

ANALYTE:
B3GALT6

NAME:	beta-1,3-galactosyltransferase 6
SYMBOL:	B3GALT6
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	beta-1,3-galactosyltransferase-6 beta3GalT6
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	26 Oct 2023 - 23:49

Source URL: <http://gentest.healthdata.be/analyte/3445>

RELATED CONTENT

Related Genetic Tests

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Dermatogenetic panel, severe, rare and hereditary genodermatoses \(gene panel - 394 genes\)](#)
- [Ehlers-Danlos syndroom, EDS \(gene panel\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders \(213 genes\)](#)
- [Osteogenesis imperfecta / Osteoporose \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [cleft lip with/without cleft palate \(virtual gene panel\)](#)

Related Diseases

- [B3GALT6-related spondylodysplastic Ehlers-Danlos syndrome](#)
- [B4GALT7-related spondylodysplastic Ehlers-Danlos syndrome](#)
- [Eye defects-arachnodactyly-cardiopathy syndrome](#)
- [Spondyloepimetaphyseal dysplasia with joint laxity](#)
- [Spondyloepimetaphyseal dysplasia with joint laxity, Beighton type](#)

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
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Ehlers-Danlos syndrome -UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Metabolic disorders (213 genes) - VUB
- Osteogenesis imperfecta (18 genes) - UGent
- Osteogenesis imperfecta and Osteoporosis (43 genes) - UGent
- Recessive Ehlers-Danlos Syndrome (11 genes) - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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