

Opis choroby *

Definicja

The term lissencephaly covers a group of rare malformations sharing the common feature of anomalies in the appearance of brain convolutions (characterised by simplification or absence of folding) associated with abnormal organisation of the cortical layers as a result of neuronal migration defects during embryogenesis.

Dane

Klasyfikacja

Kategoria

Kod ORPHA

48471

Kod OMIM

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Kod ICD10

Q04.3

Kod ICD11

LD20.1

*Źródło

orphanet