

## Opis choroby \*

### Definicja

A rare ependymal tumor characterized by the presence of a *RELN* fusion gene. This supratentorial grade II or III ependymoma most often occurs in children and young adults. Histopathological features are variable, but a distinctive vascular pattern of branching capillaries or clear-cell change are common. Patients may present with focal neurological deficits, seizures, or features of raised intracranial pressure. Prognosis is worse than in other supratentorial ependymomas.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Supratentorial C11ORF95-RELN fused ependymoma  
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#### Kod ORPHA

530792

#### Kod OMIM

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#### Kod ICD10

D43.0

#### Kod ICD11

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#### \*Źródło

orphanet