

ANALYTE:
TMEM231

NAME:	transmembrane protein 231
SYMBOL:	TMEM231
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ALYE870 FLJ22167 JBTS20 MKS11 PRO1886
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>SwissProt</u>
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- [Ataxia \(autosomic dominant and recessive / except expansion of triplets\) \(gene panel - 722 genes\)](#)
- [Ataxia Spasticity \(gene panel\)](#)
- [Ciliopathy \(gene panel\)](#)
- [Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophthisis / Bardet-Biedl syndromes and kidney cancers \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Early onset epileptic encephalopathy \(gene panel - 845 genes\)](#)
- [End-stage renal disease, ESRD \(gene panel\)](#)
- [Epilepsy gene panel](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)

Related Diseases

- Joubert syndrome with oculorenal defect
- Meckel syndrome
- Orofaciodigital syndrome type 3

Related Gene Panels

- Ataxia (348 genes) - ULB
- Ataxia Spasticity - UGent
- Ciliopathy (120 genes) - UGent
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophthisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Early onset epileptic encephalopathy (845 genes) - ULB
- End-stage renal disease (106 genes) - IPG
- Epilepsy gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Panel Nephro-ULG-V1
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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