Opis choroby *

Definicja

A rare glial tumor characterized by a grade III oligodendroglial tumour with focal or diffuse anaplastic features. It typically occurs in the supratentorial white matter. Histologically, the cells are enlarged and epithelioid with pleomorphic and increased size nuclei, a vesicular chromatin pattern and prominent nucleoli. Most patients present with seizures.

Dane

Klasyfikacja
Choroba

Kod ORPHA
251630

Kod OMIM
616568

Kod ICD10
C71.9

Kod ICD11
2A00.0Y

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*Źródło

orphanet