Hemoglobin E disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hemoglobin E disease. ORPHA:2133

Hemoglobin E disease (HbE) is a hemoglobinopathy characterized by production of abnormal variant hemoglobin known as hemoglobin E, with a generally benign, asymptomatic presentation.