

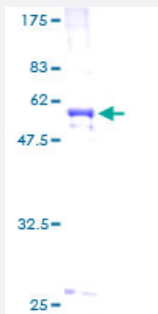
Full-Length

COL2A1 (Human) Recombinant Protein (P01)

Catalog # H00001280-P01

Size 25 ug, 10 ug

Applications



Specification

Product Description

Human COL2A1 full-length ORF (AAH07252, 1 a.a. - 268 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence

MSAFAGLGPREKGPDPDPLQYMRADQAAGGLRQHDAEVDATLKSLNNQIESIRSPEGSRKNPARTC
RDLKLCHPEWKSGDYWIDPNQGCTLDAMKVFCNMETGETCVYPNPANVPKKNWWSSKSKEKK
HIWFGETINGGFHFSYGDDNLAPNTANVQMTFLRLSTEGSQNITYHCKNSIAYLDEAAGNLKKALLI
QGSNDVEIRAEGNSRFTYTALKDGCTKHTGKWGKTVIEYRSQKTSRLPIIDIAPMDIGGPEQEFQVD
IGPVCFL

Host

Wheat Germ (in vitro)

Theoretical MW (kDa)

55.22

Interspecies Antigen Sequence

Mouse (94); Rat (94)

Preparation Method

[in vitro wheat germ expression system](#)

Purification

Glutathione Sepharose 4 Fast Flow

Quality Control Testing

12.5% SDS-PAGE Stained with Coomassie Blue.

Storage Buffer

50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.

Storage Instruction

Store at -80°C. Aliquot to avoid repeated freezing and thawing.

Note

Best use within three months from the date of receipt of this protein.

Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — COL2A1

Entrez GeneID[1280](#)**GeneBank Accession#**[BC007252](#)**Protein Accession#**[AAH07252](#)**Gene Name**

COL2A1

Gene Alias

ANFH, AOM, COL11A3, MGC131516, SEDC

Gene Description

collagen, type II, alpha 1

Omim ID[108300](#) [120140](#) [132450](#) [151210](#) [156550](#) [165720](#) [183900](#) [184250](#) [184252](#) [200610](#) [215150](#) [271700](#) [608805](#)**Gene Ontology**[Hyperlink](#)**Gene Summary**

This gene encodes the alpha-1 chain of type II collagen, a fibrillar collagen found in cartilage and the vitreous humor of the eye. Mutations in this gene are associated with achondrogenesis, chondrodysplasia, early onset familial osteoarthritis, SED congenita, Langer-Saldino achondrogenesis, Kniest dysplasia, Stickler syndrome type I, and spondyloepimetaphyseal dysplasia Strudwick type. In addition, defects in processing chondrocalcin, a calcium binding protein that is the C-propeptide of this collagen molecule, are also associated with chondrodysplasia. There are two transcripts identified for this gene. [provided by RefSeq]

Other Designations

OTTHUMP00000195063|cartilage collagen|chondrocalcin|collagen II, alpha-1 polypeptide

Pathway

- [ECM-receptor interaction](#)
- [Focal adhesion](#)

Disease

- [Asthma](#)
- [Cardiovascular Diseases](#)
- [Cleft Lip](#)
- [Cleft Palate](#)
- [Cumulative Trauma Disorders](#)
- [Diabetes Mellitus](#)
- [Edema](#)
- [Gaucher disease](#)
- [Genetic Predisposition to Disease](#)
- [Hip Dislocation](#)
- [Intervertebral Disk Displacement](#)
- [Legg-Perthes Disease](#)
- [Myopia](#)
- [Occupational Diseases](#)
- [Osteoarthritis](#)
- [Thrombophilia](#)
- [Toxoplasmosis](#)