

Product datasheet for RC210274L1V

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

Caveolin 1 (CAV1) (NM 001753) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Caveolin 1 (CAV1) (NM_001753) Human Tagged ORF Clone Lentiviral Particle

Symbol: Caveolin 1

Synonyms: BSCL3; CGL3; LCCNS; MSTP085; PPH3; VIP21

Mammalian Cell

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK
ACCN: NM 001753

ORF Size: 534 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC210274).

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 001753.3

 RefSeq Size:
 2704 bp

 RefSeq ORF:
 537 bp

 Locus ID:
 857

 UniProt ID:
 Q03135

Cytogenetics: 7q31.2

Domains: Caveolin

Protein Families: Druggable Genome, Transmembrane





Protein Pathways: Focal adhesion, Viral myocarditis

MW: 20.3 kDa

Gene Summary: The scaffolding protein encoded by this gene is the main component of the caveolae plasma

membranes found in most cell types. The protein links integrin subunits to the tyrosine kinase FYN, an initiating step in coupling integrins to the Ras-ERK pathway and promoting cell cycle progression. The gene is a tumor suppressor gene candidate and a negative regulator of the Ras-p42/44 mitogen-activated kinase cascade. Caveolin 1 and caveolin 2 are located next to each other on chromosome 7 and express colocalizing proteins that form a stable hetero-oligomeric complex. Mutations in this gene have been associated with Berardinelli-Seip congenital lipodystrophy. Alternatively spliced transcripts encode alpha and beta

isoforms of caveolin 1.[provided by RefSeq, Mar 2010]