

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	Synonimy
Choroba	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	Kod OMIM	Kod ICD10
357043	602433	G12.2

Kod ICD11
8B60.0

*Źródło

orphanet