

OriGene Technologies, Inc.

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Product datasheet for RC206991L2V

CD94 (KLRD1) (NM_002262) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type:	Lentiviral Particles
Product Name:	CD94 (KLRD1) (NM_002262) Human Tagged ORF Clone Lentiviral Particle
Symbol:	CD94
Synonyms:	CD94
Mammalian Cell Selection:	None
Vector:	pLenti-C-mGFP (PS100071)
Tag:	mGFP
ACCN:	NM_002262
ORF Size:	537 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC206991).
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. <u>More info</u>
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 002262.2</u>
RefSeq Size:	3258 bp
RefSeq ORF:	540 bp
Locus ID:	3824
UniProt ID:	<u>Q13241</u>
Cytogenetics:	12p13.2
Protein Families:	Transmembrane



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ORIGENE CD94 (KLRD1) (NM_002262) Human Tagged ORF Clone Lentiviral Particle – RC206991L2V	
Protein Pathways:	Antigen processing and presentation, Graft-versus-host disease, Natural killer cell mediated cytotoxicity
MW:	20.5 kDa
Gene Summary:	Natural killer (NK) cells are a distinct lineage of lymphocytes that mediate cytotoxic activity and secrete cytokines upon immune stimulation. Several genes of the C-type lectin superfamily, including members of the NKG2 family, are expressed by NK cells and may be involved in the regulation of NK cell function. KLRD1 (CD94) is an antigen preferentially expressed on NK cells and is classified as a type II membrane protein because it has an external C terminus. Several transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, May 2017]

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