

## 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

---

#### \*Źródło

orphanet