

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):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>SwissProt</u>
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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 \(virtual gene panel\)](#)
- [Neuromuscular disorders \(232 genes \(= myopathy, metabolic myopathy, ion channel muscle diseases, muscular dystrophy, myotonic dystrophy, rhabdomyolysis, myasthenia\)\)](#)
- [Pediatric oncopredisposition \(gene panel\)](#)
- [Primary immune deficiencies \(gene panel\)](#)
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- [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 (120 genes) - UGent
- Congenital malformation (1721 genes) - ULB
- Hematologic Familiar Forms - ULG
- Hepatology panel - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Neuromuscular disorders (232 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

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