Open Peer Review on Qeios

Hemoglobin H disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hemoglobin H disease. ORPHA:93616

Hemoglobin H (HbH) disease is a moderate to severe form of alpha-thalassemia (see this term) characterized by pronounced microcytic hypochromic hemolytic anemia.