

ANALYTE:
TCTN1

NAME:	tectonic family member 1
SYMBOL:	TCTN1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FLJ21127 JBTS13 TECT1
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
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- [Nephrogenetics / Nephropathy \(gene panel\)](#)
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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, 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)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Malformations of cortical development (235 genes) - VUB
- Nephropathy panel - UGent
- Skeletal dysplasia (genepanel) - UZA

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