

## Product datasheet for RC205901L2V

## OriGene Technologies, Inc.

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## **GDF3 (NM\_020634) Human Tagged ORF Clone Lentiviral Particle**

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: GDF3 (NM 020634) Human Tagged ORF Clone Lentiviral Particle

Symbol: GDF3

Synonyms: KFS3; MCOP7; MCOPCB6

Mammalian Cell

Selection:

None

**Vector:** pLenti-C-mGFP (PS100071)

Tag: mGFP

**ACCN:** NM\_020634 **ORF Size:** 1092 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC205901).

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.

**RefSeq:** <u>NM 020634.1</u>

 RefSeq Size:
 1224 bp

 RefSeq ORF:
 1095 bp

 Locus ID:
 9573

 UniProt ID:
 Q9NR23

 Cytogenetics:
 12p13.31

**Protein Families:** Adult stem cells, Cancer stem cells, Druggable Genome, Embryonic stem cells, ES Cell

Differentiation/IPS, Induced pluripotent stem cells, Secreted Protein, Stem cell relevant

signaling - TGFb/BMP signaling pathway





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**MW:** 41.5 kDa

**Gene Summary:** This gene encodes a secreted ligand of the TGF-beta (transforming growth factor-beta)

superfamily of proteins. Ligands of this family bind various TGF-beta receptors leading to recruitment and activation of SMAD family transcription factors that regulate gene

expression. The encoded preproprotein is proteolytically processed to generate each subunit of the disulfide-linked homodimer. This protein plays a role ocular and skeletal development.

Mutations in this gene are associated with microphthalmia, coloboma, and skeletal

abnormalities in human patients. [provided by RefSeq, Aug 2016]