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Tyrosinemia type 2

INSERM

Source

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

Tyrosinemia type 2. ORPHA:28378

Tyrosinemia type 2 is an inborn error of tyrosine metabolism characterized by hypertyrosinemia with oculocutaneous manifestations and, in some cases, intellectual deficit.