Polimikrogyria
Kod Orpha: 35981 Kod OMIM:

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
-

Kod ICD10
Q04.3

Kod ICD11
LA05.50

---

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.