Opis choroby *

Definicja

A group of rare, genetic, motor neuron disease characterized by childhood or adult onset progressive, predominantly proximal, muscular weakness and wasting. Included diseases are Autosomal dominant adult-onset proximal spinal muscular atrophy, Lower motor neuron syndrome with late-adult onset, and Autosomal dominant childhood-onset proximal spinal muscular atrophy.

Dane

Klasyfikacja
Grupa fenomenów

Kod ORPHA
211037

Kod OMIM
-

Kod ICD10
G12.1

Kod ICD11
-

---

*Źródło

orphanet