

Opis choroby *

Definicja

A severe form of lissencephaly with cerebellar hypoplasia, characterized by a microcephaly of at least - 3 SD and a thick cortex associated with complete absence of the corpus callosum.

Dane

Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA

100016

Kod OMIM

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Kod ICD10

Q04.3

Kod ICD11

LD20.1

*Źródło

orphanet