

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	Synonimy
Zespół wad wrodzonych HPE	HPE

Kod ORPHA	Kod OMIM	Kod ICD10
2162	609408	Q04.2

Kod ICD11
LA05.2

*Źródło

orphanet