

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

-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.