Hemoglobin C disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hemoglobin C disease. ORPHA:2132

Hemoglobin C disease (HbC) is a hemoglobinopathy characterized by production of abnormal variant hemoglobin known as hemoglobin C, with no or mild clinical manifestations (hemolytic anemia).