Opis choroby *

Definicja

A rare mixed neuronal-glial tumor characterized by a mostly supratentorial space-occupying lesion often involving the temporal lobe, although it may occur anywhere in the central nervous system. The tumor shows anaplastic features in its glial component and is considered WHO grade III, which may, albeit inconsistently, indicate more aggressive behavior and less favorable prognosis. Clinical symptoms vary according to the location, the most common manifestation being seizures.

Dane

Klasyfikacja
Choroba

Kod ORPHA
251957

Kod OMIM
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Kod ICD10
D43.0

Kod ICD11
2A00.21

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*Źródło

orphanet