

## 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