Belgian Genetic Tests database

ANALYTE:
COL11A2

NAME:	collagen type XI alpha 2 chain
SYMBOL:	COL11A2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HKE5
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Hearing loss (deafness), (gene panel)
- Intellectual disability (virtual gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Stickler syndrome (gene panel)
- Stickler syndrome (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Autosomal dominant otospondylomegaepiphyseal dysplasia
- Fibrochondrogenesis
- Otospondylomegaepiphyseal dysplasia
- Rare autosomal dominant non-syndromic sensorineural deafness type DFNA
- Rare autosomal recessive non-syndromic sensorineural deafness type DFNB
- Weissenbacher- Zweymuller syndrome

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
- Hearing loss (deafness) (genepanel) - UZA
- Hearing loss (deafness) syndromic (59 genes) - UZA
- Intellectual disability (gene panel)
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Stickler (6 genes) - KUL
- Stickler syndrome - UGent
