ANALYTE:
PCNT

NAME:	pericentrin
SYMBOL:	PCNT
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	KEN KIAA0402 PCN PCNTB SCKL4 Seckel syndrome 4 kendrin
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Brain malformations (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Lipodystrophy and/or hyperinsulinism (gene panel)
- Malformations of cortical development (235 genes)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Stroke (gene panel)

Related Diseases

- Microcephalic osteodysplastic primordial dwarfism type II
- Seckel syndrome

Related Gene Panels

- Brain malformations (34 genes) - ULB
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Lipodystrophy and/or hyperinsulinism (30 genes) - IPG
- Malformations of cortical development (235 genes) - VUB
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Stroke - UGent
