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Oculocutaneous albinism type 3

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Oculocutaneous albinism type 3. ORPHA:79433

Type 3 oculocutaneous albinism (OCA3) is a form of oculocutaneous albinism (OCA; see this term) characterized by rufous or brown albinism and occurring mainly in the African population.