Belgian Genetic Tests database

ANALYTE:
SBDS

NAME:	SBDS ribosome maturation factor
SYMBOL:	SBDS
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CGI-97 FLJ10917 SDO1 SDS SWDS
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ciliopathy (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Hepatology (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- Neuromuscular disorders (gene panel) (= myopathy, metabolic myopathy, ion channel muscle diseases, muscular dystrophy, myotonic dystrophy, rhabdomyolysis, myasthenia)
- Pediatric oncopredisposition (gene panel)
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Swachman-Bodian-Diamond syndrome
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Idiopathic aplastic anemia
- SBDS-related severe neonatal spondylometaphyseal dysplasia
- Shwachman-Diamond syndrome

Related Gene Panels

- Ciliopathy - UGent
- Congenital malformation (1721 genes) - ULB
- Hematologic Familiar Forms - ULG
- Hepatology panel - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability (gene panel) - UZA
- Neuromuscular disorders (261 genes) - KUL
- Pediatric oncopredisposition - UGent
- Primary immune deficiencies (444 genes) - KUL
- Primary immune deficiencies - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
