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

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*Źródło

orphanet