DISEASE:
Juvenile absence epilepsy

NAME:	Juvenile absence epilepsy
DESCRIPTION:	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.
ORPHACODE:	1941
SYNONYMS:	JAE
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	EFHC1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Epilepsy gene panel

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- EF-hand domain containing 1

Related Gene Panels

- Epilepsy gene panel - VUB