

ANALYTE:
FGD1

NAME:	FYVE, RhoGEF and PH domain containing 1
SYMBOL:	FGD1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ZFYVE3
XREF(S):	<u>Orphanet</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u> <u>Ensembl</u> <u>Genatlas</u>
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RELATED CONTENT

Related Genetic Tests

- [Aarskog-Scott syndrome](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
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Related Diseases

- [Aarskog-Scott syndrome](#)

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
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Growth retardation/short stature (genepanel) - UZA
- Intellectual Disability (104 genes) - IPG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Short Stature (46 genes) - IPG
- Skeletal dysplasia (genepanel) - UZA

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