## 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