Opis choroby *

Definicja

Lissencephaly syndrome, Norman-Roberts type is characterised by the association of lissencephaly type I with craniofacial anomalies (severe microcephaly, a low sloping forehead, a broad and prominent nasal bridge and widely set eyes) and postnatal growth retardation.

Dane

Klasyfikacja
Podtyp kliniczny

Synonimy
Microlissencephaly type A
Mikrolizencefalia typu A

Kod ORPHA
89844

Kod OMIM
257320

Kod ICD10
Q04.3

Kod ICD11
LD20.1

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*Źródło

orphanet