Opis choroby *

Definicja

A rare glial tumor characterized by a highly cellular lesion that is diffusly infiltrating at the periphery and consists of evenly-spaced monomorphic cells with the oligodendroglial phenotype. It typically occurs in the supratentorial white matter. Histologically, the cells are uniformly round to oval with round nuclei, delicate chromatin and small nucleoli. Most patients present with seizures.

Dane

Klasyfikacja
Choroba

Kod ORPHA
251627

Kod OMIM
616568

Kod ICD10
C71.9

Kod ICD11
2A00.0Y

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*Źródło

orphanet