

Product datasheet for RC206587L4V

OriGene Technologies, Inc.

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PDGF beta (PDGFB) (NM_002608) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: PDGF beta (PDGFB) (NM_002608) Human Tagged ORF Clone Lentiviral Particle

Symbol: PDGF beta

Synonyms: c-sis; IBGC5; PDGF-2; PDGF2; SIS; SSV

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM_002608

ORF Size: 723 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC206587).

Sequence:

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 002608.1

 RefSeq Size:
 3393 bp

 RefSeq ORF:
 726 bp

 Locus ID:
 5155

 UniProt ID:
 P01127

 Cytogenetics:
 22q13.1

Domains: PDGF, PDGF N

Protein Families: Druggable Genome





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Protein Pathways: Cytokine-cytokine receptor interaction, Focal adhesion, Gap junction, Glioma, MAPK signaling

pathway, Melanoma, Pathways in cancer, Prostate cancer, Regulation of actin cytoskeleton,

Renal cell carcinoma

MW: 27.3 kDa

Gene Summary: This gene encodes a member of the protein family comprised of both platelet-derived growth

factors (PDGF) and vascular endothelial growth factors (VEGF). The encoded preproprotein is proteolytically processed to generate platelet-derived growth factor subunit B, which can homodimerize, or alternatively, heterodimerize with the related platelet-derived growth factor subunit A. These proteins bind and activate PDGF receptor tyrosine kinases, which play a role in a wide range of developmental processes. Mutations in this gene are associated with meningioma. Reciprocal translocations between chromosomes 22 and 17, at sites where this

gene and that for collagen type 1, alpha 1 are located, are associated with

dermatofibrosarcoma protuberans, a rare skin tumor. Alternative splicing results in multiple

transcript variants. [provided by RefSeq, Oct 2015]