Hypotonia-cystinuria syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hypotonia-cystinuria syndrome. ORPHA:163690

Hypotonia-Cystinuria syndrome (HCS) is a rare syndrome including neonatal and infantile hypotonia and failure to thrive, cystinuria type 1 and nephrolithiasis, growth retardation due to growth hormone deficiency, and minor facial dysmorphism.