

SARS virus PUP3 Antibody (C-term) Blocking Peptide
Synthetic peptide
Catalog # BP6003a**Specification**

SARS virus PUP3 Antibody (C-term) Blocking Peptide - Product InformationPrimary Accession
Other Accession[P59634](#)
[NP_828856](#)**SARS virus PUP3 Antibody (C-term) Blocking Peptide - Additional Information****Other Names**

Non-structural protein 6, ns6, Accessory protein 6, Protein X3

Target/Specificity

The synthetic peptide sequence used to generate the antibody [AP6003a](/product/products/AP6003a) was selected from the C-term region of human SARS virus PUP3 . 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.

SARS virus PUP3 Antibody (C-term) Blocking Peptide - Protein Information**Name 6****Function**

Disrupts bidirectional nucleocytoplasmic transport by interacting with host RAE1-NUP98 complex (PubMed:[33849972](http://www.uniprot.org/citations/33849972)). Disrupts cell nuclear import complex formation also by tethering karyopherin alpha 2 and karyopherin beta 1 to the membrane (PubMed:[17596301](http://www.uniprot.org/citations/17596301)). Retention of import factors at the ER/Golgi membrane leads to a loss of transport into the nucleus (PubMed:[17596301](http://www.uniprot.org/citations/17596301)). Thereby prevents STAT1 nuclear translocation in response to interferon signaling, thus blocking the expression of interferon stimulated genes (ISGs) that display multiple antiviral activities (PubMed:[17596301](http://www.uniprot.org/citations/17596301)).

Cellular Location

Host endoplasmic reticulum membrane. Host Golgi apparatus membrane. Host cytoplasm.

Note=Localizes to virus- induced vesicular structures called double membrane vesicles

SARS virus PUP3 Antibody (C-term) Blocking Peptide - Protocols

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

- [Blocking Peptides](#)

SARS virus PUP3 Antibody (C-term) Blocking Peptide - Images

SARS virus PUP3 Antibody (C-term) Blocking Peptide - Background

An outbreak of atypical pneumonia, referred to as severe acute respiratory syndrome (SARS) and first identified in Guangdong Province, China, has spread to several countries. The severity of this disease is such that the mortality rate appears to be ~3 to 6%. A number of laboratories worldwide have undertaken the identification of the causative agent. The National Microbiology Laboratory in Canada obtained the Tor2 isolate from a patient in Toronto, and succeeded in growing a coronavirus-like agent in African Green Monkey Kidney (Vero E6) cells. This coronavirus has been named publicly by the World Health Organization and member laboratories as ?SARS virus?. The SARS membrane proteins, including the major proteins S (Spike) and M (Membrane), are inserted into the endoplasmic reticulum Golgi intermediate compartment (ERGIC) while full length replicated RNA (+ strands) assemble with the N (nucleocapsid) protein. The virus then migrates through the Golgi complex and eventually exits the cell, likely by exocytosis. The site of viral attachment to the host cell resides within the S protein. Oligomeric spike (S) glycoproteins extend from SARS membranes. These integral membrane proteins assemble within the endoplasmic reticulum of infected cells and are subsequently endoproteolyzed in the Golgi, generating noncovalently associated S1 and S2 fragments. Once on the surface of infected cells and virions, peripheral S1 fragments bind carcinoembryonic antigen-related cell adhesion molecule (CEACAM) receptors, and this triggers membrane fusion reactions mediated by integral membrane S2 fragments.

SARS virus PUP3 Antibody (C-term) Blocking Peptide - References

He, R., et al., Biochem. Biophys. Res. Commun. 316(2):476-483 (2004). Snijder, E.J., et al., J. Mol. Biol. 331(5):991-1004 (2003). Marra, M.A., et al., Science 300(5624):1399-1404 (2003).