

ANALYTE:
BBS1

NAME:	Bardet-Biedl syndrome 1
SYMBOL:	BBS1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FLJ23590
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
CREATED:	13 May 2019 - 01:01
CHANGED:	26 Oct 2023 - 23:49

Source URL: <http://gentest.healthdata.be/analyte/2205>

RELATED CONTENT

Related Genetic Tests

- [Ataxia \(autosomic dominant and recessive / except expansion of triplets\) \(gene panel - 722 genes\)](#)
- [Ciliopathy \(gene panel\)](#)
- [Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophthisis / Bardet-Biedl syndromes and kidney cancers \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Early-onset severe obesity](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)
- [Primary ciliary dyskinesia \(PCD\) Heterotaxyies \(gene panel\)](#)
- [Retinal dystrophy / RETNET \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)

Related Diseases

- [Bardet-Biedl syndrome](#)
- [Retinitis pigmentosa](#)

Related Gene Panels

- [Ataxia \(348 genes\) - ULB](#)
- [Ciliopathy \(120 genes\) - UGent](#)
- [Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophthisis, Bardet-Biedl syndromes and kidney cancers \(146 genes\) - IPG](#)
- [Congenital malformation \(1721 genes\) - ULB](#)
- [Congenital malformation gene panel - VUB](#)
- [Early-onset severe obesity \(44 genes\) - ULG](#)
- [Heterotaxie PCD - UGent](#)
- [Intellectual disability & Epilepsy - UGent](#)
- [Intellectual disability \(gene panel\)](#)
- [Intellectual disability/Epilepsy \(1091 genes\) - ULG](#)
- [Nephropathy panel - UGent](#)
- [Neurodevelopmental disorders \(1300 genes\) - ULB](#)
- [Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders \(1162 genes\) - VUB](#)
- [Panel Nephro-ULG-V1](#)
- [Retinal dystrophy - UGent](#)
- [Skeletal dysplasia \(genepanel\) - UZA](#)
- [Skeletal dysplasia - UGent](#)

Source URL: <http://gentest.healthdata.be/analyte/2205>