Hereditary spherocytosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hereditary spherocytosis. ORPHA:822

Hereditary spherocytosis is a congenital hemolytic anemia with a wide clinical spectrum (from symptom-free carriers to severe hemolysis) characterized by anemia, variable jaundice, splenomegaly and cholelithiasis.