## Opis choroby \*

## Definicja

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.

Dane

**Klasyfikacja** Choroba Synonimy CMT2I CMT2I

**Kod ORPHA** 99942

**Kod OMIM** 607677

**Kod ICD10** G60.0

**Kod ICD11** 8C20.1

## \*Źródło

orphanet