Belgian Genetic Tests database

ANALYTE:
OBSL1

NAME:	obscurin like cytoskeletal adaptor 1
SYMBOL:	OBSL1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	KIAA0657
XREF(S):	Orphanet Reactome Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/1909

RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Intellectual disability (virtual gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Short Stature (gene panel)
- Short stature/ Growth retardation/ (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- 3M syndrome

Related Gene Panels

- Cardiomyopathy, hereditary (208 genes) - VUB
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Growth retardation/short stature (genepanel) - UZA
- Intellectual disability (gene panel)
- Neurodevelopmental disorders (1300 genes) - ULB

- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Short Stature (46 genes) - IPG
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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