

ANALYTE:
ARHGAP31

NAME:	Rho GTPase activating protein 31
SYMBOL:	ARHGAP31
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CDGAP
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/analyte/1435>

RELATED CONTENT

Related Genetic Tests

- [Adams-Oliver syndrome \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Congenital structural heart defects \(gene panel\)](#)
- [Dermatogenetic panel, severe, rare and hereditary genodermatoses \(gene panel - 394 genes\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Malformations of cortical development \(235 genes\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skin disorders \(gene panel\)](#)
- [cleft lip with/without cleft palate \(virtual gene panel\)](#)

Related Diseases

- [Adams-Oliver syndrome](#)

Related Gene Panels

- [Adams-Oliver \(6 genes\) - UZA](#)
- [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
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Malformations of cortical development (235 genes) - VUB
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent

Source URL: <http://gentest.healthdata.be/analyte/1435>