

Product datasheet for RC218704L3V

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 VI (COL6A2) (NM 058174) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Collagen VI (COL6A2) (NM_058174) Human Tagged ORF Clone Lentiviral Particle

Symbol: Collagen VI

Synonyms: BTHLM1; PP3610; UCMD1

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK
ACCN: NM 058174

ORF Size: 2754 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC218704).

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 058174.1, NP 478054.1

 RefSeq Size:
 3145 bp

 RefSeq ORF:
 2757 bp

 Locus ID:
 1292

 UniProt ID:
 P12110

Protein Families: Secreted Protein

Protein Pathways: ECM-receptor interaction, Focal adhesion

21q22.3





MW: 95.3 kDa

Gene Summary:

This gene encodes one of the three alpha chains of type VI collagen, a beaded filament collagen found in most connective tissues. The product of this gene contains several domains similar to von Willebrand Factor type A domains. These domains have been shown to bind extracellular matrix proteins, an interaction that explains the importance of this collagen in organizing matrix components. Mutations in this gene are associated with Bethlem myopathy and Ullrich scleroatonic muscular dystrophy. Three transcript variants have been identified for this gene. [provided by RefSeq, Jul 2008]