

Opis choroby *

Definicja

A rare, genetic, lissencephaly with cerebellar hypoplasia subtype characterized by classical lissencephaly with thickened cortical gray matter (with either no discernable gradient, a predominantly posterior gradient, or a predominantly anterior gradient) associated with variable, predominantly midline, cerebellar hypoplasia.

Dane

Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA

100011

Kod OMIM

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Kod ICD10

Q04.3

Kod ICD11

LD20.1

*Źródło

orphanet