

### TOLLIP Antibody (N-term) Blocking Peptide Synthetic peptide

Catalog # BP1521a

## Specification

# **TOLLIP Antibody (N-term) Blocking Peptide - Product Information**

Primary Accession

<u>Q9H0E2</u>

## **TOLLIP Antibody (N-term) Blocking Peptide - Additional Information**

Gene ID 54472

Other Names Toll-interacting protein, TOLLIP

### Target/Specificity

The synthetic peptide sequence used to generate the antibody <a href=/product/products/AP1521a>AP1521a</a> was selected from the N-term region of human TOLLIP. 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.

## **TOLLIP Antibody (N-term) Blocking Peptide - Protein Information**

Name TOLLIP

### Function

Component of the signaling pathway of IL-1 and Toll-like receptors (PubMed:<a href="http://www.uniprot.org/citations/10854325" target="\_blank">10854325</a>, PubMed:<a href="http://www.uniprot.org/citations/11751856" target="\_blank">11751856</a>). Inhibits cell activation by microbial products. Recruits IRAK1 to the IL-1 receptor complex (PubMed:<a href="http://www.uniprot.org/citations/10854325" target="\_blank">10854325</a>). Inhibits cell activation by microbial products. Recruits IRAK1 to the IL-1 receptor complex (PubMed:<a href="http://www.uniprot.org/citations/10854325" target="\_blank">10854325</a>). Inhibits IRAK1 phosphorylation and kinase activity (PubMed:<a

href="http://www.uniprot.org/citations/11751856" target="\_blank">11751856</a>). Connects the ubiquitin pathway to autophagy by functioning as a ubiquitin-ATG8 family adapter and thus mediating autophagic clearance of ubiquitin conjugates (PubMed:<a

href="http://www.uniprot.org/citations/25042851" target="\_blank">25042851</a>). The TOLLIP-dependent selective autophagy pathway plays an important role in clearance of cytotoxic polyQ proteins aggregates (PubMed:<a href="http://www.uniprot.org/citations/25042851"



target="\_blank">25042851</a>). In a complex with TOM1, recruits ubiquitin-conjugated proteins onto early endosomes (PubMed:<a href="http://www.uniprot.org/citations/15047686" target="\_blank">15047686</a>). Binds to phosphatidylinositol 3-phosphate (PtdIns(3)P) (PubMed:<a href="http://www.uniprot.org/citations/26320582" target="\_blank">26320582</a>).

**Cellular Location** Cytoplasm. Endosome. Early endosome Note=Localized to endo/exosomal vesicles

### **TOLLIP Antibody (N-term) Blocking Peptide - Protocols**

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

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

#### TOLLIP Antibody (N-term) Blocking Peptide - Images

#### **TOLLIP Antibody (N-term) Blocking Peptide - Background**

Toll-interacting protein (Tollip) was initially identified as an important constituent of the IL-1R signaling pathway. Recently it was showed that this adapter protein is also involved in TLR2 and TLR4 signaling pathways. Tollip coimmunoprecipitates with TLR2 and TLR4 and overexpression of Tollip was shown to inhibit NF-kB activation in response to TLR2 and TLR4 signaling. Human TOLLIP protein is 274-amino acid, which is 97% identical to the mouse sequence. Structurally, Tollip contains a type II C2 motif. Immunoblot analysis showed expression of a 30-kD protein in numerous tissues and cell lines. Functionally, Tollip inhibits cell activation by microbial products, recruits IRAK1 to the IL-1 receptor complex and inhibits IRAK1 phosphorylation and kinase activity. Since the inhibition by Tollip is mediated through its ability to suppress the activity of IRAK, Tollip may act as moderator of the inflammatory response following TLR activation.

### **TOLLIP Antibody (N-term) Blocking Peptide - References**

Zhang, G., et al., J. Biol. Chem. 277(9):7059-7065 (2002).Bulut, Y., et al., J. Immunol. 167(2):987-994 (2001).Burns, K., et al., Nat. Cell Biol. 2(6):346-351 (2000).Volpe, F., et al., FEBS Lett. 419(1):41-44 (1997).