

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

*Źródło

orphonet