ANALYTE:
MYCN

NAME:	MYCN proto-oncogene, bHLH transcription factor
SYMBOL:	MYCN
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	MYCNOT N-myc bHLHe37
XREF(S):	Orphanet Reactome OMIM SwissProt Ensembl Genatlas HGNC
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Feingold syndrome
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual 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)

Related Diseases

- Feingold syndrome type 1
- Hereditary retinoblastoma
- Neuroblastoma
- Non-hereditary retinoblastoma

Related Gene Panels

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

- Intellectual disability/Epilepsy (1091 genes) - ULG
- 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
