Belgian Genetic Tests database

ANALYTE:
SH3PXD2B

NAME:	SH3 and PX domains 2B
SYMBOL:	SH3PXD2B
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FLJ20831
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Glaucoma (gene panel)
- Intellectual disability (virtual gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Frank-Ter Haar syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Glaucoma - UGent
- Intellectual disability (gene panel)
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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