

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	Synonimy
Zespół wad wrodzonych	Nova syndrome
	Zespół Nova

Kod ORPHA	Kod OMIM	Kod ICD10
2703	-	Q03.8

Kod ICD11
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*Źródło

orphanet