Kearns-Sayre syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Kearns-Sayre syndrome. ORPHA:480

Kearns-Sayre syndrome (KSS) is a mitochondrial disease characterized by progressive external ophthalmoplegia (PEO), pigmentary retinitis and an onset before the age of 20 years. Common additional features include deafness, cerebellar ataxia and heart block.