Opis choroby *

Definicja

A rare, genetic neurological disorder characterized by the presence of diffuse pachygyria and arachnoid cysts, psychomotor developmental delay and intellectual disability. Seizures (absence, atonic and generalized tonic-clonic) and, on occasion, headache are also associated.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Kuzniecky syndrome
Zespół Kuzniecky'ego

Kod ORPHA
2798

Kod OMIM
600176

Kod ICD10
G40.4

Kod ICD11
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*Źródło

orphanet