

ANALYTE:
TCTN3

NAME:	tectonic family member 3
SYMBOL:	TCTN3
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SYNONYMS:	DKFZP564D116 JBTS18 TECT3
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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- [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](#)
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- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [cleft lip with/without cleft palate \(virtual gene panel\)](#)

Related Diseases

- [Joubert syndrome](#)
- [Meckel syndrome](#)
- [Orofaciodigital syndrome type 4](#)
- [Orofaciodigital syndrome type 6](#)

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](#)
- [Cleft lip and palate / dysmorphic facial features / craniofacial anomalies \(255 genes\)\) - UCL](#)
- [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](#)
- [Nephropathy panel - UGent](#)
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- [Skeletal dysplasia \(394 genes\) - VUB](#)
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