Opis choroby *

Definicja

A rare genetic movement disorder characterized by dystonia affecting at first an upper limb, less frequently beginning in the head and neck region, before slowly spreading to other locations. The clinical spectrum, like age of onset, is variable with focal, segmental, or generalized distribution, but cranial involvement with speech difficulties and cervical involvement are typical, whereas lower limbs are often spared. With progression of the disease, many patients suffer from generalized dystonia while mostly remaining ambulatory.

Dane

Klasyfikacja

Choroba

Synonimy

DYT6

Dystonia uogólniona rozpoczynająca się od szyi i

kończyn górnych

DYT6

Idiopatyczna dystonia torsyjna typu mieszanego

Generalized cervical and upper-limb-onset

dystonia

Idiopathic torsion dystonia of mixed type

Kod ORPHA

Kod OMIM

Kod ICD10

98806

602629

G24.1

Kod ICD11 8A02.0Y

*Źródło

orphanet