Autosomal dominant Charcot-Marie-Tooth disease type 2A1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant Charcot-Marie-Tooth disease type 2A1. ORPHA:99946

Autosomal dominant Charcot-Marie-Tooth disease type 2A1 (CMT2A1) is a form of axonal Charcot-Marie-Tooth disease, a peripheral sensorimotor neuropathy. CMT2A presents with a more prominent muscle weakness in lower than upper limbs and frequent postural tremor.