Opis choroby *

Definicja

A rare low-grade astrocytoma characterized by a high degree of cellular differentiation, slow growth, and diffuse infiltration of adjacent brain structures, and corresponding to WHO grade II. The tumor typically affects young adults and has an intrinsic tendency for progression to high-grade glioma. Histological variants are fibrillary, gemistocytic, and protoplasmic astrocytoma. Patients most commonly present with seizures, but also with other neurological or neuropsychological abnormalities, depending on the location.

Dane

Klasyfikacja
Choroba

Kod ORPHA
251595

Kod OMIM
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Kod ICD10
C71.9

Kod ICD11
2A00.0Y

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*Źródło

orphanet