

## 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](#)

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