Autosomal dominant Charcot-Marie-Tooth disease type 2N

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant Charcot-Marie-Tooth disease type 2N. ORPHA:228174

Autosomal dominant Charcot-Marie-Tooth disease type 2N (CMT2N) is a mild form of axonal Charcot-Marie-Tooth disease, a peripheral sensorimotor neuropathy, characterized by distal legs sensory loss and weakness that can be asymmetric. Tendon reflexes are reduced in the knees and absent in ankles. Progression is slow.