

ANALYTE:
EVC2

NAME:	EvC ciliary complex subunit 2
SYMBOL:	EVC2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	LBN limbin
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

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- [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](#)
- [Congenital structural heart defects \(gene panel\)](#)
- [Dermatogenetic panel, severe, rare and hereditary genodermatoses \(gene panel - 394 genes\)](#)
- [Ellis-van Creveld syndrome \(2 genes\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Primary ciliary dyskinesia \(PCD\) Heterotaxyies \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
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- [Skin disorders \(gene panel\)](#)

Related Diseases

- [Acrofacial dysostosis, Weyers type](#)
- [Ellis Van Creveld syndrome](#)

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
- Congenital structural heart defects - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Ellis-van Creveld (2 genes) - IPG
- Heterotaxie PCD - UGent
- Intellectual disability (gene panel)
- Nephropathy panel - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent

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