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Oculocutaneous albinism type 1

INSERM

Source

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

Oculocutaneous albinism type 1. ORPHA:352731

Type 1 oculocutaneous albinism (OCA1) describes a group of tyrosine related OCAs (see this term) that includes OCA1A, OCA1B, type 1 minimal pigment oculocutaneous albinism (OCA1-MP) and type 1 temperature sensitive oculocutaneous albinism (OCA1-TS) (see these terms).

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