

## APOBEC3F Antibody (N-term) Blocking Peptide

Synthetic peptide Catalog # BP9176a

### **Specification**

## APOBEC3F Antibody (N-term) Blocking Peptide - Product Information

**Primary Accession** 

**Q8IUX4** 

# APOBEC3F Antibody (N-term) Blocking Peptide - Additional Information

Gene ID 200316

#### **Other Names**

DNA dC->dU-editing enzyme APOBEC-3F, 354-, Apolipoprotein B mRNA-editing enzyme catalytic polypeptide-like 3F, A3F, APOBEC3F

## Target/Specificity

The synthetic peptide sequence used to generate the antibody <a href=/products/AP9176a>AP9176a</a> was selected from the N-term region of human APOBEC3F. 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.

### APOBEC3F Antibody (N-term) Blocking Peptide - Protein Information

### Name APOBEC3F

### **Function**

DNA deaminase (cytidine deaminase) which acts as an inhibitor of retrovirus replication and retrotransposon mobility via deaminase- dependent and -independent mechanisms. Exhibits antiviral activity against viruse such as HIV-1 or HIV-2 (PubMed:<a href="http://www.uniprot.org/citations/15141007" target="\_blank">15141007</a>, PubMed:<a href="http://www.uniprot.org/citations/15152192" target="\_blank">15152192</a>, PubMed:<a href="http://www.uniprot.org/citations/23001005" target="\_blank">23001005</a>, PubMed:<a href="http://www.uniprot.org/citations/34774569" target="\_blank">34774569</a>). After the penetration of retroviral nucleocapsids into target cells of infection and the initiation of reverse transcription, it can induce the conversion of cytosine to uracil in the minus-sense single-strand viral DNA, leading to G-to-A hypermutations in the subsequent plus-strand viral DNA (PubMed:<a href="http://www.uniprot.org/citations/15141007" target="\_blank">15141007</a>). The resultant



detrimental levels of mutations in the proviral genome, along with a deamination-independent mechanism that works prior to the proviral integration, together exert efficient antiretroviral effects in infected target cells. Selectively targets single-stranded DNA and does not deaminate double-stranded DNA or single- or double-stranded RNA. Exhibits antiviral activity also against hepatitis B virus (HBV), equine infectious anemia virus (EIAV), xenotropic MuLV-related virus (XMRV) and simian foamy virus (SFV) and may inhibit the mobility of LTR and non-LTR retrotransposons. May also play a role in the epigenetic regulation of gene expression through the process of active DNA demethylation.

**Cellular Location** 

Cytoplasm. Cytoplasm, P-body.

**Tissue Location** 

Widely expressed. Highly expressed in ovary.

## APOBEC3F Antibody (N-term) Blocking Peptide - Protocols

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

• Blocking Peptides

APOBEC3F Antibody (N-term) Blocking Peptide - Images

# APOBEC3F Antibody (N-term) Blocking Peptide - Background

This protein is a member of the cytidine deaminase gene family. It is one of seven related genes or pseudogenes found in a cluster, thought to result from gene duplication, on chromosome 22. Members of the cluster encode proteins that are structurally and functionally related to the C to U RNA-editing cytidine deaminase APOBEC1. It is thought that the proteins may be RNA editing enzymes and have roles in growth or cell cycle control.

# APOBEC3F Antibody (N-term) Blocking Peptide - References

Khatua, A.K., et.al., Virology 400 (1), 68-75 (2010) Koning, F.A., et.al., J. Virol. 83 (18), 9474-9485 (2009)