

**KLRC1 Antibody (N-term) Blocking peptide**  
**Synthetic peptide**  
**Catalog # BP12570a****Specification**

---

**KLRC1 Antibody (N-term) Blocking peptide - Product Information**Primary Accession [P26715](#)**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-nonsel self 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/9486650" target="\_blank">9486650</a>, PubMed:<a href="http://www.uniprot.org/citations/18083576" target="\_blank">18083576</a>, PubMed:<a href="http://www.uniprot.org/citations/9430220" target="\_blank">9430220</a>, PubMed:<a href="http://www.uniprot.org/citations/37264229" target="\_blank">37264229</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/9485206" target="\_blank">9485206</a>, PubMed:<a href="http://www.uniprot.org/citations/12165520" target="\_blank">12165520</a>). Key

inhibitory receptor on natural killer (NK) cells that regulates their activation and effector functions (PubMed:<a href="http://www.uniprot.org/citations/9486650" target="\_blank">9486650</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/30860984" target="\_blank">30860984</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:9430220, PubMed:9485206, PubMed:20952657). 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.

- [Blocking Peptides](#)

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

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

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. The protein encoded by this gene belongs to the killer cell lectin-like receptor family, also called NKG2 family, which is a group of transmembrane proteins preferentially expressed in NK cells. This family of proteins is characterized by the type II membrane orientation and the presence of a C-type lectin domain. This protein forms a complex with another family member, KLRD1/CD94, and has been implicated in the recognition of the MHC class I HLA-E molecules in NK cells. The genes of NKG2 family members form a killer cell lectin-like receptor gene cluster on chromosome 12. Four alternatively spliced transcript variants encoding two distinct isoforms have been observed. [provided by RefSeq].

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