Hereditary coproporphyria

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hereditary coproporphyria. ORPHA:79273

Hereditary coproporphyria is a form of acute hepatic porphyria (see this term) characterized by the occurrence of neuro-visceral attacks and, more rarely, by the presence of cutaneous lesions.