Opis choroby *

Definicja

A rare developmental defect during embryogenesis characterized by the presence of linear clefts containing cerebrospinal fluid lined by abnormal grey matter that extend from the lateral ventricles to the pial surface of the cortex. Schizencephaly can involve one or both cerebral hemispheres and may lead to a variety of neurological symptoms such as epilepsy, motor deficits, and psychomotor retardation.

Dane

Klasyfikacja
Choroba

Kod ORPHA
799

Kod OMIM
269160

Kod ICD10
Q04.6

Kod ICD11
LA05.61

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*Źródło

orphanet