Belgian Genetic Tests database

ANALYTE:
DDR2

NAME:	discoidin domain receptor tyrosine kinase 2
SYMBOL:	DDR2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	TKT
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital structural heart defects (gene panel)
- Epidermal nevus syndrome (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Maffucci syndrome (gene panel)
- Overgrowth & vascular anomalies (gene panel)
- Primary cardiac arrhythmias (Atrial fibrillation / Brugada syndome / Catech. polymorphic ventricular tachycardia / Early repolaristion syndrome / Ideopathic ventricular fibrillation / Long QT syndrome / Sick sinus syndrome / Short QT syndrome) (gene pane)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Sturge-Weber syndrome (gene panel)

Related Diseases

- Spondyloepimetaphyseal dysplasia-short limb-abnormal calcification syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital structural heart defects - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)

- Maffucci syndrome (65 genes) - KUL
- Overgrowth & vascular anomalies (65 genes) - KUL
- Primary cardiac arrhythmias (113 genes) - VUB
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Sturge-Weber syndrome (65 genes) - KUL
- epidermal nevus syndrome (65 genes) - KUL

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