Opis choroby *

Definicja

A rare hereditary demyelinating motor and sensory neuropathy characterized by slowed nerve conduction velocities, in the absence of clinically apparent neurological deficits, gait abnormalities or muscular atrophy, associated with a germline mutation in the ARGHEF10 gene.

Dane

Klasyfikacja
Choroba

Kod ORPHA
140481

Kod OMIM
608236

Kod ICD10
G60.0

Kod ICD11
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*Źródło

orphanet