

#### OriGene Technologies, Inc.

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# Product datasheet for TL311862V

### NKG2C (KLRC2) Human shRNA Lentiviral Particle (Locus ID 3822)

## **Product data:**

Product Type:	shRNA Lentiviral Particles
Product Name:	NKG2C (KLRC2) Human shRNA Lentiviral Particle (Locus ID 3822)
Locus ID:	3822
Synonyms:	CD159c; NKG2-C; NKG2C
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	KLRC2 - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, >10^7 TU/ml.
RefSeq:	<u>NM_002260</u> , <u>NM_002260.1, NM_002260.2, NM_002260.3, BC112039</u> , <u>BC112039.1, BC093644</u> , <u>BC106069</u>
UniProt ID:	<u>P26717</u>
Summary:	Natural killer (NK) cells are lymphocytes that can mediate lysis of certain tumor cells and virus-infected cells without previous activation. They can also regulate specific humoral and cell-mediated immunity. NK cells preferentially express several calcium-dependent (C-type) lectins, which have been implicated in the regulation of NK cell function. The group, designated KLRC (NKG2) are expressed primarily in natural killer (NK) cells and encodes a family of transmembrane proteins characterized by a type II membrane orientation (extracellular C terminus) and the presence of a C-type lectin domain. The KLRC (NKG2) gene family is located within the NK complex, a region that contains several C-type lectin genes preferentially expressed on NK cells. KLRC2 alternative splice variants have been described but their full-length nature has not been determined. [provided by RefSeq, Jul 2008]
shRNA Design:	These shRNA constructs were designed against multiple splice variants at this gene locus. To be certain that your variant of interest is targeted, please contact <u>techsupport@origene.com</u> . If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u> .



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#### STATES ORIGENE NKG2C (KLRC2) Human shRNA Lentiviral Particle (Locus ID 3822) – TL311862V

Performance Guaranteed: OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).

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