Hereditary methemoglobinemia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hereditary methemoglobinemia. ORPHA:621

Hereditary methemoglobinemia (HM) is a rare red cell disorder classified principally into two clinical phenotypes: autosomal recessive congenital (or hereditary) methemoglobinemia types I and II (RCM/RHM type 1; RCM/RHM type 2, see these terms).