Mohr-Tranebjaerg syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Mohr-Tranebjaerg syndrome. ORPHA:52368

Mohr-Tranebjaerg syndrome (MTS) is an X-linked recessive neurodegenerative syndrome characterized by clinical manifestations commencing with early childhood onset hearing loss, followed by adolescent onset progressive dystonia or ataxia, visual impairment from early adulthood onwards and dementia from the 4th decade onwards.