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Autosomal spastic paraplegia type 72

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base. Autosomal spastic paraplegia type 72. ORPHA:401849*

Autosomal spastic paraplegia type 72 is a rare, genetic, pure hereditary spastic paraplegia disorder characterized by early childhood onset of slowly progressive crural spastic paraparesis presenting with spastic gait, mild stiffness at rest, hyperreflexia (in lower limbs), extensor plantar responses and, in some, mild postural tremor, pes cavus, sphincter disturbances and sensory loss at ankles.