

### KLRC1 Antibody (N-term) Blocking peptide Synthetic peptide

Catalog # BP12570a

## Specification

# KLRC1 Antibody (N-term) Blocking peptide - Product Information

Primary Accession

<u>P26715</u>

# KLRC1 Antibody (N-term) Blocking peptide - Additional Information

Gene ID 3821

**Other Names** 

NKG2-A/NKG2-B type II integral membrane protein, CD159 antigen-like family member A, NK cell receptor A, NKG2-A/B-activating NK receptor, CD159a, KLRC1, NKG2A

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions** This product is for research use only. Not for use in diagnostic or therapeutic procedures.

## KLRC1 Antibody (N-term) Blocking peptide - Protein Information

Name KLRC1

## **Synonyms** NKG2A {ECO:0000303|PubMed:18083576}

Function

Immune inhibitory receptor involved in self-nonself discrimination. In complex with KLRD1 on cytotoxic and regulatory lymphocyte subsets, recognizes non-classical major histocompatibility (MHC) class Ib molecule HLA-E loaded with self-peptides derived from the signal sequence of classical MHC class Ia molecules. Enables cytotoxic cells to monitor the expression of MHC class I molecules in healthy cells and to tolerate self (PubMed:<a

href="http://www.uniprot.org/citations/18083576" target="\_blank">18083576</a>, PubMed:<a href="http://www.uniprot.org/citations/37264229" target="\_blank">37264229</a>, PubMed:<a href="http://www.uniprot.org/citations/9430220" target="\_blank">9430220</a>, PubMed:<a href="http://www.uniprot.org/citations/9486650" target="\_blank">9486650</a>). Upon HLA-E-peptide binding, transmits intracellular signals through two immunoreceptor tyrosine-based inhibition motifs (ITIMs) by recruiting INPP5D/SHP-1 and INPPL1/SHP-2 tyrosine phosphatases to ITIMs, and ultimately opposing signals transmitted by activating receptors through dephosphorylation of proximal signaling molecules (PubMed:<a

href="http://www.uniprot.org/citations/12165520" target="\_blank">12165520</a>, PubMed:<a href="http://www.uniprot.org/citations/9485206" target="\_blank">9485206</a>). Key inhibitory



receptor on natural killer (NK) cells that regulates their activation and effector functions (PubMed:<a href="http://www.uniprot.org/citations/30860984" target=" blank">30860984</a>, PubMed:<a href="http://www.uniprot.org/citations/9430220" target=" blank">9430220</a>, PubMed:<a href="http://www.uniprot.org/citations/9485206" target="\_blank">9485206</a>, PubMed:<a href="http://www.uniprot.org/citations/9486650" target=" blank">9486650</a>). Dominantly counteracts T cell receptor signaling on a subset of memory/effector CD8-positive T cells as part of an antigen-driven response to avoid autoimmunity (PubMed:<a href="http://www.uniprot.org/citations/12387742" target=" blank">12387742</a>). On intraepithelial CD8-positive gamma-delta regulatory T cells triggers TGFB1 secretion, which in turn limits the cytotoxic programming of intraepithelial CD8-positive alpha-beta T cells, distinguishing harmless from pathogenic antigens (PubMed:<a href="http://www.uniprot.org/citations/18064301" target=" blank">18064301</a>). In HLA-E-rich tumor microenvironment, acts as an immune inhibitory checkpoint and may contribute to progressive loss of effector functions of NK cells and tumor-specific T cells, a state known as cell exhaustion (PubMed:<a href="http://www.uniprot.org/citations/30503213" target=" blank">30503213</a>, PubMed:<a href="http://www.uniprot.org/citations/30860984" target=" blank">30860984</a>).

## **Cellular Location**

Cell membrane; Single-pass type II membrane protein

#### **Tissue Location**

Predominantly expressed in NK cells (at protein level) (PubMed:20952657, PubMed:9430220, PubMed:9485206). Expressed in intraepithelial CD8-positive T cell subsets with higher frequency in gamma-delta T cells than alpha-beta T cells (at protein level) (PubMed:18064301). Expressed in memory gamma-delta T cells (at protein level) (PubMed:20952657). Restricted to a subset of memory/effector CD8-positive alpha-beta T cells (at protein level) (PubMed:12387742) Expressed in intratumoral NK and CD8-positive T cells (PubMed:30503213). Expressed in melanoma-specific cytotoxic T cell clones (at protein level) (PubMed:9485206). KLRD1-KLRC1 and KLRD1-KLRC2 are differentially expressed in NK and T cell populations, with only minor subsets expressing both receptor complexes (at protein level) (PubMed:20952657).

## KLRC1 Antibody (N-term) Blocking peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

#### <u>Blocking Peptides</u>

## KLRC1 Antibody (N-term) Blocking peptide - Images

## KLRC1 Antibody (N-term) Blocking peptide - Background

Natural killer (NK) cells are lymphocytes that can mediatelysis of certain tumor cells and virus-infected cells withoutprevious activation. They can also regulate specific humoral andcell-mediated immunity. The protein encoded by this gene belongs tothe killer cell lectin-like receptor family, also called NKG2family, which is a group of transmembrane proteins preferentiallyexpressed in NK cells. This family of proteins is characterized bythe type II membrane orientation and the presence of a C-typelectin domain. This protein forms a complex with another familymember, KLRD1/CD94, and has been implicated in the recognition of the MHC class I HLA-E molecules in NK cells. The genes of NKG2family members form a killer cell lectin-like receptor gene clusteron chromosome 12. Four alternatively spliced transcript variantsencoding two distinct isoforms have been observed. [provided byRefSeq].

## KLRC1 Antibody (N-term) Blocking peptide - References

Ucisik-Akkaya, E., et al. Mol. Hum. Reprod. 16(10):770-777(2010)Ma, J., et al. J. Med. Virol. 82(9):1501-1507(2010)Harrison, R.J., et al. Clin. Exp. Immunol. 161(2):306-314(2010)Rose, J.E., et al. Clin. Exp. Immunol. Al. Clin. Exp. Immunol



al. Mol. Med. 16 (7-8), 247-253 (2010) :Beziat, V., et al. PLoS ONE 5 (8), E11966 (2010) :