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
-

Kod ICD10
Q04.3

Kod ICD11
LD20.1

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*Źródło

orphanet