

Opis choroby *

Definicja

A rare glial tumor characterized by a highly cellular lesion that is diffusely 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

*Źródło

orphanet