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

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*Źródło

orphanet