

## Opis choroby \*

### Definicja

A rare mixed neuronal-glia 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