Opis choroby *

Definicja

A rare low-grade astrocytoma characterized by superficial location in the cerebral hemispheres with involvement of the meninges, composed of GFAP-expressing cells showing nuclear and cytoplasmic pleomorphism and xanthomatous change, surrounded by a reticulin network. The tumor corresponds to WHO grade II and typically affects children and young adults, who often present with a long history of seizures. Extent of resection and mitotic index are important prognostic factors.

Dane

Klasyfikacja
Choroba

Synonimy
PXA
PXA

Kod ORPHA
251607

Kod OMIM
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Kod ICD10
C71.9

Kod ICD11
2A00.0Y

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*Źródło

orphanet