Opis choroby *

Definicja

A rare, genetic, movement disorder characterized by involuntary movements on one side of the body that mirror intentional movements on the opposite side of the body, which are present in various first-degree members of a family, persist beyond the first decade of life, and have no associated comorbidities.

Dane

Klasyfikacja
Choroba

Synonimy
Familial congenital controlateral synkinesia
Dziedziczna wrodzona synkinezja kontrlateralna
Dziedziczne wrodzone ruchy lustrzane
Izolowana wrodzona synkinezja kontrlateralna
Izolowane wrodzone ruchy lustrzane
Rodzinna wrodzona synkinezja kontrlateralna
Hereditary congenital controlateral synkinesia
Hereditary congenital mirror movements
Isolated congenital controlateral synkinesia
Isolated congenital mirror movements

Kod ORPHA
238722

Kod OMIM
616059

Kod ICD10
G25.8

Kod ICD11
8A0Y

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*Źródło

orphanet