Hemochromatosis type 4

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hemochromatosis type 4. ORPHA:139491

Hemochromatosis type 4 (also called ferroportin disease) is a form of rare hereditary hemochromatosis (HH; see this term), a group of diseases characterized by excessive tissue iron deposition of genetic origin.