Congenital cataract-hypertrophic cardiomyopathy-mitochondrial myopathy syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital cataract-hypertrophic cardiomyopathy-mitochondrial myopathy syndrome. ORPHA:1369

Congenital cataract - hypertrophic cardiomyopathy - mitochrondrial myopathy (CCM) is a mitochondrial disease (see this term) characterized by cataracts, hypertrophic cardiomyopathy, muscle weakness and lactic acidosis after exercise.