

## 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

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Nova syndrome Zespół Nova

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
2703	-	Q03.8

**Kod ICD11**  
-

---

### \*Źródło

orphanet