ANALYTE:
PDE6D

NAME:	phosphodiesterase 6D
SYMBOL:	PDE6D
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	JBTS22 retinal rod rhodopsin-sensitive cGMP 3',5'-cyclic phosphodiesterase subunit delta
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	26 Oct 2023 - 23:49

RELATED CONTENT

Related Genetic Tests

- Ciliopathy (gene panel)
- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / 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)
- Epilepsy gene panel
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)

Related Diseases

- Joubert syndrome
- Orofaciodigital syndrome type 6

Related Gene Panels

- Ciliopathy (120 genes) - UGent
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, 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
- Epilepsy gene panel - VUB

- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1

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