

ANALYTE:
TCTN2

NAME:	tectonic family member 2
SYMBOL:	TCTN2
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SYNONYMS:	FLJ12975 JBTS24 MKS8 Meckel syndrome, type 8 TECT2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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Related Diseases

- [Joubert syndrome](#)
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Related Gene Panels

- [Ataxia \(348 genes\) - ULB](#)

- 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
- Epilepsy gene panel - VUB
- Heterotaxie PCD - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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