Opis choroby *

Definicja

A form of lissencephaly with cerebellar hypoplasia characterized by subtle microcephaly, hypotonia and neurological and cognitive development delay. Hippocampal malformation is a characteristic imaging feature of this disorder.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
100012

Kod OMIM
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Kod ICD10
Q04.3

Kod ICD11
LD20.1

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*Źródło

orphanet