

**ANALYTE:**  
**RPS19**

<b>NAME:</b>	ribosomal protein S19
<b>SYMBOL:</b>	RPS19
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	DBA Diamond-Blackfan anemia S19 eS19
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">Ensembl</a> <a href="#">Genatlas</a> <a href="#">HGNC</a> <a href="#">OMIM</a> <a href="#">Reactome</a> <a href="#">SwissProt</a>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	22 Jun 2023 - 16:14

## RELATED CONTENT

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### Related Genetic Tests

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation \(gene panel\)](#)
- [Congenital structural heart defects \(gene panel\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Myeloid/lymphoid neoplasms with germline predisposition \(gene panel\)](#)
- [Pediatric oncopredisposition \(gene panel\)](#)
- [cleft lip with/without cleft palate \(virtual gene panel\)](#)
- [« Inherited bone marrow failures syndromes » with or without organ dysfunction](#)

### Related Diseases

- [Diamond-Blackfan anemia](#)

### Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Hematologic Familiar Forms - ULG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Pediatric oncopredisposition - UGent

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