

## Opis choroby \*

### Definicja

A rare cerebral malformation characterized by an almost or complete lack of cortex, specifically the cerebral hemispheres, with the cranium and meninges completely intact. In most cases, death occurs in utero or in the first weeks of life. Developmental delay, drug-resistant seizures, spastic diplegia, severe growth failure, deafness and blindness are typical.

### Dane

### Klasyfikacja

Zespół wad wrodzonych

#### Kod ORPHA

2177

#### Kod OMIM

-

#### Kod ICD10

Q04.3

#### Kod ICD11

LA05.62

---

### \*Źródło

orphanet