Coffin-Lowry syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Coffin-Lowry syndrome. ORPHA:192

Coffin-Lowry syndrome (CLS) is a rare genetic neurological disorder characterized by psychomotor and growth retardation, facial dysmorphism, digit abnormalities, and progressive skeletal changes.