Autosomal dominant spastic paraplegia type 13

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant spastic paraplegia type 13. ORPHA:100994

A rare hereditary spastic paraplegia characterized by progressive spastic paraplegia with pyramidal signs in the lower limbs, decreased vibration sense, and increased reflexes in the upper limbs.