Periodic paralysis with later-onset distal motor neuropathy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Periodic paralysis with later-onset distal motor neuropathy. ORPHA:397750

Periodic paralysis with later-onset distal motor neuropathy is a rare, genetic, neuromuscular disease characterized by acute episodic muscle weakness in upper and lower extremities (which responds to acetazolamide treatment) associated with later-onset, chronic, slowly progressive, distal, axonal neuropathy.