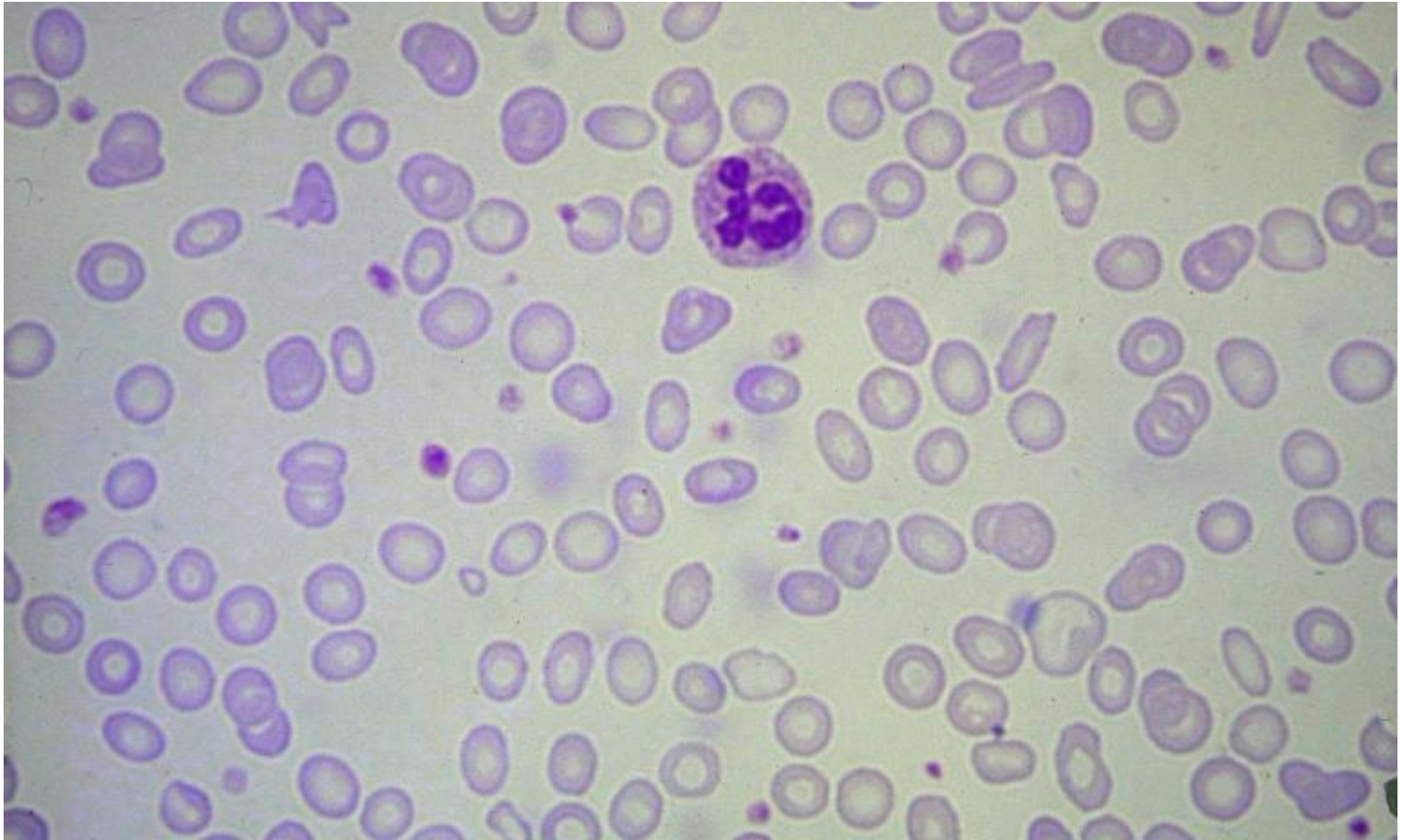


Morphology in Anemia

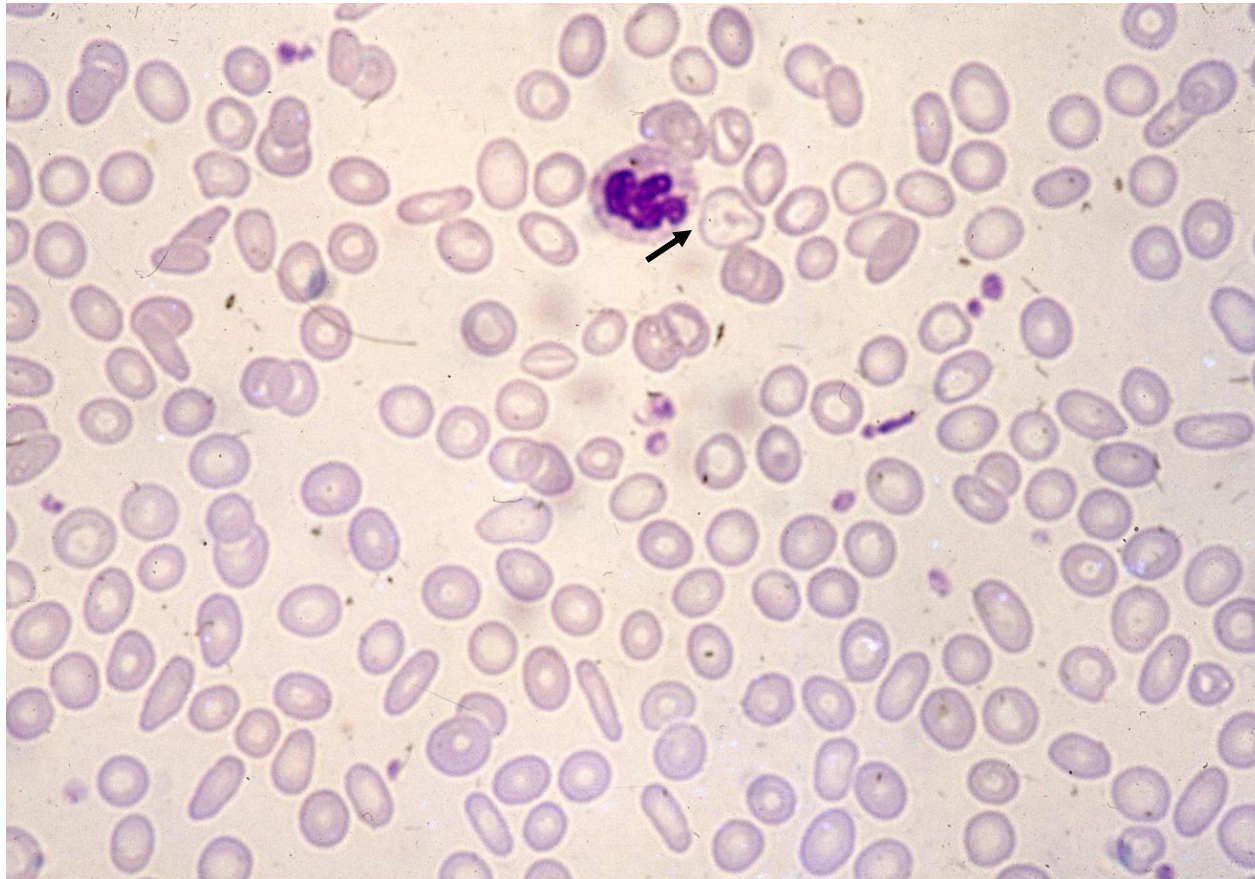
Prof Peretz Resnitzky

Iron deficiency anemia



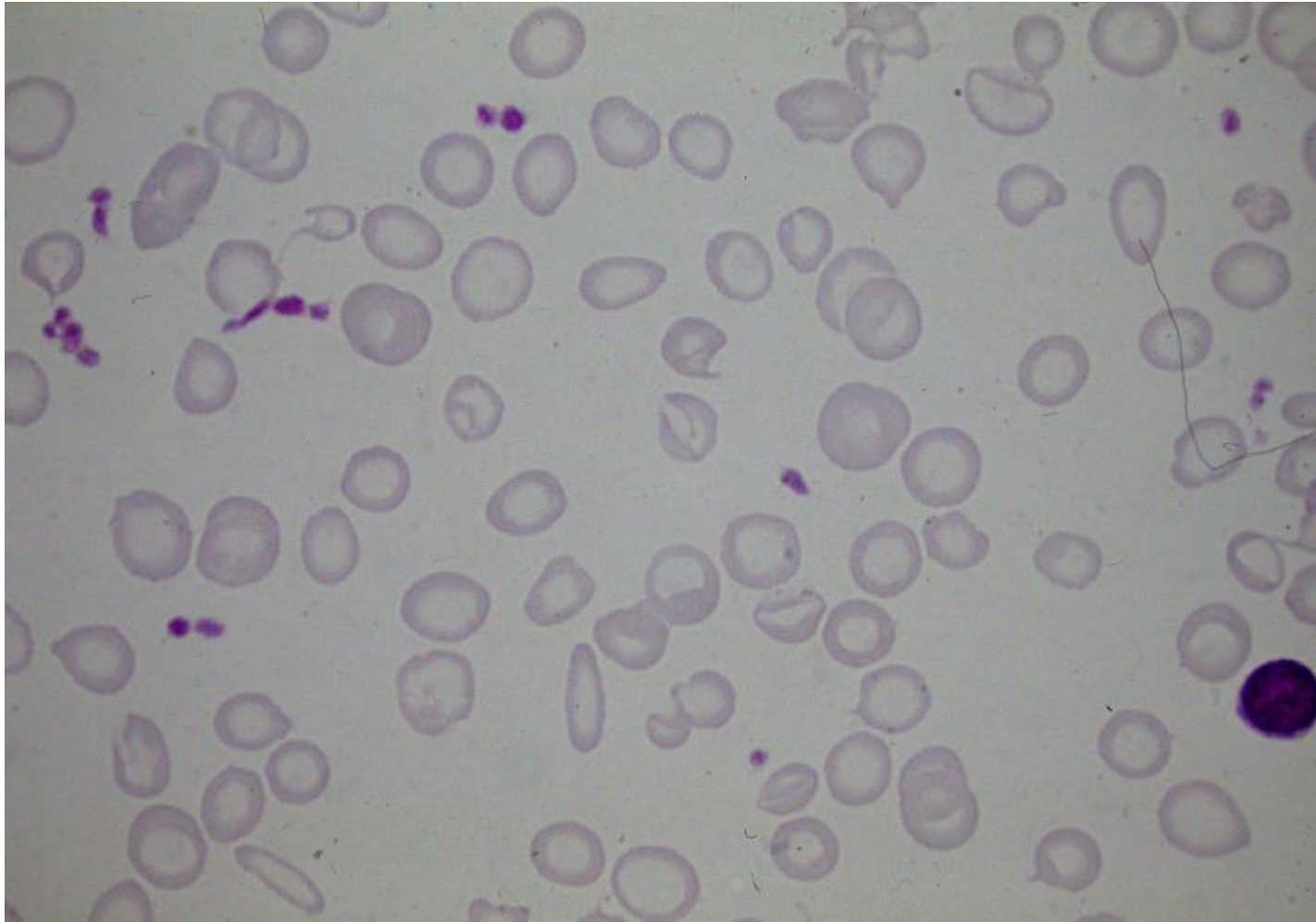
Anisocytosis, poikilocytosis, ovalocytes, hypochromia

Iron deficiency anemia



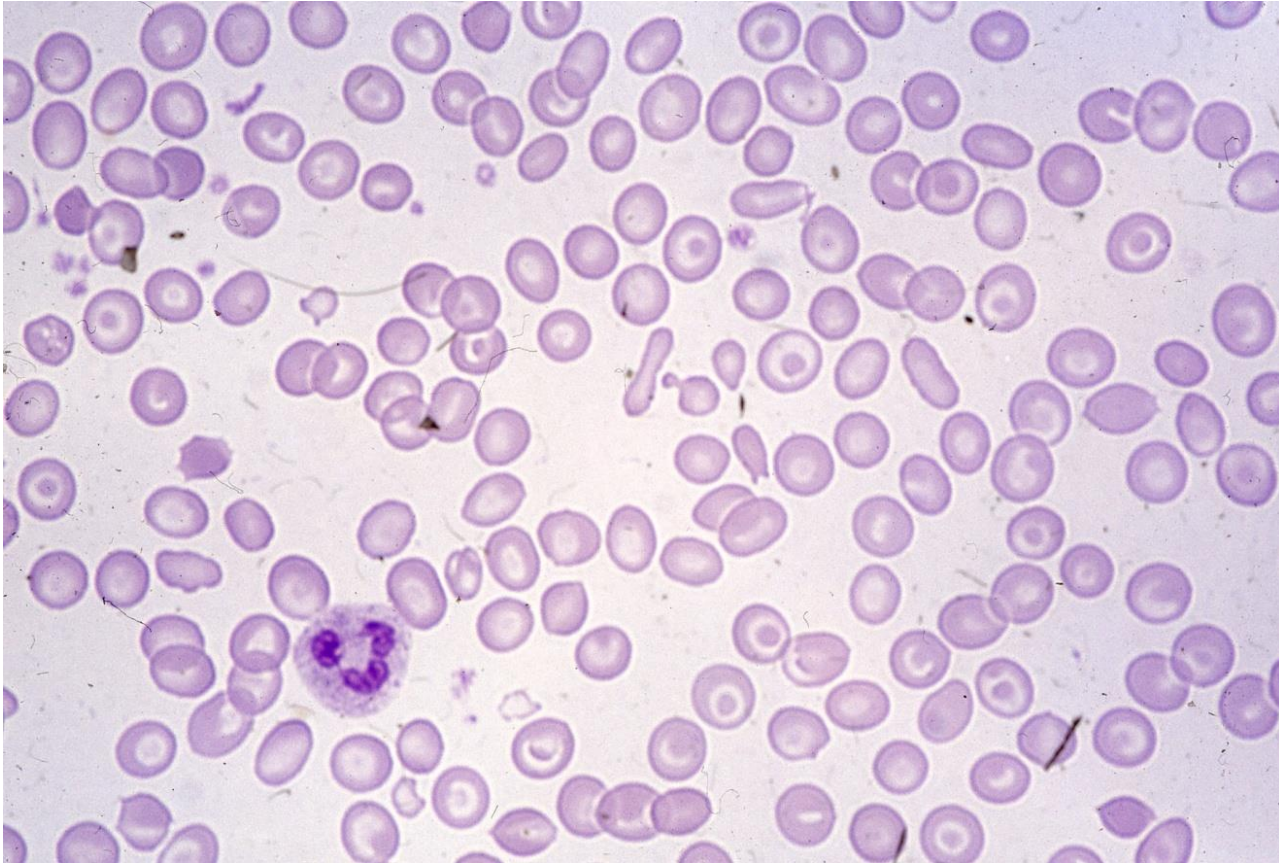
Anisocytosis, ovalocytes, hypochromia,
target cell (arrow)

Iron deficiency anemia



Anisocytosis, poikilocytosis, hypochromia, ovalocytes,

Thalassemia minor



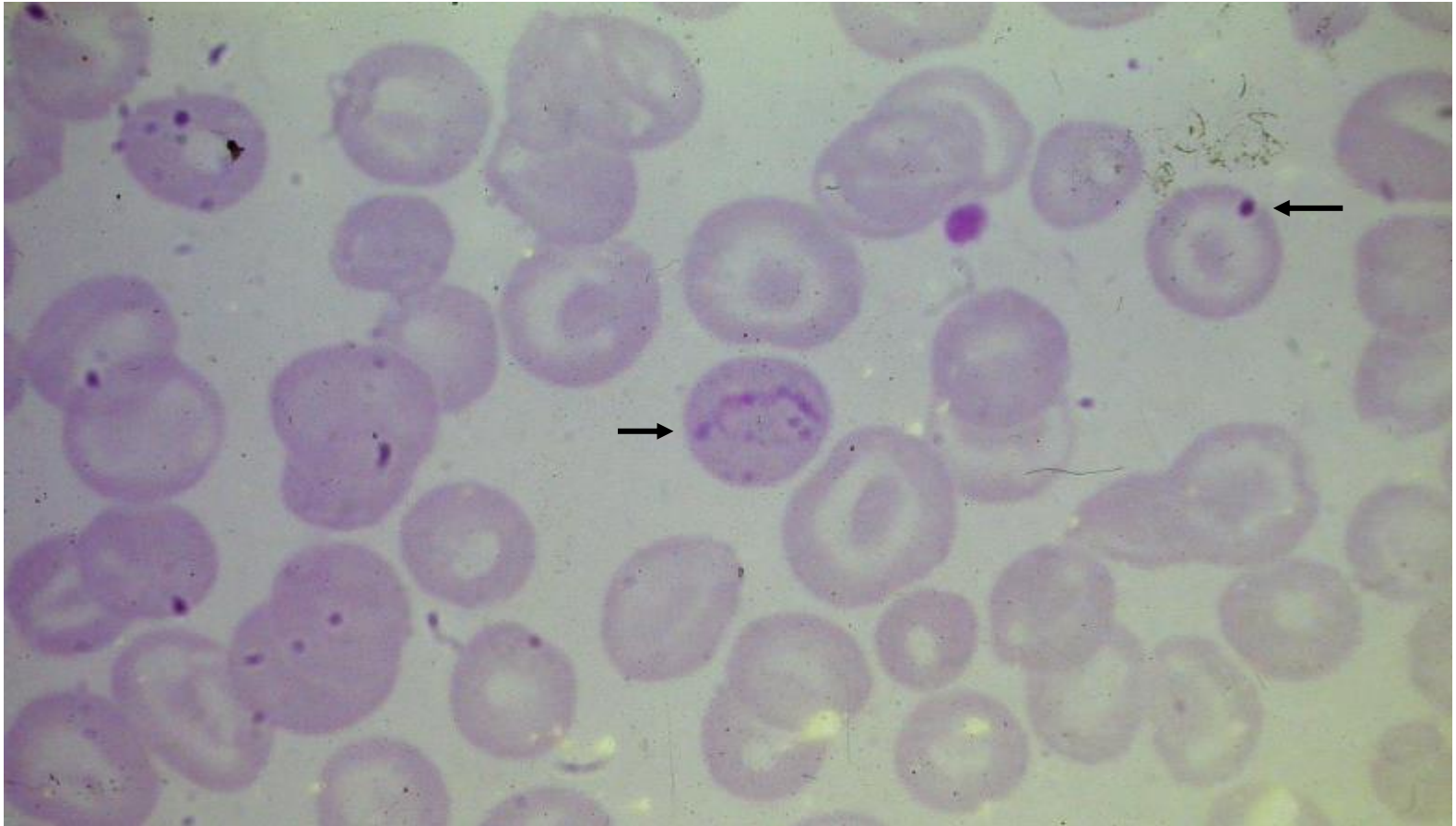
Target cells' ovalocytes

Thalassemia minor



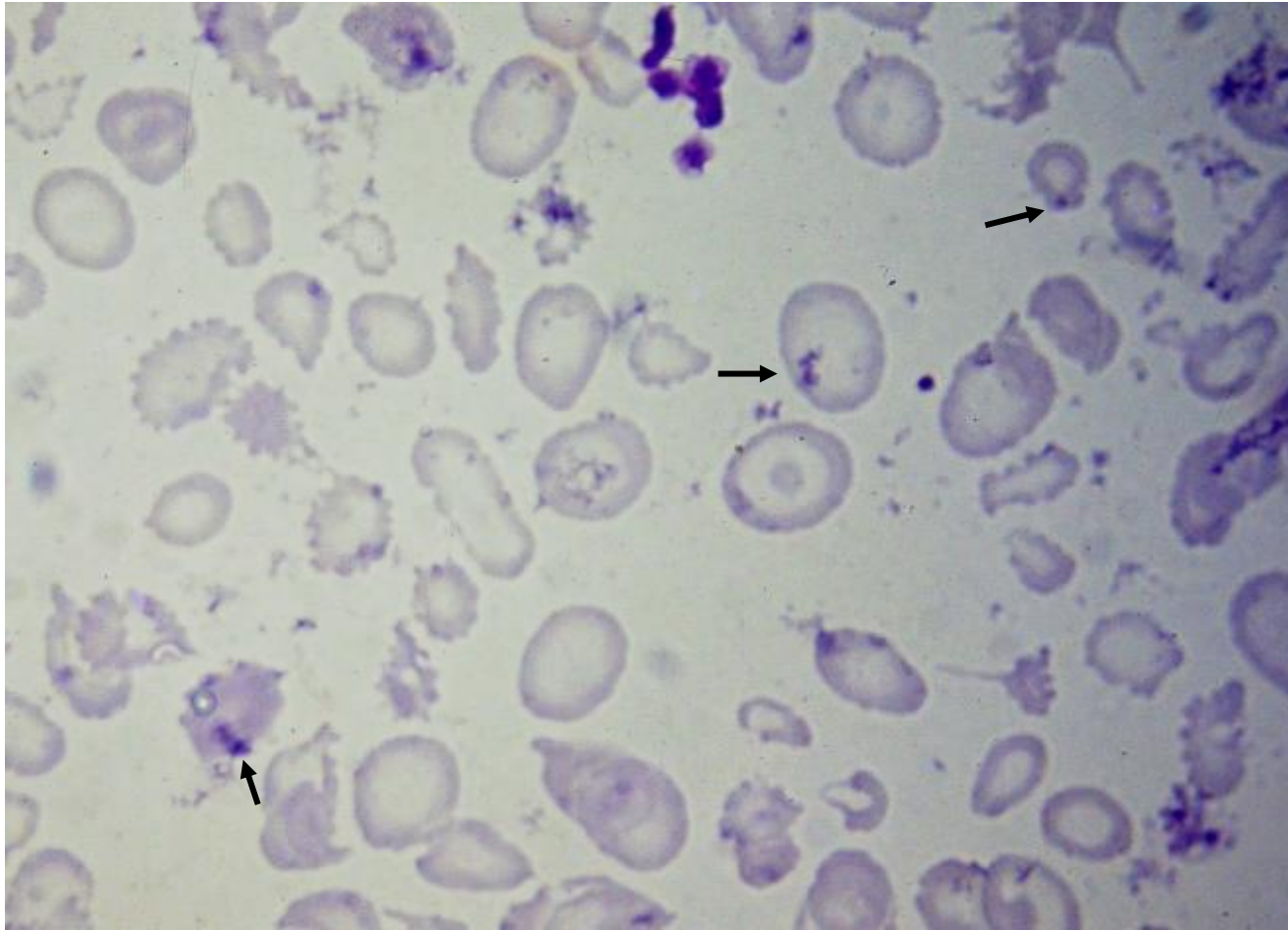
Hypochromia, target cells, mycrocytes, Howell Jolly body (arrow)

Thalassemia minor



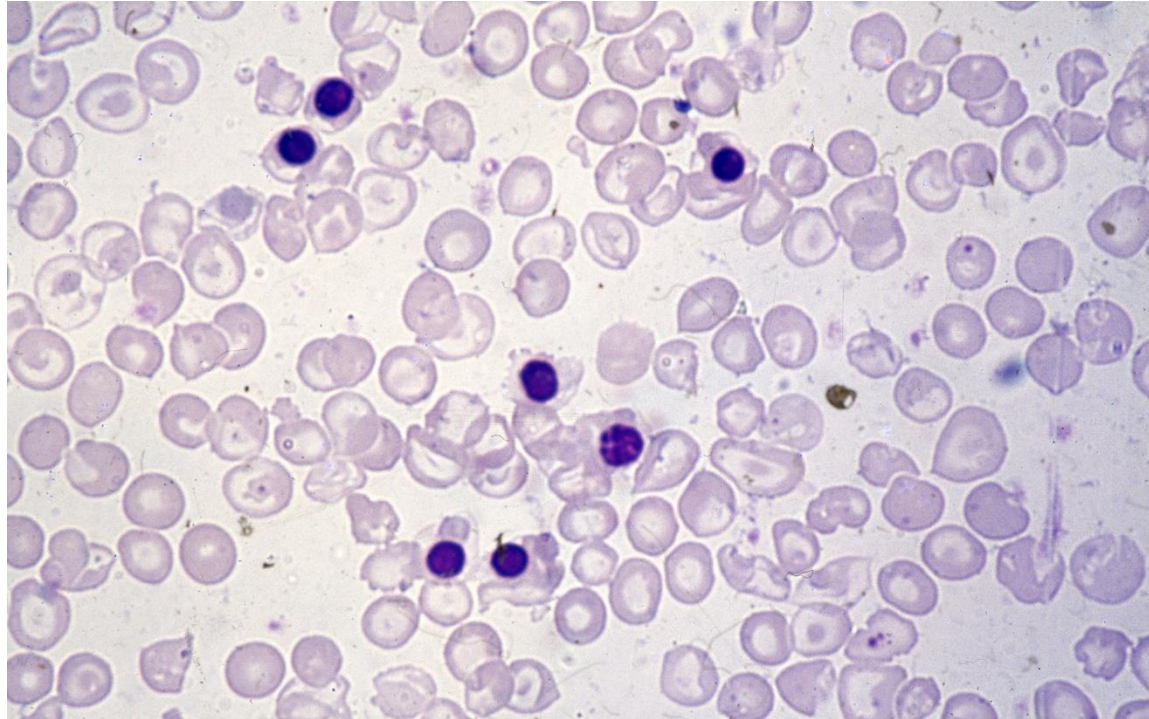
Hypochromia, target cells, Cabot ring & basophilic stippling (arrow),
Howell-Jolly body (arrow)

Thalassemia major



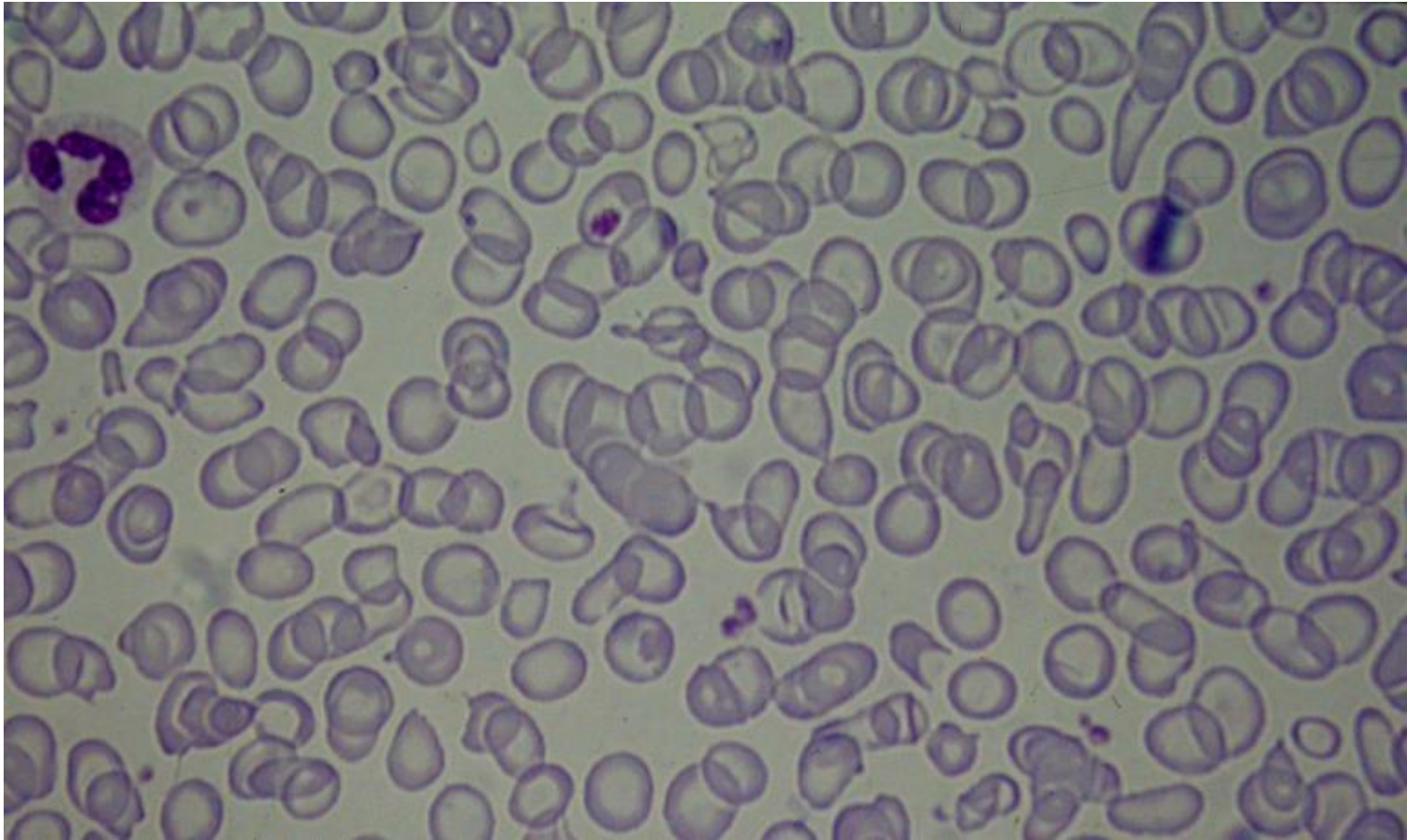
Aniso-poikilocytosis, hypochromia, target cells, Howell-Jolly bodies (arrows)

Thalassemia major after Sx



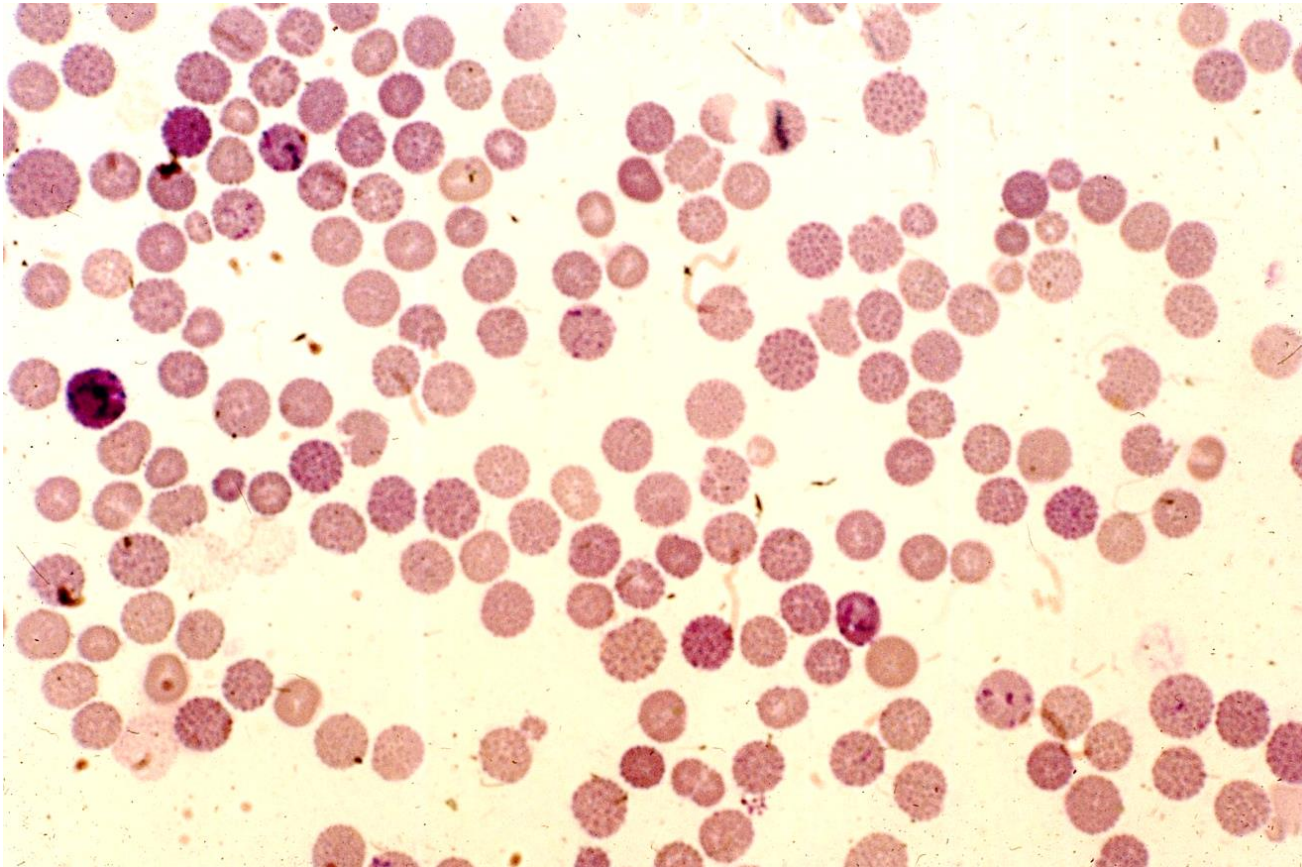
Nucleated rbc, target cells Howell Jolly bodies

Hemoglobin H disease



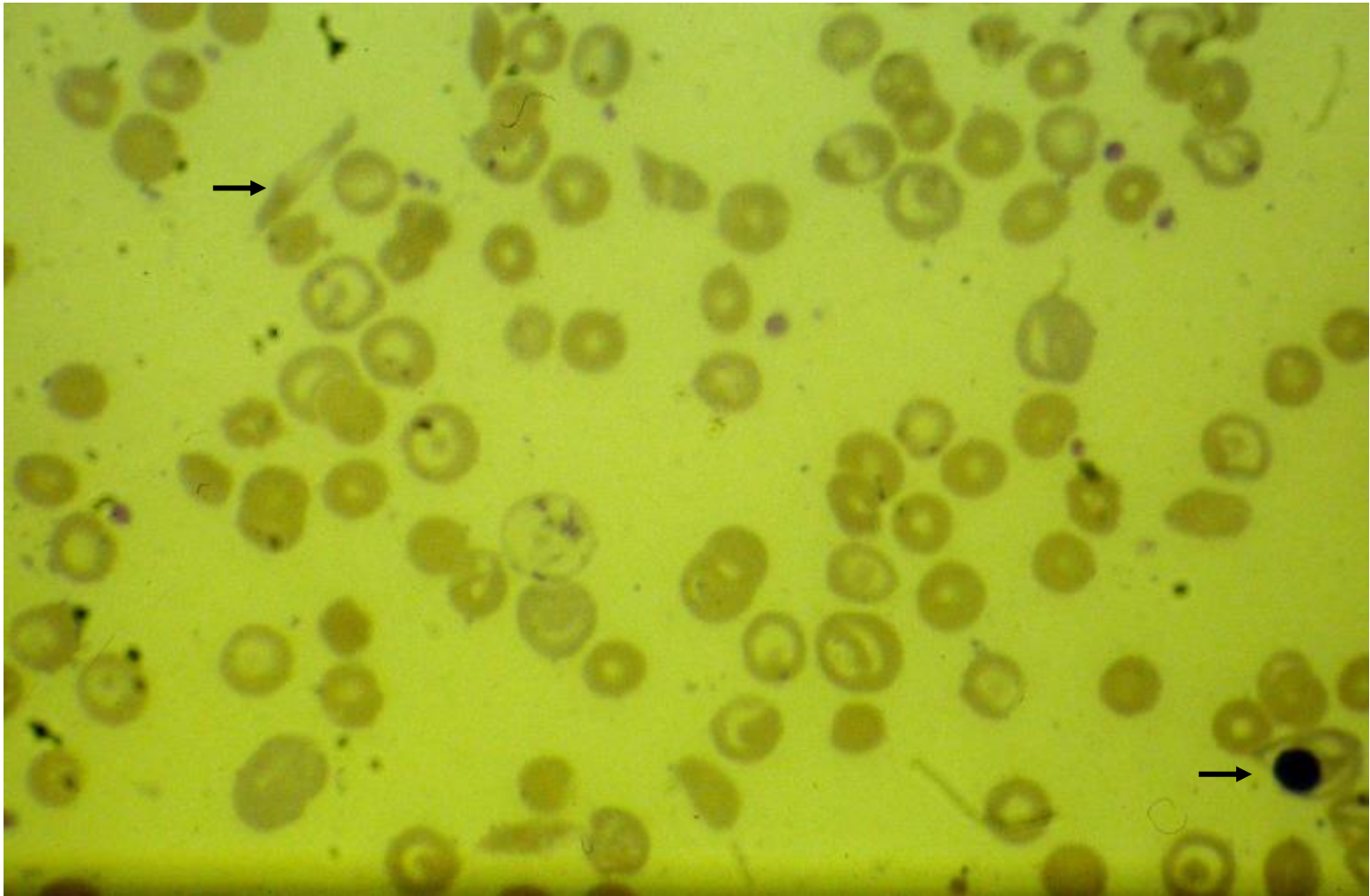
Hypochromia , anisopoikilocytosis, ovalocytes

Hemoglobin H disease



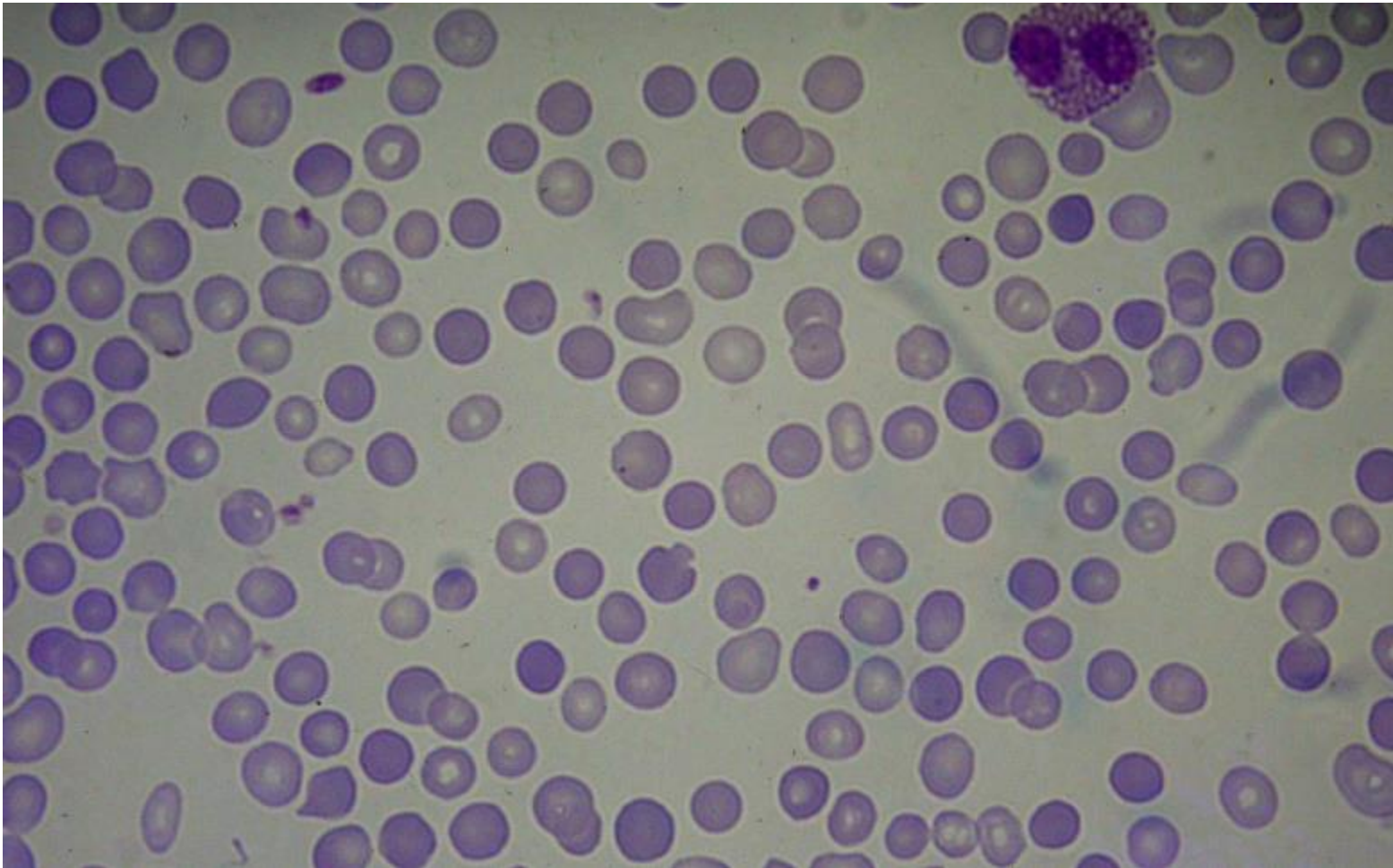
Brilliant cresyl blue staining: inclusions in rbc
(reticulocytes are also stained)

Sickle cell-Thalassemia



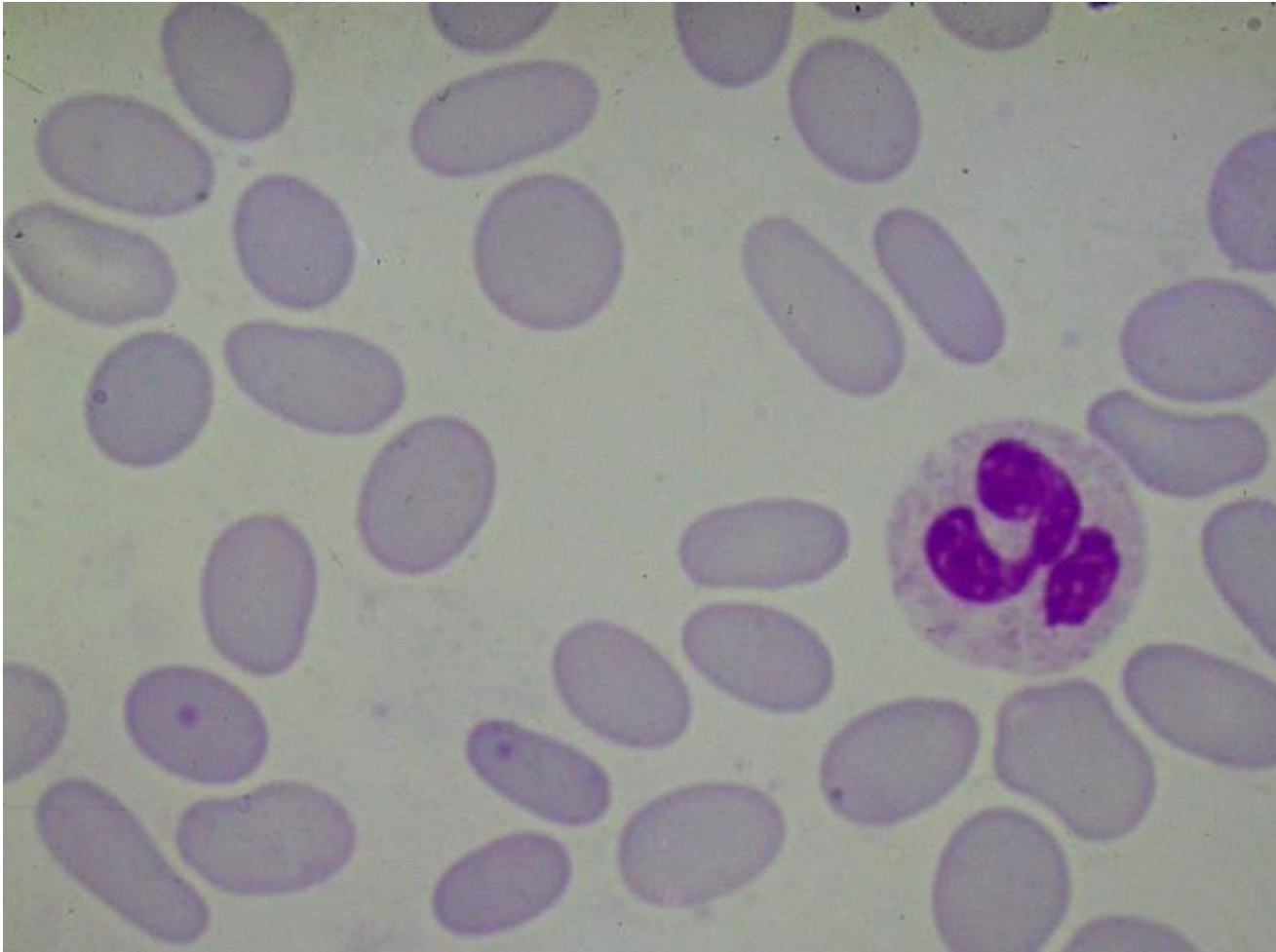
Hypochromia, target cells, sickle cell (arrow), nucleated rbc (arrow)

Congenital spherocytosis



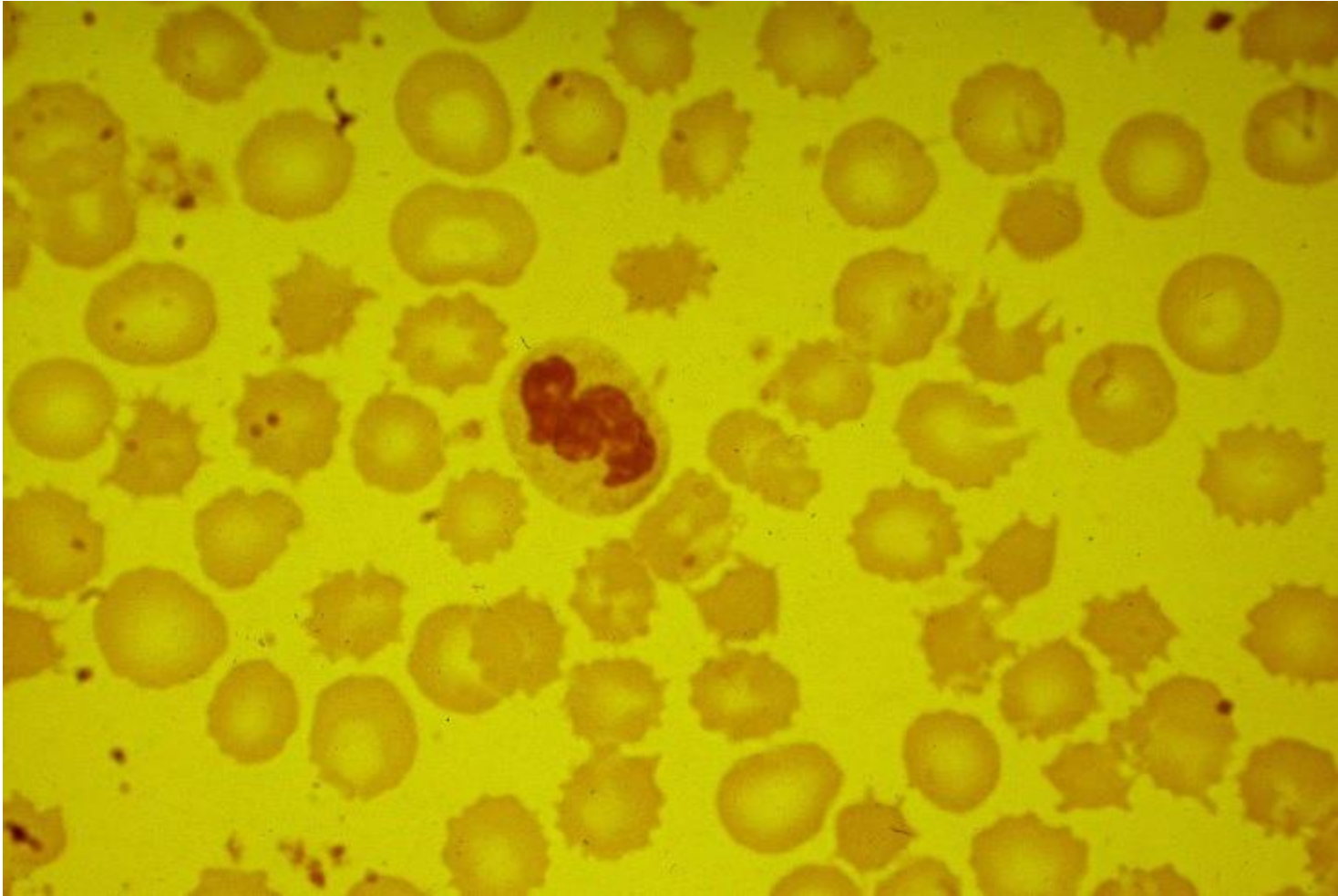
Spherocytes, and micro-spherocytes

Congenital elliptocytosis (ovalocytosis)

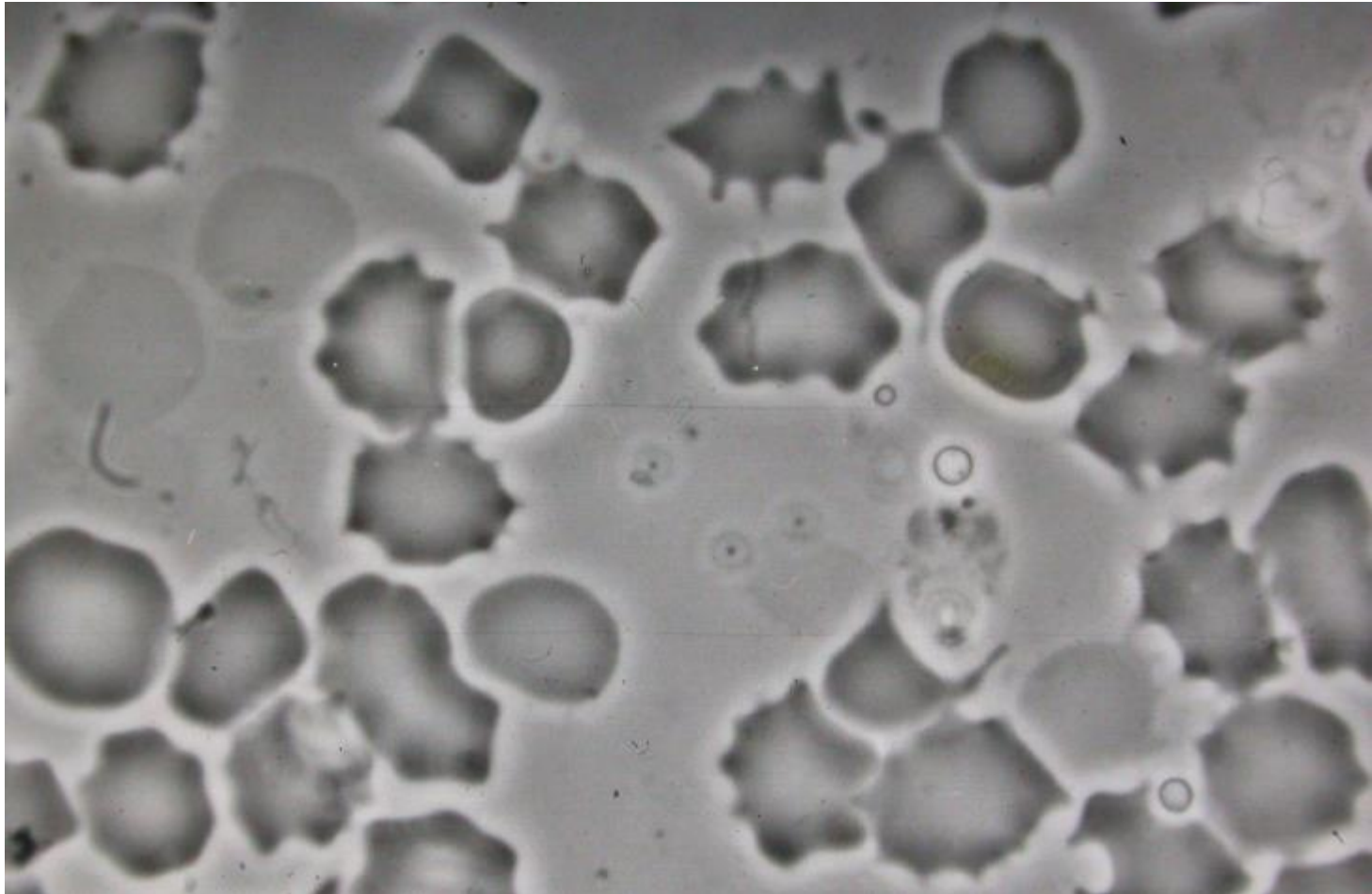


Up to 15% elliptocytes can be found in: IDA, Megaloblastic anemia, Thalassemia

Acanthocytosis

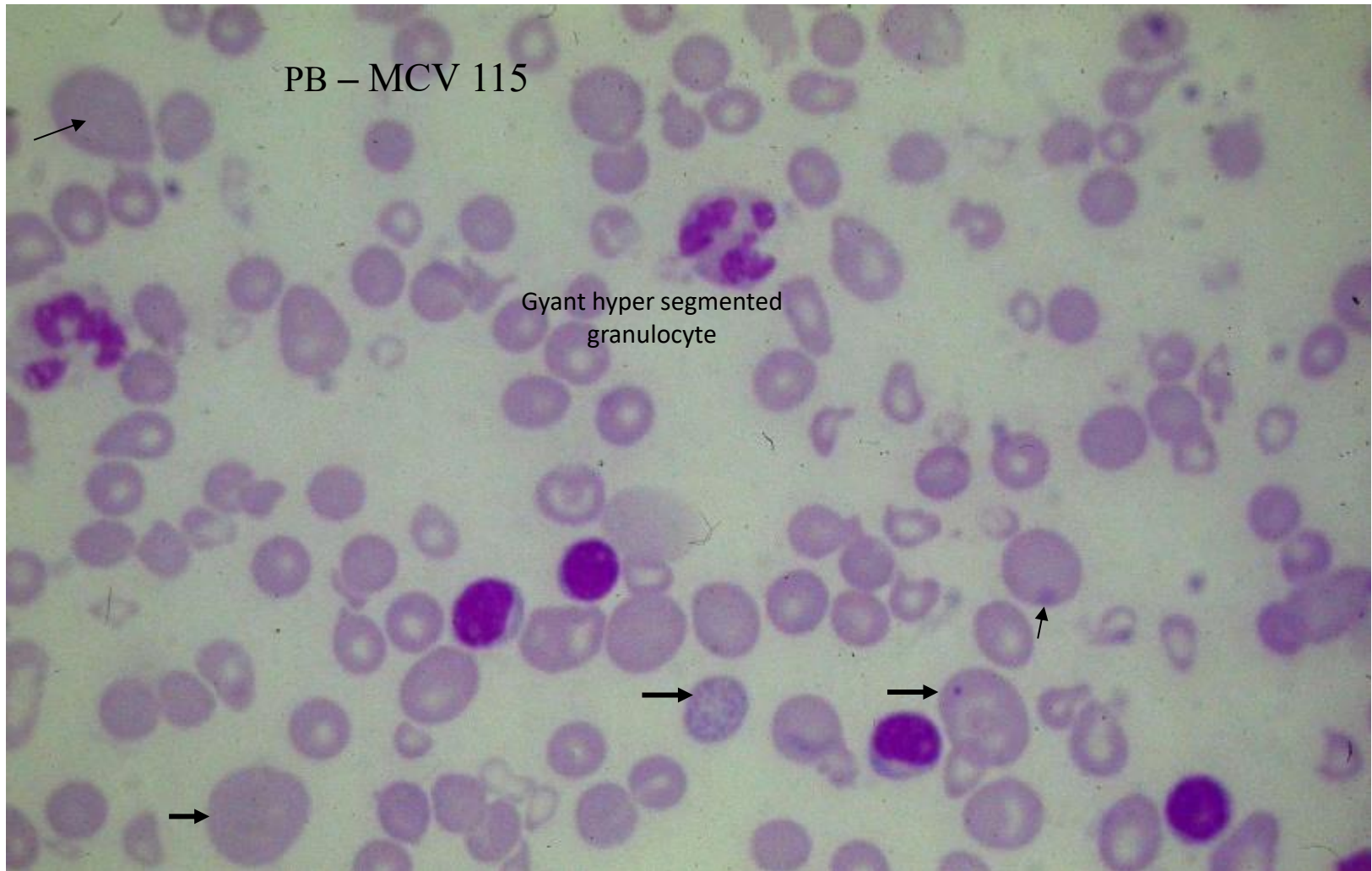


Acanthocytosis

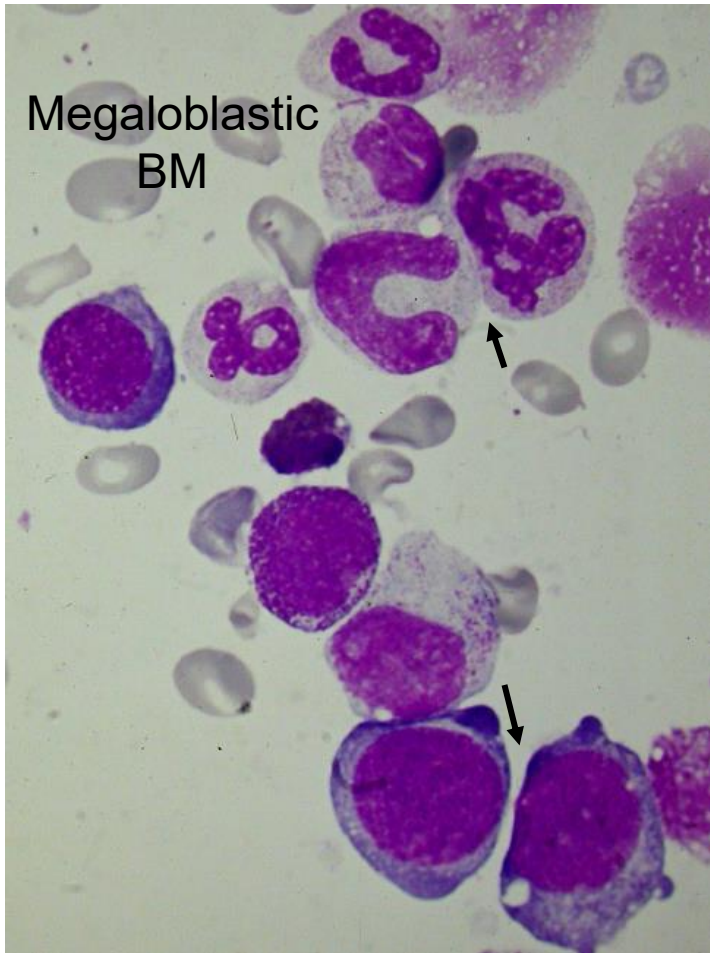


Phase contrast microscopy

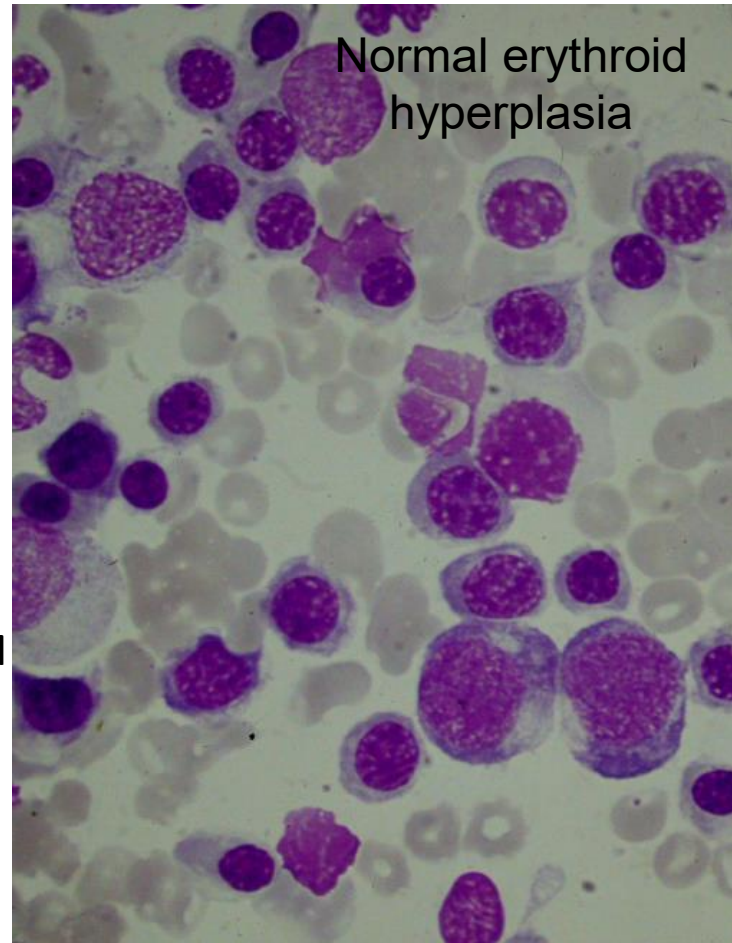
PB in Megaloblastic anemia



Macrocytes, Howell Jolly bodies, basophilic stippling (arrows). Marked aniso-poikilocytosis

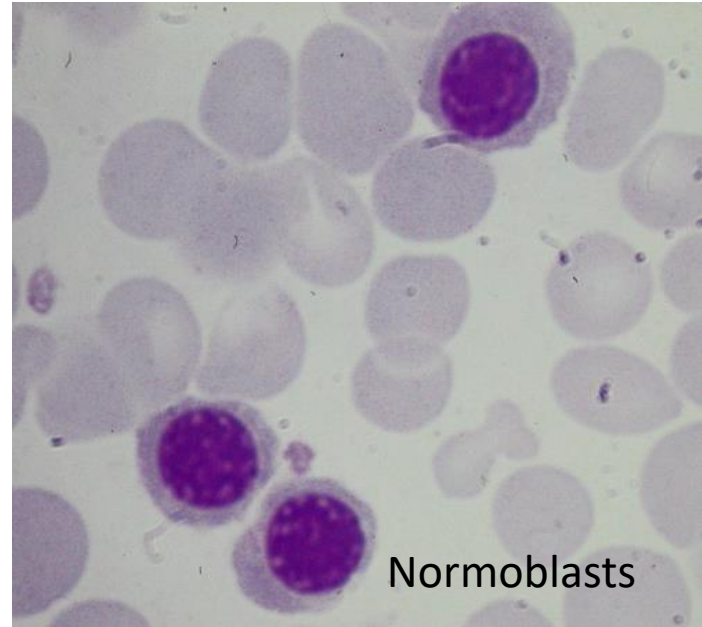


BM

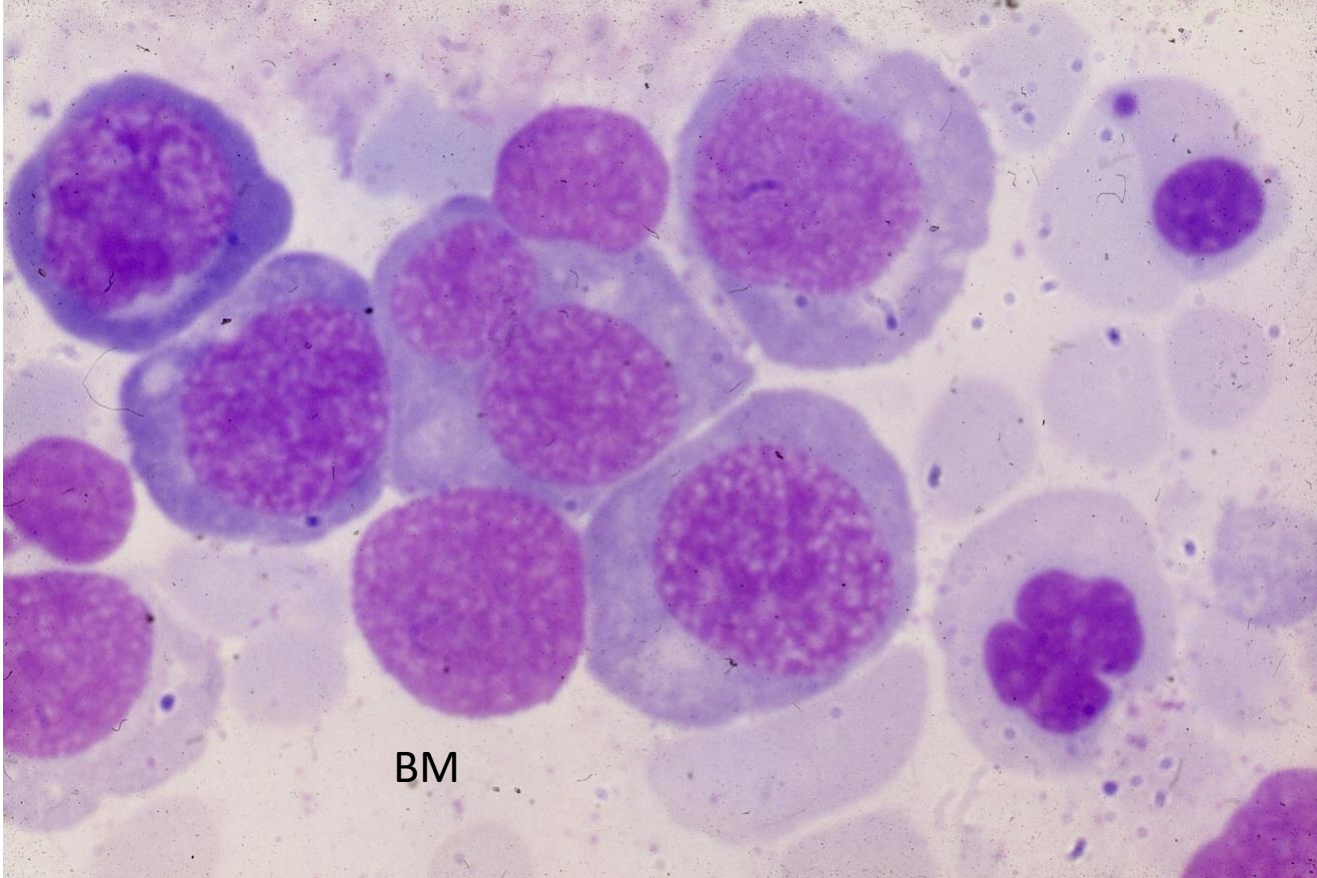


Gyant stab, gyant
hypersegmented neutrophil
and two pro-megaloblasts

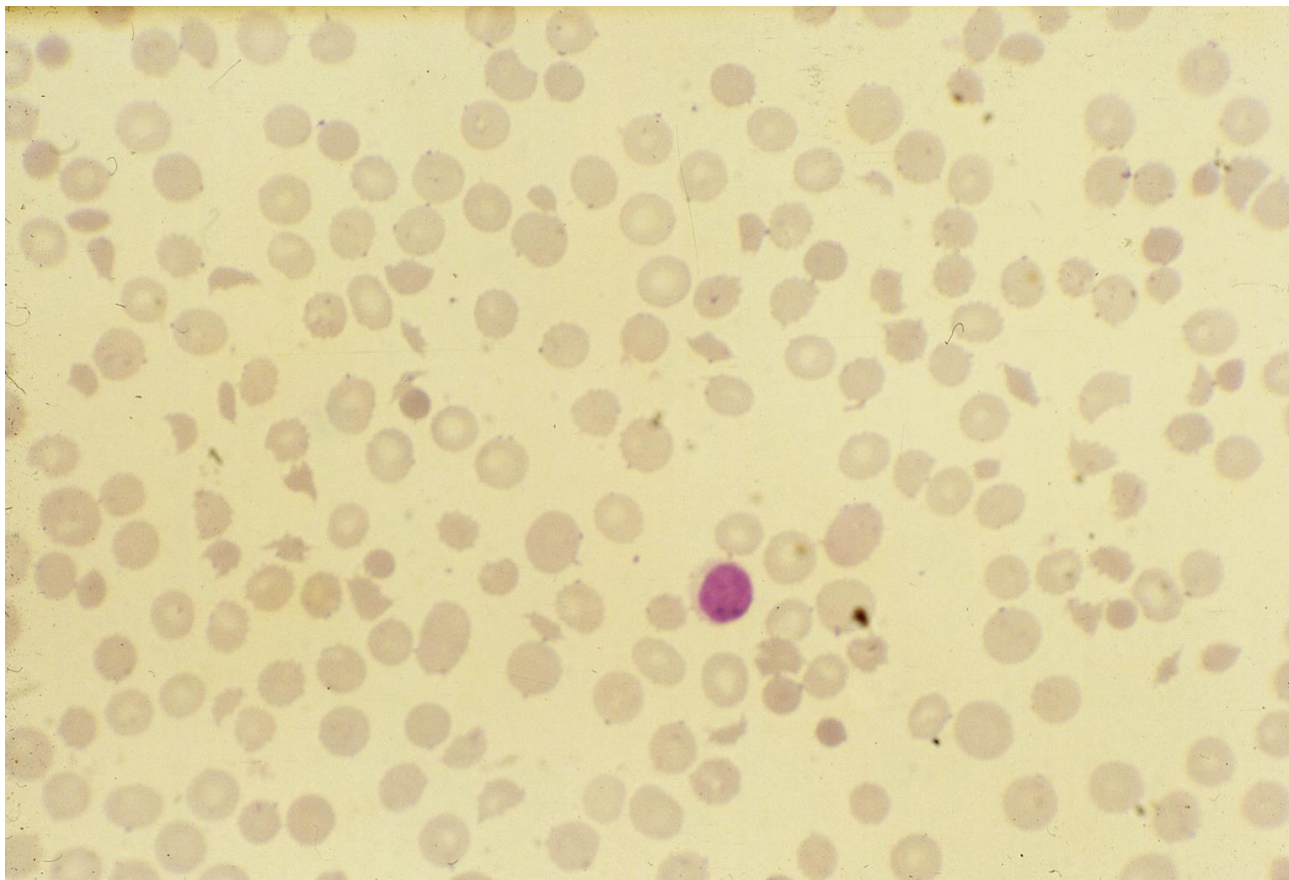
Megaloblasts vs normoblasts



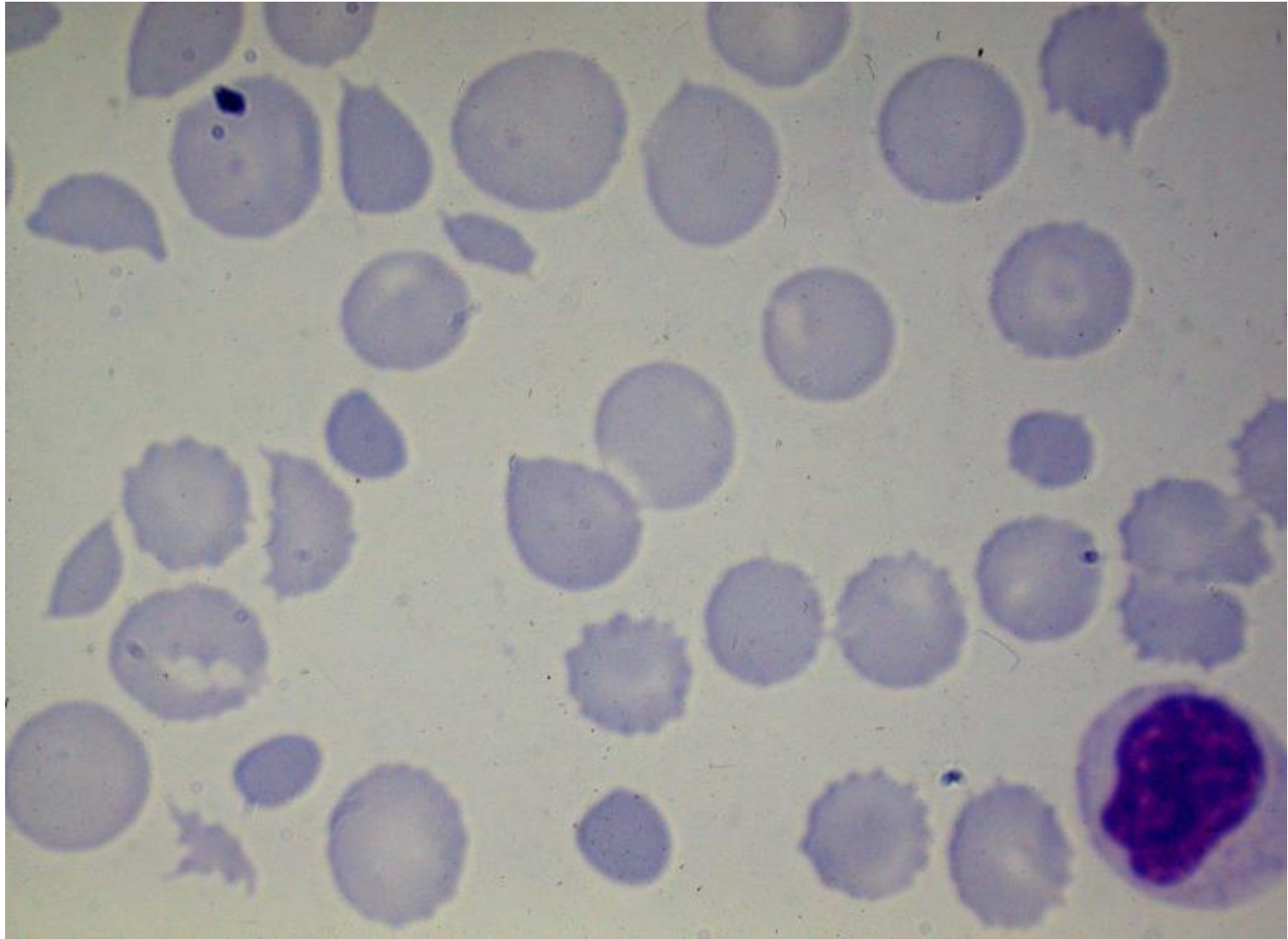
Megaloblasts in different stages of maturation



Microangiopathic hemolytic anemia

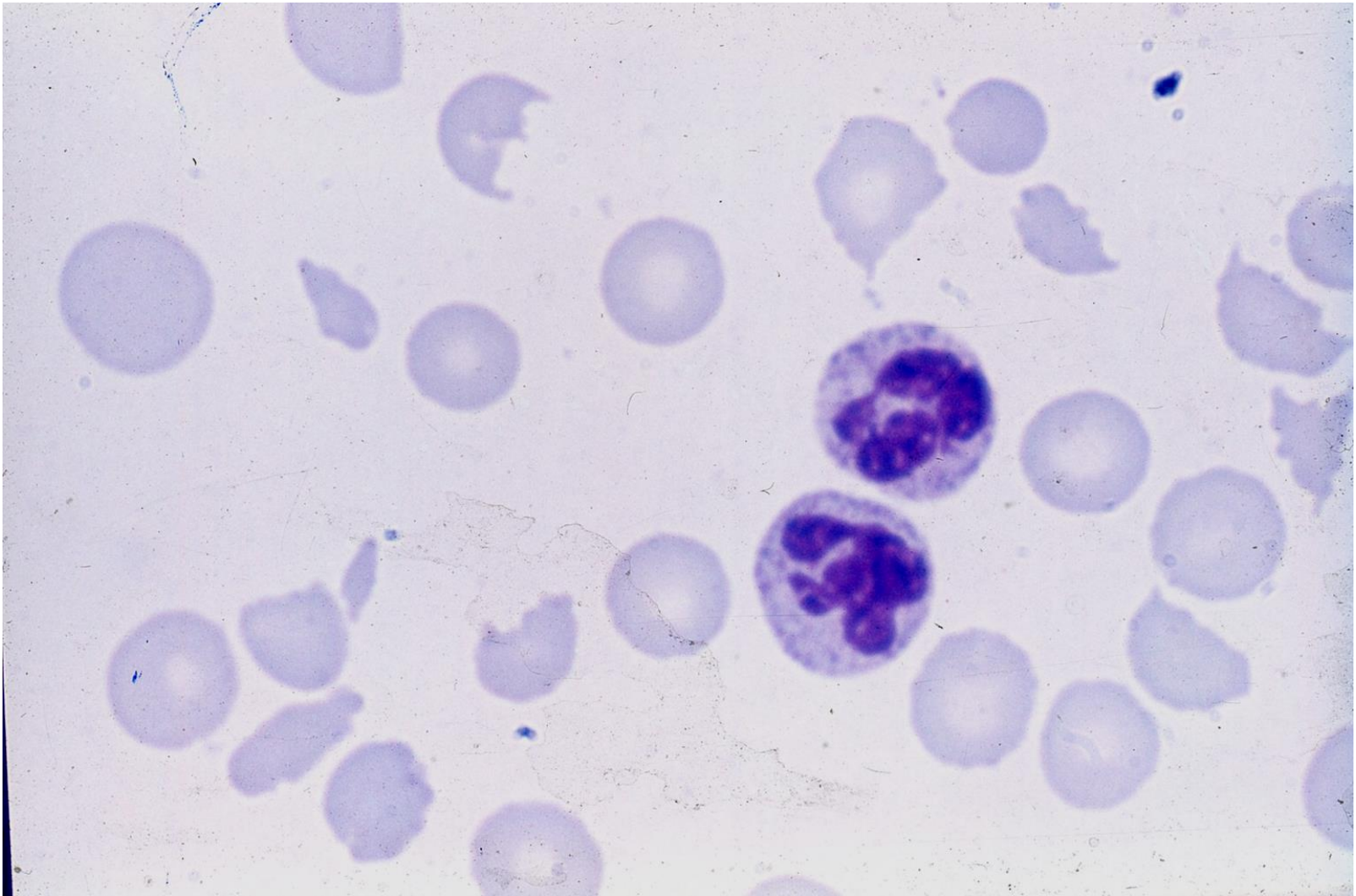


Microangiopathic hemolytic anemia

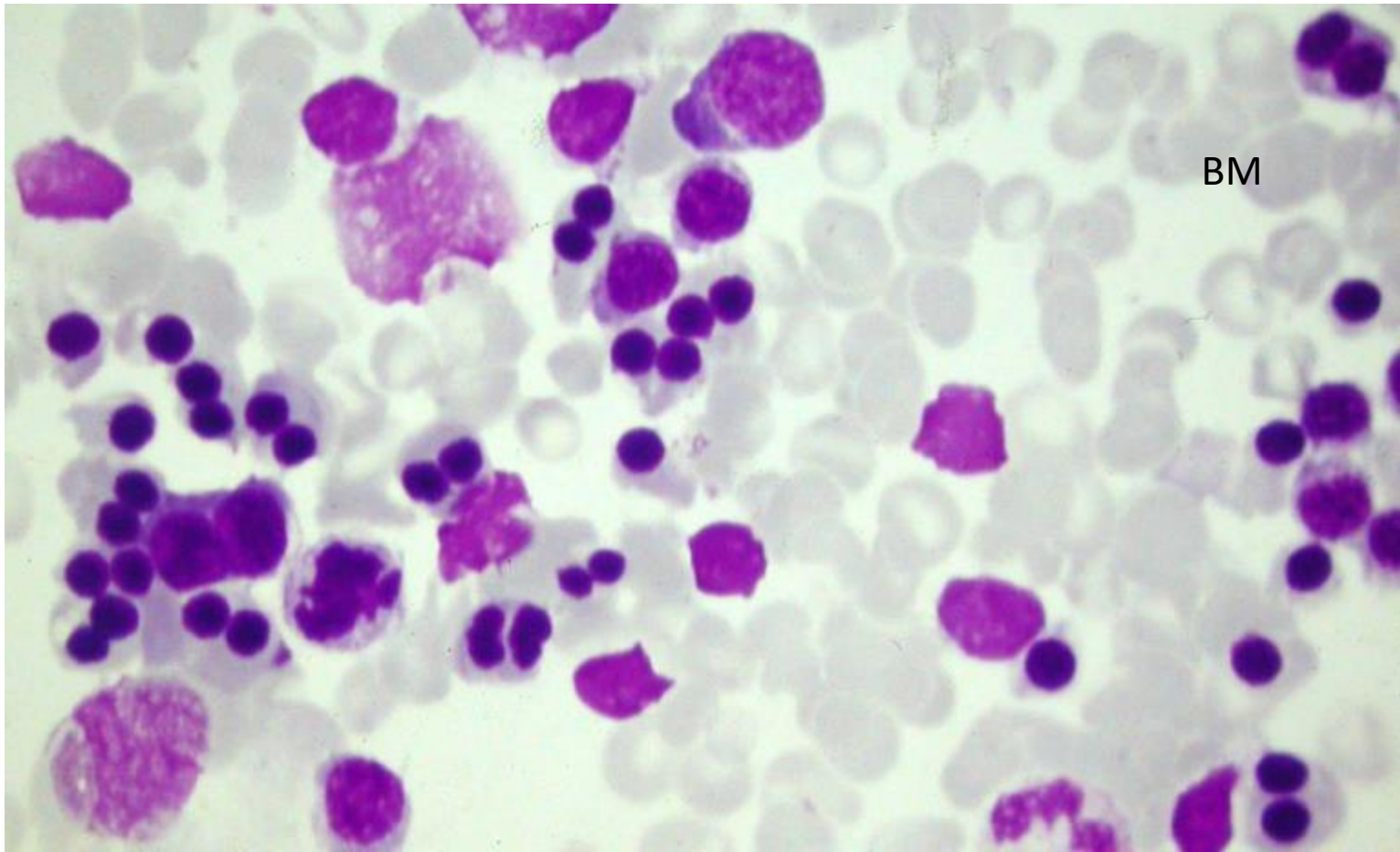


Helmet and fragmented rbc, microspherocytes

Microangiopathic hemolytic anemia

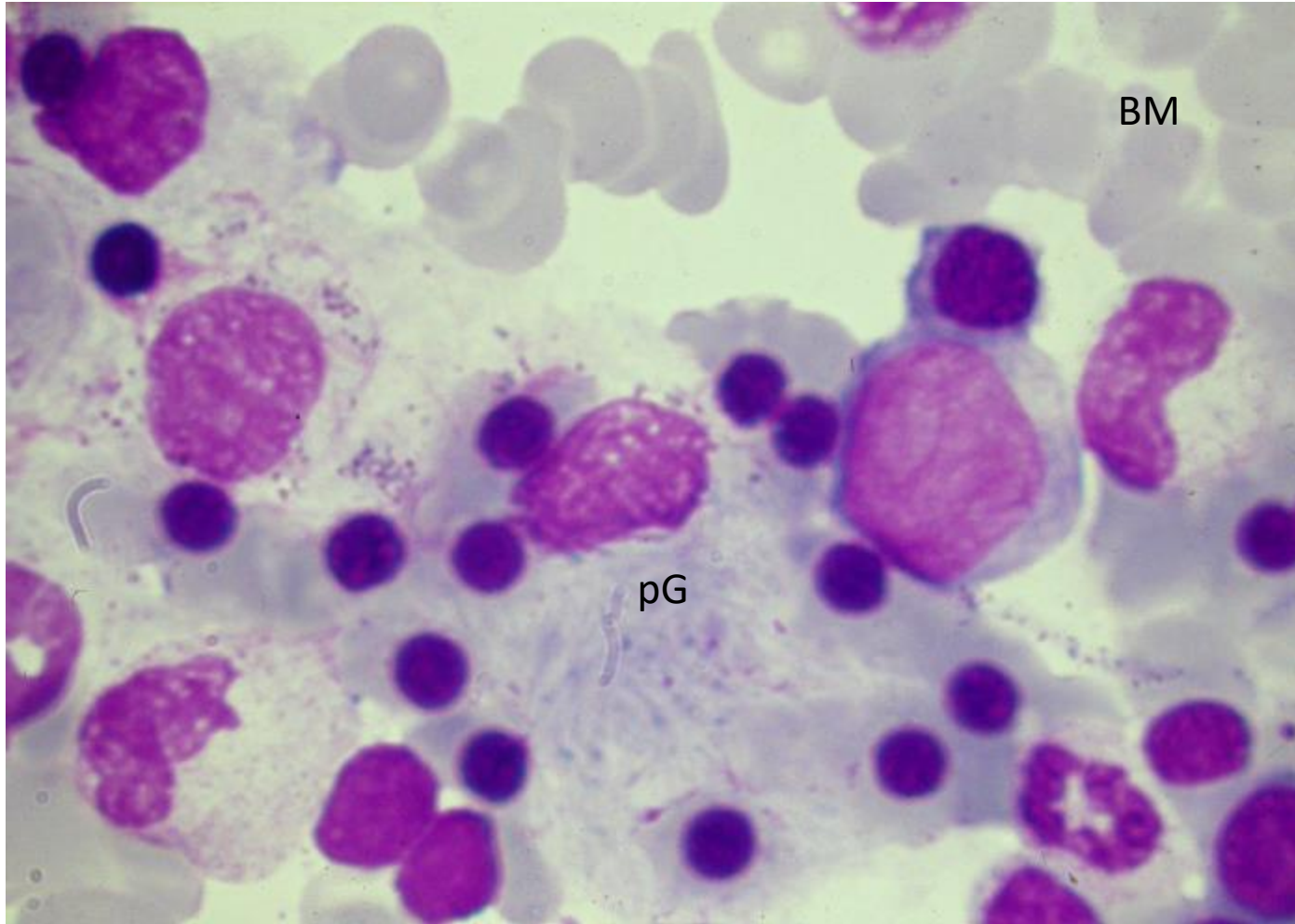


Congenital dyserythropoietic anemia type II (CDA II - HEMPAS)

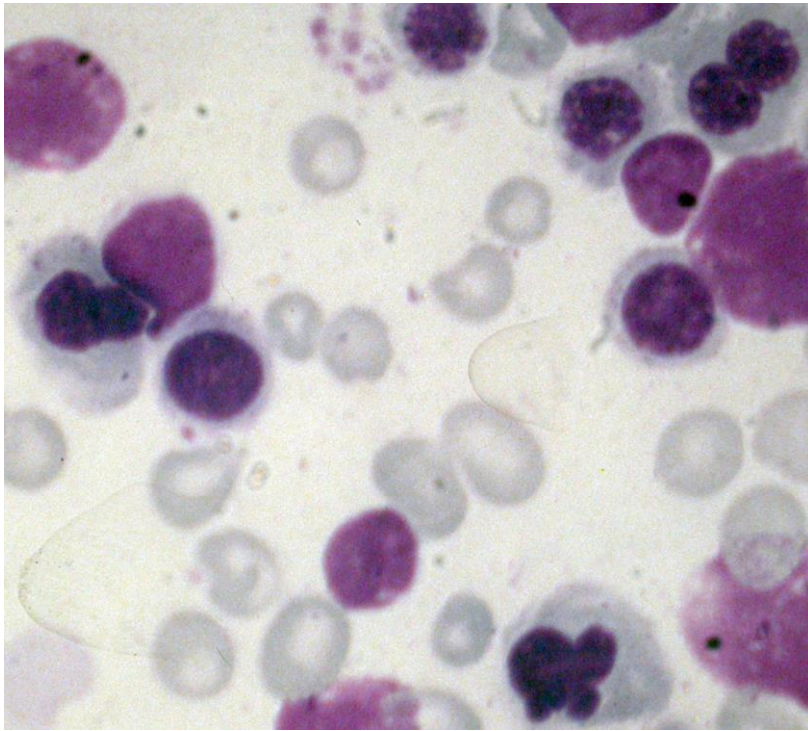


Erythroid hyperplasia with many binucleated late normoblasts

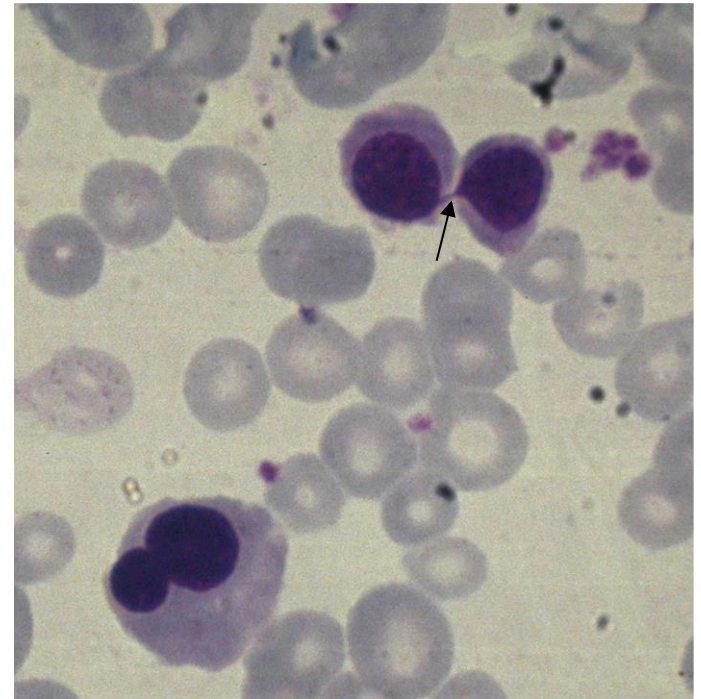
CDA II - HEMPAS



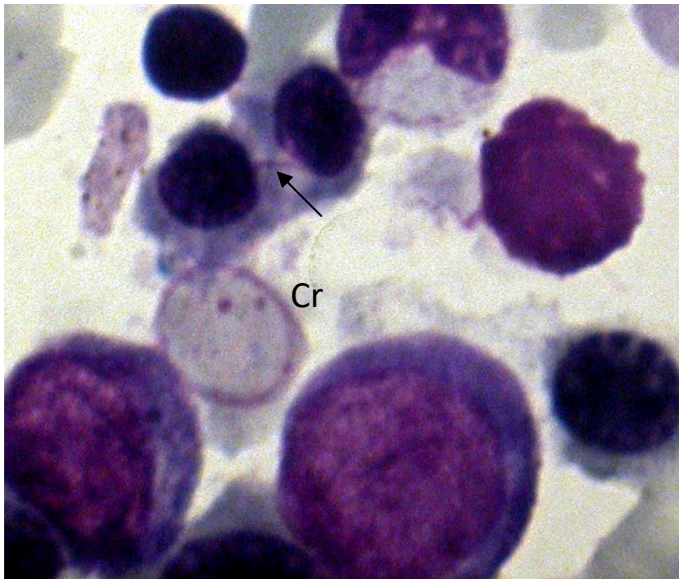
Binucleated normoblasts and pseudo-Gaucher cell (G)



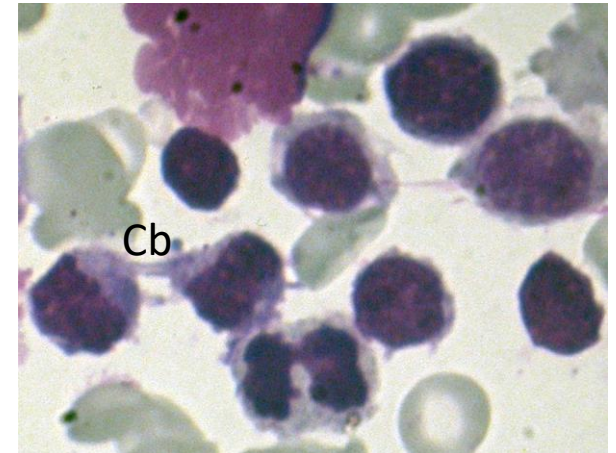
CDAI



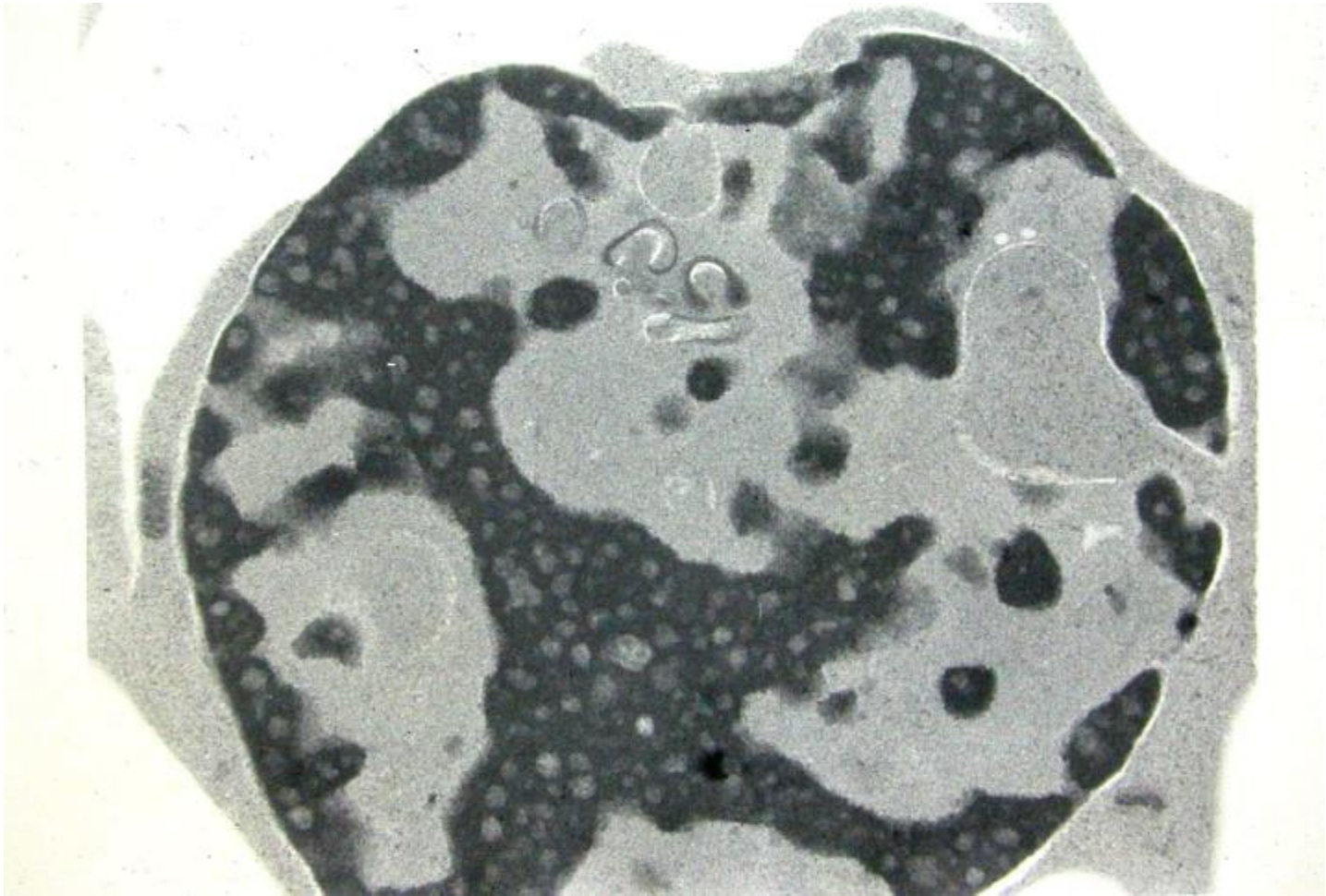
BM



Dyserythropoiesis with binucleated erythroblasts of different sizes and internuclear chromatin bridges (arrow). A Cabot ring in a rbc (Cr), and cytoplasmic bridges Cb) between erythroblasts.



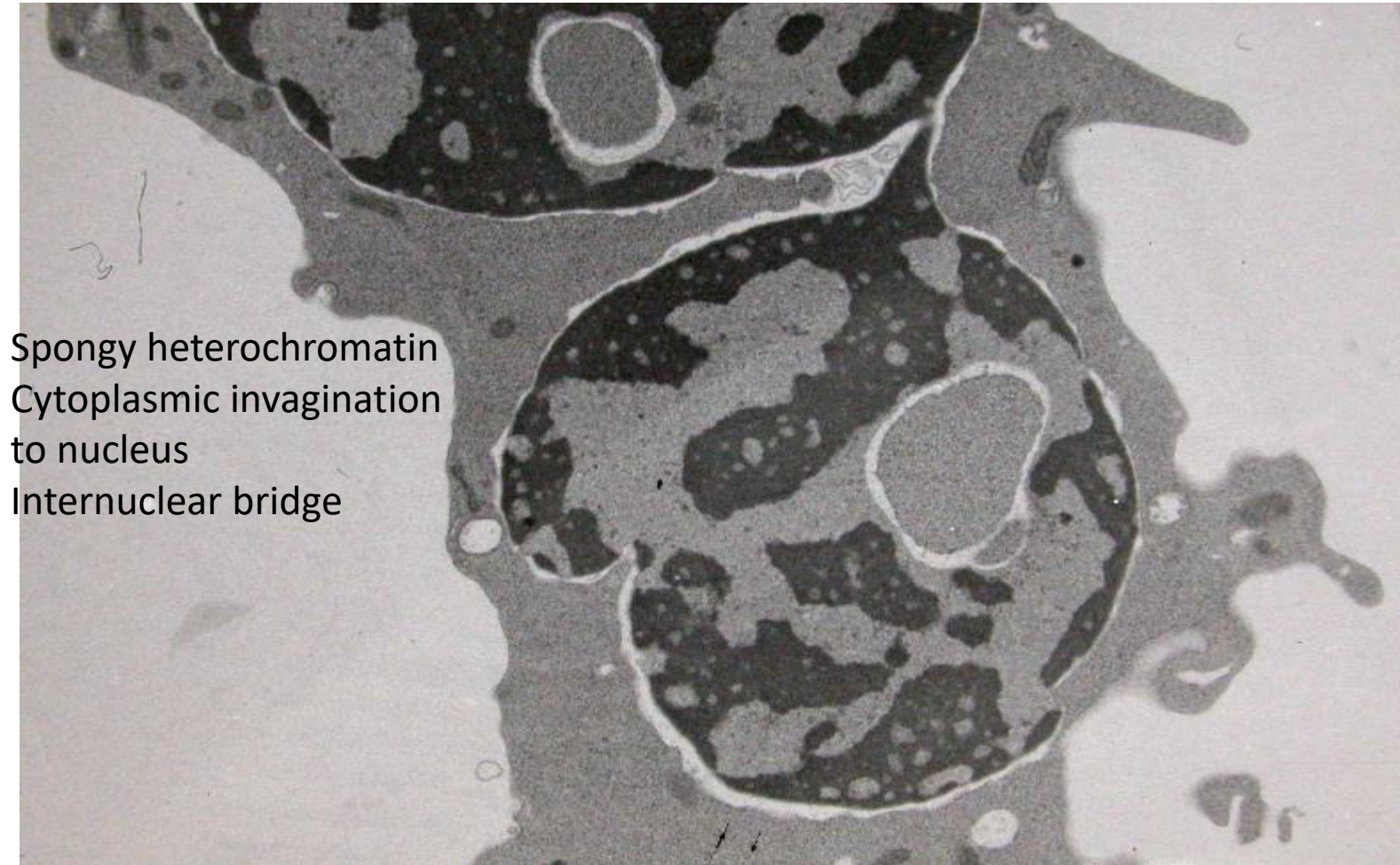
CDA I



EM

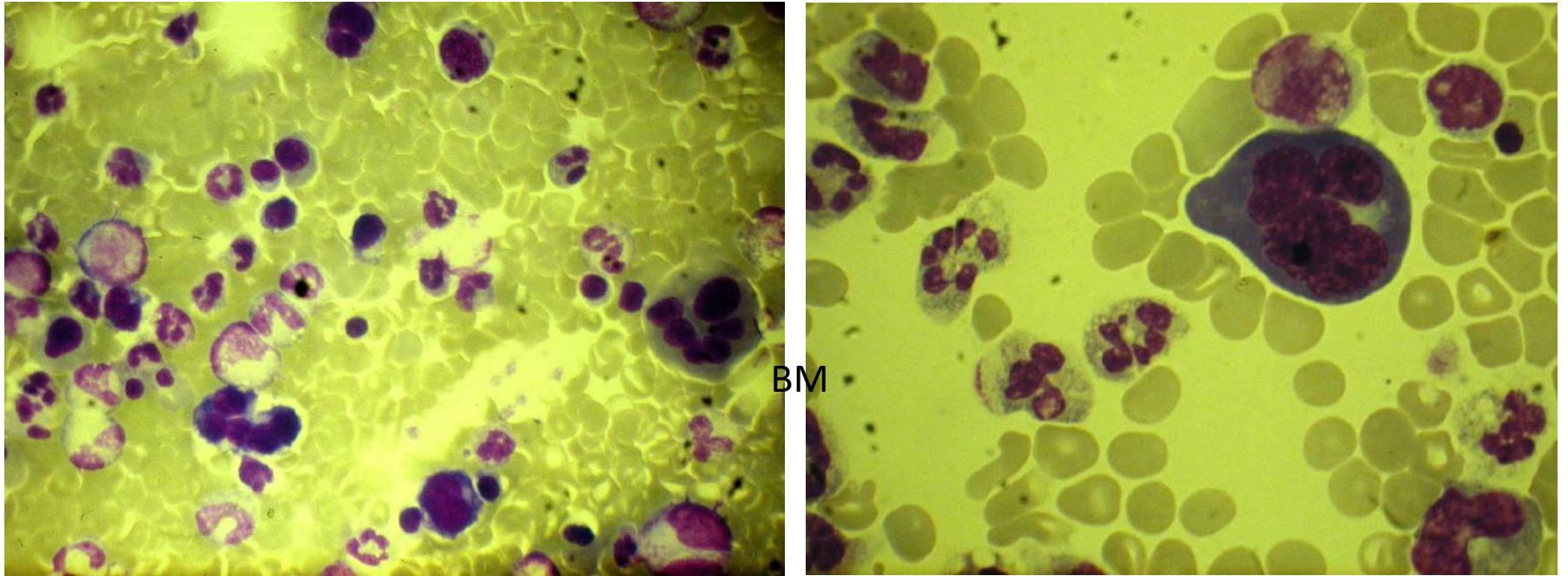
**“spongy” heterochromatin, dilatation of nuclear pores
and cytoplasmic invagination into the nucleus**

CDA I diagnosed by EM



Spongy heterochromatin
Cytoplasmic invagination
to nucleus
Internuclear bridge

CD III



Dyserythropoiesis with gyant multinuclear erythroblasts