ANALYTE:
RIT1

NAME:	Ras like without CAAX 1
SYMBOL:	RIT1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	GTP-binding protein Roc1 MGC125864 MGC125865 RIBB ROC1 Ric-like, expressed in many tissues
XREF(S):	Orphanet Reactome SwissProt Ensembl Genatlas HGNC OMIM
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (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)
- Hereditary Spastic Paraplegia (94 genes)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Pediatric oncopredisposition (gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- RASopathy (gene panel)
- Short Stature (gene panel)
- Short stature/ Growth retardation/ (gene panel)
- Skin disorders (gene panel)

Related Diseases

- Noonan syndrome

Related Gene Panels

- Cardiomyopathy, hereditary (208 genes) - VUB
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Growth retardation/short stature (genepanel) - UZA
- Hereditary Spastic Paraplegia (94 genes) - KUL
- Heterotaxie PCD - UGent
- 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
- Pediatric oncopredisposition - UGent
- RASopathy - KUL
- Short Stature (46 genes) - IPG
- Skin disorders - UGent
