Open Peer Review on Qeios

Autosomal recessive spastic paraplegia type 60

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal recessive spastic paraplegia type 60. ORPHA:401800

Autosomal recessive spastic paraplegia type 60 is a rare, complex hereditary spastic paraplegia disorder characterized by infantile onset of progressive lower limb spasticity, inability to walk, hypertonia and impaired vibration sense at ankles, with complicating signs including sensory impairment, nystagmus, motor axonal neuropathy and mild intellectual disability.