Opis choroby *

Definicja

A rare developmental defect during embryogenesis syndrome characterized by a glabellar capillary malformation, congenital communicating hydrocephalus, and posterior fossa brain abnormalities, including Dandy-Walker malformation, cerebellar vermis agenesis, and mega cisterna magna. Seizures are occasionally associated. There have been no further descriptions in the literature since 1979.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Nova syndrome
Zespół Nova

Kod ORPHA
2703

Kod OMIM
-

Kod ICD10
Q03.8

Kod ICD11
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*Źródło

orphanet