Opis choroby *

Definicja

A rare mixed neuronal-glial tumor characterized by a benign, usually supratentorial lesion with predominantly cortical location and multinodular architecture. The tumor typically becomes symptomatic in the second or third decade of life with drug-resistant partial seizures. Histological hallmark is the specific glioneuronal element, columns oriented perpendicularly to the cortical surface, formed by bundles of axons attached to oligodendroglia-like cells, while neurons appear to float in an abundant eosinophilic matrix.

Dane

Klasyfikacja
Choroba

Synonimy
DNET
DNET

Kod ORPHA
251946

Kod OMIM
-

Kod ICD10
D33.0

Kod ICD11
2A00.21

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*Źródło

orphanet