

ANALYTE:
EYA1

NAME:	EYA transcriptional coactivator and phosphatase 1
SYMBOL:	EYA1
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- [Cataract \(gene panel\)](#)
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- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Hearing loss \(deafness\), \(gene panel\)](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Microphthalmia / Anophthalmia / Coloboma-Anterior Segment Dysgenesis \(MAC-ASD\) \(gene panel\)](#)
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- [Nephropathies, hereditary \(gene panel\)](#)
- [Renal or urinary tract malformation \(CAKUT\) \(gene panel\)](#)
- [cleft lip with/without cleft palate \(virtual gene panel\)](#)

Related Diseases

- [BOR syndrome](#)
- [Branchiootic syndrome](#)
- [Otofaciocervical syndrome](#)

Related Gene Panels

- [Cakut \(congenital anomalies of the kidney and urinary tract-1\) \(69 genes\) - IPG](#)
- [Cataract - 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
- Hearing loss (deafness) (genepanel) - UZA
- Hearing loss (deafness) syndromic (59 genes) - UZA
- Intellectual disability (gene panel)
- Microphthalmia/Anophthalmia/Coloboma - Anterior Segment Dysgenesis - UGent
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1

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