

# COL1A2 Knockout 293T Cell Lysate, Homozygous

**Catalog No.: RM50018**

## Basic Information

**Catalog No.**

RM50018

**Category**

Cell Lysate

**Parental Cell line**

293T

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

COL1A2

**Species**

Human

**Gene ID**

1278

**Swiss Prot**

P08123

**Synonyms**OI4; EDSCV; EDSARTH2; Collagen  
I/COL1A2

## Contact

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## Background

This gene encodes the pro-alpha2 chain of type I collagen whose triple helix comprises two alpha1 chains and one alpha2 chain. Type I is a fibril-forming collagen found in most connective tissues and is abundant in bone, cornea, dermis and tendon. Mutations in this gene are associated with osteogenesis imperfecta types I-IV, Ehlers-Danlos syndrome type VIIB, recessive Ehlers-Danlos syndrome Classical type, idiopathic osteoporosis, and atypical Marfan syndrome. Symptoms associated with mutations in this gene, however, tend to be less severe than mutations in the gene for the alpha1 chain of type I collagen (COL1A1) reflecting the different role of alpha2 chains in matrix integrity. Three transcripts, resulting from the use of alternate polyadenylation signals, have been identified for this gene.

## Product Information

**Description**

COL1A2 Knockout cell line is engineered from 293T cell line with Gene-Editing Technology.

Allele-1:exon1 was deleted

Allele-2:exon1 was deleted

Mammalian cells such as human, rat and mouse cells are normally diploid with two alleles. Homozygote: both alleles were knocked out, mRNA has no signal, no expression of proteins. Heterozygote: only one allele was knocked out, the mRNA transcript levels was decreased compared to wild type, and the protein expression levels was also lower than that of the wild type.

**Packaging**

1 vial parental cell Lysate and 1 vial knockout cell Lysate

**Shipping Conditions**

4°C

**Amount**

50µL, 2µg/µL.

**Storage**

Lysate is stable for 12 months when stored at -20°C. Minimizing freeze-thaw cycles.

**Protocol**

To be used as WB control. Lysate is supplied in 1× SDS sample buffer (2% SDS, 60 mM Tris-HCl pH 6.8, 10% Glycerol, 0.02% Bromophenol blue, 60 mM beta-mercaptoethanol). Lysate should be boiled for 3 - 5 minutes before loading onto gel.

## Sequencing data

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WT GGGGCTCTGCGAC\*\*\*\*\*TAAGTGCCTTCAGC  
Mut GGGGCTCTGCGAC\*\*\*Deletion\*\*\*TAAGTGCCTTCAGC  
Allele-1: exon1 was deleted

Genome sequence analysis of PCR products from parental (WT) and COL1A2 knockout (KO) 293T cells, using sanger sequencing.

WT GGGGCTCTGCGAC\*\*\*\*\*TAAGTGCCTTCAGC  
Mut GGGGCTCTGCGAC\*\*\*Deletion\*\*\*TAAGTGCCTTCAGC  
Allele-2: exon1 was deleted