwald-Giemsa stain.