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Hemoglobin D disease

INSERM

Source

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

Hemoglobin D disease. ORPHA:90039

Hemoglobin D disease(HbD) is a hemoglobinopathy characterized by production of abnormal variant hemoglobin known as hemoglobin D, with no or mild clinical manifestations (splenomegaly, very mild anemia).