

ANALYTE:
COL1A2

NAME:	collagen type I alpha 2 chain
SYMBOL:	COL1A2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	alpha 2(I)-collagen alpha-2 collagen type I collagen I, alpha-2 polypeptide collagen of skin, tendon and bone, alpha-2 chain type I procollagen
XREF(S):	Orphanet OMIM Reactome SwissProt Ensembl Genatlas HGNC
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RELATED CONTENT

Related Genetic Tests

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Congenital structural heart defects \(gene panel\)](#)
- [Dermatogenetic panel, severe, rare and hereditary genodermatoses \(gene panel - 394 genes\)](#)
- [Ehlers-Danlos syndroom, EDS \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)
- [Osteogenesis Imperfecta \(gene panel\)](#)
- [Osteogenesis imperfecta / Osteoporose \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
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Related Diseases

- Arthrochalasia Ehlers-Danlos syndrome
- Cardiac-valvular Ehlers-Danlos syndrome
- Ehlers-Danlos/osteogenesis imperfecta syndrome
- High bone mass osteogenesis imperfecta
- Osteogenesis imperfecta type 1
- Osteogenesis imperfecta type 2
- Osteogenesis imperfecta type 3
- Osteogenesis imperfecta type 4

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Ehlers-Danlos syndrome -UGent
- Intellectual disability (gene panel)
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Osteogenesis Imperfecta (25 genes) - KUL
- Osteogenesis imperfecta and Osteoporosis (43 genes) - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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