

CD69 Blocking Peptide (Center)

Synthetic peptide Catalog # BP5413c

Specification

CD69 Blocking Peptide (Center) - Product Information

Primary Accession Other Accession NP_001772.1

CD69 Blocking Peptide (Center) - Additional Information

Gene ID 969

Other Names

Early activation antigen CD69, Activation inducer molecule, AIM, BL-AC/P26, C-type lectin domain family 2 member C, EA1, Early T-cell activation antigen p60, GP32/28, Leukocyte surface antigen Leu-23, MLR-3, CD69, CD69, CLEC2C

Target/Specificity

The synthetic peptide sequence is selected from aa 134-148 of HUMAN CD69

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.

CD69 Blocking Peptide (Center) - Protein Information

Name CD69

Synonyms CLEC2C

Function

Transmembrane protein expressed mainly on T-cells resident in mucosa that plays an essential role in immune cell homeostasis. Rapidly expressed on the surface of platelets, T-lymphocytes and NK cells upon activation by various stimuli, such as antigen recognition or cytokine signaling, stimulates different signaling pathways in different cell types (PubMed:24752896, PubMed:14752896, PubMed:14752896, Pub

href="http://www.uniprot.org/citations/26296369" target="_blank">26296369, PubMed:35930205). Negatively regulates Th17 cell differentiation through its carbohydrate dependent interaction with

galectin-1/LGALS1 present on immature dendritic cells (PubMed:24752896). Association



of CD69 cytoplasmic tail with the JAK3/STAT5 signaling pathway regulates the transcription of RORgamma/RORC and, consequently, differentiation toward the Th17 lineage (By similarity). Also acts via the S100A8/S100A9 complex present on peripheral blood mononuclear cells to promote the conversion of naive CD4 T-cells into regulatory T-cells (PubMed:26296369). Acts as an oxidized low-density lipoprotein (oxLDL) receptor in CD4 T- lymphocytes and negatively regulates the inflammatory response by inducing the expression of PDCD1 through the activation of NFAT (PubMed:35930205). Participates in adipose tissue-derived mesenchymal stem cells (ASCs)-mediated protection against P.aeruginosa infection. Mechanistically, specifically recognizes P.aeruginosa to promote ERK1 activation, followed by granulocyte-macrophage colony-stimulating factor (GM-CSF) and other inflammatory cytokines secretion (PubMed:34841721). In eosinophils, induces IL-10 production through the ERK1/2 pathway (By similarity). Negatively regulates the chemotactic responses of effector lymphocytes and dendritic cells (DCs) to sphingosine 1 phosphate/S1P by acting as a S1PR1 receptor agonist and facilitating the internalization and degradation of the receptor (PubMed: 37039481).

Cellular Location

Cell membrane; Single-pass type II membrane protein

Tissue Location

Expressed on the surface of activated T-cells, B- cells, natural killer cells, neutrophils, eosinophils, epidermal Langerhans cells and platelets

CD69 Blocking Peptide (Center) - Protocols

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

• Blocking Peptides

CD69 Blocking Peptide (Center) - Images

CD69 Blocking Peptide (Center) - Background

This gene encodes a member of the calcium dependent lectin superfamily of type II transmembrane receptors. Expression of the encoded protein is induced upon activation of T lymphocytes, and may play a role in proliferation. Furthermore, the protein may act to transmit signals in natural killer cells and platelets. Alternative splicing results in multiple transcript variants.

CD69 Blocking Peptide (Center) - References

Davila, S., et al. Genes Immun. (2010) In press:

Kolenko, P., et al. Acta Crystallogr. Sect. F Struct. Biol. Cryst. Commun. 65 (PT 12), 1258-1260 (2009) :

Hu, M., et al. J. Mol. Recognit. 22(6):516-520(2009)

Radstake, T.R., et al. PLoS ONE 4 (6), E5981 (2009) :

Natarajan, K., et al. Biochemistry 39(48):14779-14786(2000)

Vance, B.A., et al. Arch. Biochem. Biophys. 368(2):214-220(1999)

Lopez-Cabrera, M., et al. J. Biol. Chem. 270(37):21545-21551(1995)

Bezouska, K., et al. Biochem. Biophys. Res. Commun. 208(1):68-74(1995)

Santis, A.G., et al. Eur. J. Immunol. 24(7):1692-1697(1994)

Lopez-Cabrera, M., et al. J. Exp. Med. 178(2):537-547(1993)

Ziegler, S.F., et al. Eur. J. Immunol. 23(7):1643-1648(1993)





Cambiaggi, C., et al. Immunogenetics 36(2):117-120(1992)