

## Opis choroby \*

### Definicja

A rare syndromic form of lissencephaly characterized by severe microcephaly, agyria, agenesis of the corpus callosum, cerebellar hypoplasia, facial dysmorphology and epiphyseal stippling of the metacarpal bones. The syndrome may be an allelic variant of Neu-Laxova syndrome and Lissencephaly type III with cystic dilations of the cerebellum and foetal akinesia sequence.

### Dane

### Klasyfikacja

Zespół wad wrodzonych

#### Kod ORPHA

86822

#### Kod OMIM

601160

#### Kod ICD10

Q04.3

#### Kod ICD11

LD20.1

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### \*Źródło

orphanet