Opis choroby *

Definicja

A rare hereditary motor and sensory neuropathy disorder characterized by the typical CMT phenotype (slowly progressive distal muscle atrophy and weakness in upper and lower limbs, distal sensory loss in extremities, reduced or absent deep tendon reflexes and foot deformities) with nerve biopsy demonstrating demyelinating and axonal changes and nerve conduction velocities varying from the demyelinating to axonal range.

Dane

Klasyfikacja
Choroba

Synonimy
CMTDIF
CMTDIF

Kod ORPHA
352670

Kod OMIM
615185

Kod ICD10
G60.0

Kod ICD11
8C20.2

---

*Źródło

orphanet