Opis choroby *

Definicja

A very rare glial neoplasm of the central nervous system, most often with an intra-axial peripheral supratentorial location in one hemisphere of the frontal or parietal lobes and usually presenting in infants and young adults with symptoms of vomiting, loss of consciousness, epileptic seizures and headaches.

Dane

Klasyfikacja
Choroba

Kod ORPHA
251679

Kod OMIM
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Kod ICD10
C71.9

Kod ICD11
2A00.4

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*Źródło

orphanet