

BAT3 Antibody (C-term) Blocking Peptide

Synthetic peptide

Catalog # BP16737b

Specification

BAT3 Antibody (C-term) Blocking Peptide - Product Information

Primary Accession

[P46379](#)**BAT3 Antibody (C-term) Blocking Peptide - Additional Information**

Gene ID 7917

Other Names

Large proline-rich protein BAG6, BAG family molecular chaperone regulator 6, BCL2-associated athanogene 6, BAG-6, BAG6, HLA-B-associated transcript 3, Protein G3, Protein Scythe, BAG6, BAT3, G3

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.

BAT3 Antibody (C-term) Blocking Peptide - Protein InformationName BAG6 ([HGNC:13919](#))**Function**

ATP-independent molecular chaperone preventing the aggregation of misfolded and hydrophobic patches-containing proteins (PubMed:[21636303](http://www.uniprot.org/citations/21636303)). Functions as part of a cytosolic protein quality control complex, the BAG6/BAT3 complex, which maintains these client proteins in a soluble state and participates in their proper delivery to the endoplasmic reticulum or alternatively can promote their sorting to the proteasome where they undergo degradation (PubMed:[20516149](http://www.uniprot.org/citations/20516149), PubMed:[21636303](http://www.uniprot.org/citations/21636303), PubMed:[21743475](http://www.uniprot.org/citations/21743475), PubMed:[28104892](http://www.uniprot.org/citations/28104892)). The BAG6/BAT3 complex is involved in the post-translational delivery of tail-anchored/type II transmembrane proteins to the endoplasmic reticulum membrane. Recruited to ribosomes, it interacts with the transmembrane region of newly synthesized tail-anchored proteins and together with SGTA and ASNA1 mediates their delivery to the endoplasmic reticulum (PubMed:[20516149](http://www.uniprot.org/citations/20516149), PubMed:[20676083](http://www.uniprot.org/citations/20676083), PubMed:[20676083](http://www.uniprot.org/citations/20676083)).

<http://www.uniprot.org/citations/28104892> target="_blank">28104892, PubMed:25535373). Client proteins that cannot be properly delivered to the endoplasmic reticulum are ubiquitinated by RNF126, an E3 ubiquitin-protein ligase associated with BAG6 and are sorted to the proteasome (PubMed:24981174, PubMed:28104892, PubMed:27193484). SGTA which prevents the recruitment of RNF126 to BAG6 may negatively regulate the ubiquitination and the proteasomal degradation of client proteins (PubMed:23129660, PubMed:25179605, PubMed:27193484). Similarly, the BAG6/BAT3 complex also functions as a sorting platform for proteins of the secretory pathway that are mislocalized to the cytosol either delivering them to the proteasome for degradation or to the endoplasmic reticulum (PubMed:21743475). The BAG6/BAT3 complex also plays a role in the endoplasmic reticulum-associated degradation (ERAD), a quality control mechanism that eliminates unwanted proteins of the endoplasmic reticulum through their retrotranslocation to the cytosol and their targeting to the proteasome. It maintains these retrotranslocated proteins in an unfolded yet soluble state condition in the cytosol to ensure their proper delivery to the proteasome (PubMed:21636303). BAG6 is also required for selective ubiquitin-mediated degradation of defective nascent chain polypeptides by the proteasome. In this context, it may participate in the production of antigenic peptides and play a role in antigen presentation in immune response (By similarity). BAG6 is also involved in endoplasmic reticulum stress-induced pre-emptive quality control, a mechanism that selectively attenuates the translocation of newly synthesized proteins into the endoplasmic reticulum and reroutes them to the cytosol for proteasomal degradation. BAG6 may ensure the proper degradation of these proteins and thereby protects the endoplasmic reticulum from protein overload upon stress (PubMed:26565908). By inhibiting the polyubiquitination and subsequent proteasomal degradation of HSPA2 it may also play a role in the assembly of the synaptonemal complex during spermatogenesis (By similarity). Also positively regulates apoptosis by interacting with and stabilizing the proapoptotic factor AIFM1 (By similarity). By controlling the steady-state expression of the IGF1R receptor, indirectly regulates the insulin-like growth factor receptor signaling pathway (PubMed:26692333).

Cellular Location

Cytoplasm, cytosol. Nucleus. Secreted, extracellular exosome Note=Normally localized in cytosol and nucleus, it can also be released extracellularly, in exosomes, by tumor and myeloid dendritic cells (PubMed:18055229, PubMed:18852879). Cytoplasmic retention is due to interaction with GET4 (PubMed:29042515).

Tissue Location

Expressed by immature dendritic cells (at protein level).

BAT3 Antibody (C-term) Blocking Peptide - Protocols

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

- [Blocking Peptides](#)

BAT3 Antibody (C-term) Blocking Peptide - Images

BAT3 Antibody (C-term) Blocking Peptide - Background

This gene was first characterized as part of a cluster of genes located within the human major

histocompatibility complex class III region. This gene encodes a nuclear protein that is cleaved by caspase 3 and is implicated in the control of apoptosis. In addition, the protein forms a complex with E1A binding protein p300 and is required for the acetylation of p53 in response to DNA damage. Multiple transcript variants encoding different isoforms have been found for this gene.

BAT3 Antibody (C-term) Blocking Peptide - References

Bailey, S.D., et al. *Diabetes Care* 33(10):2250-2253(2010) Ucisik-Akkaya, E., et al. *Mol. Hum. Reprod.* 16(10):770-777(2010) Minami, R., et al. *J. Cell Biol.* 190(4):637-650(2010) Liu, C.Y., et al. *Carcinogenesis* 31(7):1259-1263(2010) Hsieh, Y.Y., et al. *J. Clin. Lab. Anal.* 24(4):262-268(2010)