

Product datasheet for RC224868L1V

OriGene Technologies, Inc.

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OPN1LW (NM_020061) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: OPN1LW (NM_020061) Human Tagged ORF Clone Lentiviral Particle

Symbol: OPN1LW

Synonyms: CBBM; CBP; COD5; RCP; ROP

Mammalian Cell

Selection:

None

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

 Tag:
 Myc-DDK

 ACCN:
 NM_020061

 ORF Size:
 1092 bp

ORF Nucleotide

1032 bp

Sequence:

The ORF insert of this clone is exactly the same as(RC224868).

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 020061.2

RefSeq Size: 1356 bp
RefSeq ORF: 1095 bp
Locus ID: 5956
Cytogenetics: Xq28

Protein Families: Druggable Genome, Transmembrane

MW: 40.4 kDa







Gene Summary:

This gene encodes for a light absorbing visual pigment of the opsin gene family. The encoded protein is called red cone photopigment or long-wavelength sensitive opsin. Opsins are G-protein coupled receptors with seven transmembrane domains, an N-terminal extracellular domain, and a C-terminal cytoplasmic domain. This gene and the medium-wavelength opsin gene are tandemly arrayed on the X chromosome and frequent unequal recombination and gene conversion may occur between these sequences. X chromosomes may have fusions of the medium- and long-wavelength opsin genes or may have more than one copy of these genes. Defects in this gene are the cause of partial, protanopic colorblindness. [provided by RefSeq, Jul 2008]