Oculocerebrorenal syndrome of Lowe

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Oculocerebrorenal syndrome of Lowe. ORPHA:534

Oculocerebrorenal syndrome of Lowe (OCRL) is a multisystem disorder characterized by congenital cataracts, glaucoma, intellectual disabilities, postnatal growth retardation and renal tubular dysfunction with chronic renal failure.