Belgian Genetic Tests database

ANALYTE:
CDKN1C

NAME:	cyclin dependent kinase inhibitor 1C
SYMBOL:	CDKN1C
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	KIP2 P57
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt Reactome
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Overgrowth (gene panel)
- Pediatric oncopredisposition (gene panel)
- Short stature/ Growth retardation/ (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skin disorders (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Beckwith-Wiedemann syndrome due to CDKN1C mutation
- IMAGe syndrome
- Intrauterine growth restriction-short stature-early adult-onset diabetes syndrome
- Silver-Russell syndrome due to a point mutation

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
- Growth retardation/short stature (genepanel) - UZA
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Overgrowth (24 genes) - IPG
- Pediatric oncopredisposition - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent
