

LIM Kinase 2B (LIMK2B) Antibody (N-term) Blocking peptide
Synthetic peptide
Catalog # BP7814a**Specification**

LIM Kinase 2B (LIMK2B) Antibody (N-term) Blocking peptide - Product Information

Primary Accession [P53671](#)
Other Accession [Q99464](#)

LIM Kinase 2B (LIMK2B) Antibody (N-term) Blocking peptide - Additional Information

Gene ID 3985

Other Names

LIM domain kinase 2, LIMK-2, LIMK2

Target/Specificity

The synthetic peptide sequence used to generate the antibody [AP7814a](/product/products/AP7814a) was selected from the N-term region of human LIMK2B. A 10 to 100 fold molar excess to antibody is recommended. Precise conditions should be optimized for a particular assay.

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.

LIM Kinase 2B (LIMK2B) Antibody (N-term) Blocking peptide - Protein Information

Name LIMK2

Function

Serine/threonine-protein kinase that plays an essential role in the regulation of actin filament dynamics (PubMed: [10436159](http://www.uniprot.org/citations/10436159), PubMed: [11018042](http://www.uniprot.org/citations/11018042)). Acts downstream of several Rho family GTPase signal transduction pathways (PubMed: [10436159](http://www.uniprot.org/citations/10436159), PubMed: [11018042](http://www.uniprot.org/citations/11018042)). Involved in astral microtubule organization and mitotic spindle orientation during early stages of mitosis by mediating phosphorylation of TPPP (PubMed: [22328514](http://www.uniprot.org/citations/22328514)). Displays serine/threonine-specific phosphorylation of myelin basic protein and histone (MBP) in vitro (PubMed: [8537403](http://www.uniprot.org/citations/8537403)).

Suppresses ciliogenesis via multiple pathways; phosphorylation of CFL1, suppression of directional trafficking of ciliary vesicles to the ciliary base, and by facilitating YAP1 nuclear localization where it acts as a transcriptional corepressor of the TEAD4 target genes AURKA and PLK1 (PubMed:25849865).

Cellular Location

Cytoplasm, cytoskeleton, spindle. Cytoplasm, cytoskeleton, microtubule organizing center, centrosome [Isoform LIMK2b]: Cytoplasm. Cytoplasm, perinuclear region. Nucleus Note=Mainly present in the cytoplasm and is scarcely translocated to the nucleus.

LIM Kinase 2B (LIMK2B) Antibody (N-term) Blocking peptide - Protocols

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

- [Blocking Peptides](#)

LIM Kinase 2B (LIMK2B) Antibody (N-term) Blocking peptide - Images**LIM Kinase 2B (LIMK2B) Antibody (N-term) Blocking peptide - Background**

There are approximately 40 known eukaryotic LIM proteins, so named for the LIM domains they contain. LIM domains are highly conserved cysteine-rich structures containing 2 zinc fingers. Although zinc fingers usually function by binding to DNA or RNA, the LIM motif probably mediates protein-protein interactions. LIM kinase-1 and LIM kinase-2 belong to a small subfamily with a unique combination of 2 N-terminal LIM motifs and a C-terminal protein kinase domain. The LIMK2 protein is phosphorylated and activated by ROCK, a downstream effector of Rho, and LIM kinase 2, in turn, phosphorylates cofilin, inhibiting its actin-depolymerizing activity. It is thought that this pathway contributes to Rho-induced reorganization of the actin cytoskeleton. Two alternative splice variants of LIMK2 that utilize alternative promoters have been identified.

LIM Kinase 2B (LIMK2B) Antibody (N-term) Blocking peptide - References

Blume-Jensen P, et al. Nature 2001. 411: 355. Cantrell D, J. Cell Sci. 2001. 114: 1439. Jhiang S. Oncogene 2000. 19: 5590. Manning G, et al. Science 2002. 298: 1912. Moller, D, et al. Am. J. Physiol. 1994. 266: C351-C359. Robertson, S. et al. Trends Genet. 2000. 16: 368. Robinson D, et al. Oncogene 2000. 19: 5548. Van der Ven, P, et al. Hum. Molec. Genet. 1993. 2: 1889. Vanhaesebroeck, B, et al. Biochem. J. 2000. 346: 561. Van Weering D, et al. Recent Results Cancer Res. 1998. 154: 271.