

Product datasheet for RC221976L2V

OriGene Technologies, Inc.

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CD36 (NM_001001548) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: CD36 (NM_001001548) Human Tagged ORF Clone Lentiviral Particle

Symbol: CD36

Synonyms: BDPLT10; CHDS7; FAT; GP3B; GP4; GPIV; PASIV; SCARB3

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_001001548

ORF Size: 1416 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC221976).

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.

RefSeq: <u>NM 001001548.1</u>

RefSeq Size:4727 bpRefSeq ORF:1419 bp

 Locus ID:
 948

 UniProt ID:
 P16671

 Cytogenetics:
 7q21.11

Protein Families: Druggable Genome, Transmembrane



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Protein Pathways: Adipocytokine signaling pathway, ECM-receptor interaction, Hematopoietic cell lineage, PPAR

signaling pathway

MW: 53.1 kDa

Gene Summary: The protein encoded by this gene is the fourth major glycoprotein of the platelet surface and

serves as a receptor for thrombospondin in platelets and various cell lines. Since

thrombospondins are widely distributed proteins involved in a variety of adhesive processes, this protein may have important functions as a cell adhesion molecule. It binds to collagen, thrombospondin, anionic phospholipids and oxidized LDL. It directly mediates cytoadherence of Plasmodium falciparum parasitized erythrocytes and it binds long chain fatty acids and may function in the transport and/or as a regulator of fatty acid transport. Mutations in this gene cause platelet glycoprotein deficiency. Multiple alternatively spliced transcript variants

have been found for this gene. [provided by RefSeq, Feb 2014]