

Product datasheet for RC211080L2V

OriGene Technologies, Inc.

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COMP (NM_000095) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: COMP (NM 000095) Human Tagged ORF Clone Lentiviral Particle

Symbol: COMP

Synonyms: CTS2; EDM1; EPD1; MED; PSACH; THBS5; TSP5

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_000095 **ORF Size:** 2271 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC211080).

OTI Disclaimer:

Sequence:

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 000095.2

 RefSeq Size:
 2471 bp

 RefSeq ORF:
 2274 bp

 Locus ID:
 1311

 UniProt ID:
 P49747

 Cytogenetics:
 19p13.11

Domains: EGF_CA, tsp_3, EGF, EGF

Protein Families: Druggable Genome, Secreted Protein





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Protein Pathways: ECM-receptor interaction, Focal adhesion, TGF-beta signaling pathway

MW: 82.9 kDa

Gene Summary: The protein encoded by this gene is a noncollagenous extracellular matrix (ECM) protein. It

consists of five identical glycoprotein subunits, each with EGF-like and calcium-binding (thrombospondin-like) domains. Oligomerization results from formation of a five-stranded coiled coil and disulfides. Binding to other ECM proteins such as collagen appears to depend on divalent cations. Contraction or expansion of a 5 aa aspartate repeat and other mutations can cause pseudochondroplasia (PSACH) and multiple epiphyseal dysplasia (MED). [provided

by RefSeq, Jul 2016]