

## Product datasheet for RC206234L1V

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## Collagen I (COL1A1) (NM\_000088) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: Collagen I (COL1A1) (NM\_000088) Human Tagged ORF Clone Lentiviral Particle

Symbol: Collagen I

Synonyms: CAFYD; EDSARTH1; EDSC; OI1; OI2; OI3; OI4

**Mammalian Cell** 

Selection:

None

Vector:

pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK

ACCN: NM\_000088

ORF Size: 4392 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC206234).

Sequence:

Cytogenetics:

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through

naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

**OTI Annotation:** This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

**RefSeg:** NM 000088.3, NP 000079.2

 RefSeq Size:
 5927 bp

 RefSeq ORF:
 4395 bp

 Locus ID:
 1277

 UniProt ID:
 P02452

**Domains:** COLFI, VWC, Collagen

17q21.33

**Protein Families:** Druggable Genome





**Protein Pathways:** ECM-receptor interaction, Focal adhesion

**MW:** 139.01 kDa

**Gene Summary:** This gene encodes the pro-alpha1 chains 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 VIIA, Ehlers-Danlos syndrome Classical type, Caffey Disease and idiopathic osteoporosis. Reciprocal translocations between chromosomes 17 and 22, where this gene and the gene for platelet-derived growth factor beta are located, are associated with a particular type of skin tumor called dermatofibrosarcoma protuberans, resulting from unregulated expression of the growth factor. Two transcripts, resulting from the use of alternate polyadenylation

signals, have been identified for this gene. [provided by R. Dalgleish, Feb 2008]