

Product datasheet for RC204576L1V

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

Vinculin (VCL) (NM_003373) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Vinculin (VCL) (NM_003373) Human Tagged ORF Clone Lentiviral Particle

Symbol: Vinculin

Synonyms: CMD1W; CMH15; HEL114; MV; MVCL

NM 003373

Mammalian Cell

Selection:

ACCN:

None

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

Tag: Myc-DDK

ORF Size: 3198 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC204576).

•

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 003373.3

RefSeq Size: 5443 bp
RefSeq ORF: 3201 bp
Locus ID: 7414
UniProt ID: P18206

Cytogenetics: 10q22.2

Domains: Vinculin

Protein Families: Druggable Genome





Vinculin (VCL) (NM_003373) Human Tagged ORF Clone Lentiviral Particle - RC204576L1V

Protein Pathways: Adherens junction, Focal adhesion, Leukocyte transendothelial migration, Regulation of actin

cytoskeleton

MW: 116.7 kDa

Gene Summary: Vinculin is a cytoskeletal protein associated with cell-cell and cell-matrix junctions, where it is

thought to function as one of several interacting proteins involved in anchoring F-actin to the membrane. Defects in VCL are the cause of cardiomyopathy dilated type 1W. Dilated

cardiomyopathy is a disorder characterized by ventricular dilation and impaired systolic function, resulting in congestive heart failure and arrhythmia. Multiple alternatively spliced transcript variants have been found for this gene, but the biological validity of some variants

has not been determined. [provided by RefSeq, Jul 2008]