Oculocutaneous albinism type 4

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Oculocutaneous albinism type 4. ORPHA:79435

Oculocutaneous albinism type 4 (OCA4) is a type of OCA (see this term) characterized by varying degrees of skin and hair hypopigmentation, numerous ocular changes and misrouting of the optic nerves at the chiasm.