Belgian Genetic Tests database

ANALYTE:
ADGRG1

NAME:	adhesion G protein-coupled receptor G1
SYMBOL:	ADGRG1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	TM7LN4 TM7XN1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Ataxia Spasticity (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Malformations of cortical development (235 genes)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Polymicrogyria, bilateral perisylvian, frontoparietal

Related Diseases

- Bilateral frontoparietal polymicrogyria
- Bilateral perisylvian polymicrogyria

Related Gene Panels

- Ataxia (348 genes) - ULB
- Ataxia Spasticity - UGent

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Leukodystrophy - UGent
- Malformations of cortical development (235 genes) - VUB
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
