Astroblastoma

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Astroblastoma. ORPHA:251679

Astroblastoma is a very rare glial neoplasm of the central nervous system, most often with an intra-axial peripheral supratentorial location in one hemisphere of the frontal or parietal lobes and usually presenting in infants and young adults with symptoms of vomiting, loss of consciousness, epileptic seizures and headaches.