Autosomal recessive spastic paraplegia type 27

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal recessive spastic paraplegia type 27. ORPHA:101007

Autosomal recessive spastic paraplegia type 27 is a rare, pure or complex hereditary spastic paraplegia characterized by a variable onset of slowly progressive lower limb spasticity, hyperreflexia and extensor plantar responses, that may be associated with sensorimotor polyneuropathy, decreased vibration sense, lower limb distal muscle wasting, dysarthria and mild to moderate intellectual disability.