Opis choroby *

Definicja

A rare, genetic motor neuron disease characterized by late childhood- or adolescent-onset of slowly progressive, severe, distal limb muscle weakness and wasting, in association with pyramidal signs, normal sensation, and absence of bulbar involvement, leading to degeneration of motor neurons in the brain and spinal cord.

Dane

Klasyfikacja
Choroba

Synonimy
ALS4
ALS4
dHMN z objawami górnego neuronu ruchowego
Dystalna dziedziczna neuropatia ruchowa z objawami górnego neuronu ruchowego
Distal hereditary motor neuropathy with upper motor neuron signs
dHMN with upper motor neuron signs

Kod ORPHA
357043

Kod OMIM
602433

Kod ICD10
G12.2

Kod ICD11
8B60.0

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*Źródło

orphanet