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Primary hyperoxaluria type 3

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Primary hyperoxaluria type 3. ORPHA:93600

Primary hyperoxaluria type 3 (PH3) is a disorder of glyoxylate metabolism that can be asymptomatic or characterized by oxalate nephrolithiasis.