Open Peer Review on Qeios

Hemoglobin E-beta-thalassemia syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hemoglobin E-beta-thalassemia syndrome. ORPHA:231249

Hemoglobin E - beta-thalassemia (HbE - BT) is a form of beta-thalassemia (see this term) that results in a mild to severe clinical presentation ranging from a condition indistinguishable from beta-thalassemia major to a mild form of beta-thalassemia intermedia (see these terms).