Mitochondrial DNA-associated Leigh syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mitochondrial DNA-associated Leigh syndrome. ORPHA:255210

Maternally inherited Leigh syndrome is a rare subtype of Leigh syndrome (see this term) characterized clinically by encephalopathy, lactic acidosis, seizures, cardiomyopathy, respiratory disorders and developmental delay, with onset in infancy or early childhood, and resulting from maternally-inherited mutations in mitochondrial DNA.