Distal hereditary motor neuropathy type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Distal hereditary motor neuropathy type 1. ORPHA:139518

Distal hereditary motor neuropathy type 1 is a rare neuromuscular disease characterized by slowly-progressive lower limb muscular weakness and atrophy, without sensory impairment. Additional clinical features may include pes cavus, hammertoe and increased muscle tone.