Opis choroby *

Definicja

A rare complex brain malformation characterized by incomplete cleavage of the prosencephalon, and affecting both the forebrain and face and resulting in neurological manifestations and facial anomalies of variable severity.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
HPE
HPE

Kod ORPHA
2162

Kod OMIM
609408

Kod ICD10
Q04.2

Kod ICD11
LA05.2

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*Źródło

orphanet