Opis choroby *

Definicja

A form of holoprosencephaly characterized by fusion of the left and right frontal and parietal lobes with only a posterior interhemispheric fissure. Craniofacial features variably include ocular hypotelorism, midline cleft lip (complete or partial) and a flat nose.

Dane

Klasyfikacja
Podtyp kliniczny

Kod ORPHA
220386

Kod OMIM
610829

Kod ICD10
Q04.2

Kod ICD11
LA05.2

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*Źródło

orphanet