Opis choroby *

Definicja

A rare acquired motor neuron disease characterized by an initial unilateral weakness in the intrinsic hand muscles that eventually spreads to the opposite limb (with an asymmetrical distribution) and that has a very slow progression of muscular atrophy over a 20 year period.

Dane

Klasyfikacja

Choroba

Kod ORPHA 99965

Kod OMIM

Kod ICD10

G12.8

Kod ICD11

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*Źródło

orphanet