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Oculocerebrocutaneous syndrome

INSERM

Source

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

Oculocerebrocutaneous syndrome. ORPHA:1647

Oculocerebrocutaneous syndrome (OCCS) is a rare congenital disorder associated with an intellectual disability and is typically characterized by the triad of eye, central nervous system and skin malformations.