DISEASE:
Autosomal dominant nocturnal frontal lobe epilepsy

NAME:	Autosomal dominant nocturnal frontal lobe epilepsy
DESCRIPTION:	A rare seizure disorder characterized by intermittent dystonia and/or choreoathetoid movements that occur during sleep. The clusters of nocturnal motor seizures are often stereotyped and brief.
ORPHACODE:	98784
SYNONYMS:	ADNFLE Autosomal dominant sleep-related hypermotor epilepsy
XREF(S):	Orphanet OMIM OMIM OMIM OMIM OMIM ICD-10

ANALYTE(S):	CHRNA4 CHRNA2 CHRNB2 KCNT1 DEPDC5 CRH CABP4
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Epilepsy (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- calcium binding protein 4
- cholinergic receptor nicotinic alpha 2 subunit
- cholinergic receptor nicotinic alpha 4 subunit
- cholinergic receptor nicotinic beta 2 subunit
- corticotropin releasing hormone
- DEP domain containing 5, GATOR1 subcomplex subunit
- potassium sodium-activated channel subfamily T member 1

Related Gene Panels

- Rare epilepsy with developmental delay (> 240 genes) - UZA