ANALYTE:
POLR3B

NAME:	RNA polymerase III subunit B
SYMBOL:	POLR3B
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	C128 FLJ10388 RPC2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Ataxia Spasticity (gene panel)
- Cerebral palsy (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Hereditary spastic paraplegia (gene panel - 249 genes)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Neuromuscular disorders (548 genes)
- Peripheral neuropathy (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Endosteal sclerosis-cerebellar hypoplasia syndrome
- Hypomyelination-hypogonadotropic hypogonadism-hypodontia syndrome

Related Gene Panels

- Ataxia (348 genes) - ULB
- Ataxia Spasticity - UGent
- Cerebral palsy (212 genes) - UZA
- Congenital malformation (1721 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hereditary spastic paraplegia (188 genes) - ULB
- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Leukodystrophy - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Neuromuscular disorders (548 genes) - ULB
- Neuropathy (genepanel) - UZA
- Skeletal dysplasia (genepanel) - UZA
