Opis choroby *

Definicja

An extremely rare slow-growing glial neoplasm of the central nervous system, usually arising in a superficial location in the cerebrum, affecting all ages and both sexes, and characterized by intractable seizures and headaches, with most cases being cured by surgical incision alone and therefore having a good prognosis.

Dane

Klasyfikacja
Choroba

Kod ORPHA
251671

Kod OMIM
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Kod ICD10
C71.9

Kod ICD11
2A00.0Y

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*Źródło

orphanet