

Opis choroby *

Definicja

A rare, genetic or acquired, cerebral malformation characterized by an intracerebral fluid-filled cyst or cavity with or without communication between the ventricle and subarachnoid space. Clinical manifestations depend on location and severity and may include hemiparesis, seizures, intellectual disability, and dystonia.

Dane

Klasyfikacja

Choroba

Kod ORPHA

2940

Kod OMIM

614483

Kod ICD10

Q04.6

Kod ICD11

LA05.60

*Źródło

orphanet