

Opis choroby *

Definicja

A heterogenous group of cerebral cortical malformations characterized by excessive cortical folding and abnormal cortical layering that, depending on its topographic distribution, presents with variable combinations of neurological symptoms of varying severity such as epilepsy, developmental delay, intellectual disability, motor dysfunction (e.g. spasticity), and pseudobulbar palsy.

Dane

Klasyfikacja

Grupa fenomenów

Kod ORPHA

35981

Kod OMIM

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Kod ICD10

Q04.3

Kod ICD11

LA05.50

*Źródło

orphanet