Diprosopia
Kod Orpha: 1681 Kod OMIM:

Opis choroby *

Definicja
Diprosopus is a rare, life-threatening developmental defect during embryogenesis, and a subtype of conjoined twins, characterized by partial or complete duplication of the facial structures on a single head, neck, trunk and body. It may be associated with congenital anomalies involving the cardiovascular, gastrointestinal, respiratory and central nervous systems. Cleft lip and palate have been reported in rare cases.
Dane

Klasyfikacja
Wada morfologiczna

Synonimy
Craniofacial duplication
Diprosopia

Kod ORPHA
1681

Kod OMIM
-

Kod ICD10
Q89.4

Kod ICD11
-

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.