

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

-

Kod ICD10

C71.9

Kod ICD11

2A00.0Y

*Źródło

orphanet