Dyschromatosis symmetrica hereditaria

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Dyschromatosis symmetrica hereditaria. ORPHA:41

Acropigmentation of Dohi is a genodermatosis characterised by the presence of hyperpigmented and hypopigmented macules, principally located on the extremities and limbs.