Juvenile absence epilepsy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Juvenile absence epilepsy. ORPHA:1941

Juvenile absence epilepsy (JAE) is a genetic epilepsy with onset occurring around puberty. JAE is characterized by sporadic occurrence of absence seizures, frequently associated with a long-life prevalence of generalized tonic-clonic seizures (GTCS) and sporadic myoclonic jerks.