

**ANALYTE:
ACTG1**

NAME:	actin gamma 1
SYMBOL:	ACTG1
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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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RELATED CONTENT

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- [Congenital structural heart defects \(gene panel\)](#)
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- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Malformations of cortical development \(235 genes\)](#)
- [Microphthalmia / Anophthalmia / Coloboma-Anterior Segment Dysgenesis \(MAC-ASD\) \(gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
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- [Skeletal dysplasia \(gene panel\)](#)
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Related Diseases

- [Baraitser-Winter cerebrofrontofacial syndrome](#)
- [Coloboma of choroid and retina](#)
- [Coloboma of iris](#)
- [Rare autosomal dominant non-syndromic sensorineural deafness type DFNA](#)

Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hearing loss (deafness) (genepanel) - UZA
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Malformations of cortical development (235 genes) - VUB
- Microphthalmia/Anophthalmia/Coloboma - Anterior Segment Dysgenesis - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Rare epilepsy with developmental delay (> 240 genes) - UZA
- Skeletal dysplasia (genepanel) - UZA
- test

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