Hereditary elliptocytosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hereditary elliptocytosis. ORPHA:288

Hereditary elliptocytosis (HE) is a rare clinically and genetically heterogeneous disorder of the red cell membrane characterized by manifestations ranging from mild to severe transfusion-dependent hemolytic anemia but with the majority of patients being asymptomatic.