# Autosomal dominant Charcot-Marie-Tooth disease type 2I

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant Charcot-Marie-Tooth disease type 2I. ORPHA:99942*

Autosomal dominant Charcot-Marie-Tooth disease type 2I (CMT2I) is a form of axonal Charcot-Marie-Tooth disease, a peripheral sensorimotor neuropathy, characterized by a late onset with severe sensory loss (paresthesia and hypoesthesia) associated with distal weakness, mainly of the legs, and absent or reduced deep tendon reflexes.