## Opis choroby *

Definicja

A rare low-grade astrocytoma characterized by a benign, slowly growing lesion typically arising in the wall of the lateral ventricles, composed of large ganglioid astrocytes. The tumor corresponds to WHO grade I and typically occurs during the first two decades of life in patients with tuberous sclerosis complex. Most patients present with worsening of epilepsy or symptoms of increased intracranial pressure.

Dane

**Klasyfikacja**
Choroba

Synonimy
SEGA
SEGA

**Kod ORPHA**
251618

**Kod OMIM**
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**Kod ICD10**
D43.2

**Kod ICD11**
2A00.0Y

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*Źródło

orphanet