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 Synonimy

Choroba PXA

 PXA

Kod ORPHA Kod OMIM Kod ICD10

251607 - C71.9

Kod ICD11 2A00.0Y

*Źródło

orphanet