

## Phospho-MAPT(\$720) Antibody Blocking peptide

Synthetic peptide Catalog # BP3381a

# **Specification**

## Phospho-MAPT(S720) Antibody Blocking peptide - Product Information

**Primary Accession** 

P10636

# Phospho-MAPT(S720) Antibody Blocking peptide - Additional Information

**Gene ID 4137** 

#### **Other Names**

Microtubule-associated protein tau, Neurofibrillary tangle protein, Paired helical filament-tau, PHF-tau, MAPT, MAPTL, MTBT1, TAU

## **Target/Specificity**

The synthetic peptide sequence used to generate the antibody <a href=/product/products/AP3381a>AP3381a</a> was selected from the region of human Phospho-MAPT-S720. 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.

### Phospho-MAPT(S720) Antibody Blocking peptide - Protein Information

Name MAPT (HGNC:6893)

Synonyms MAPTL, MTBT1, TAU

### **Function**

Promotes microtubule assembly and stability, and might be involved in the establishment and maintenance of neuronal polarity (PubMed:<a href="http://www.uniprot.org/citations/21985311" target="\_blank">21985311</a>). The C-terminus binds axonal microtubules while the N-terminus binds neural plasma membrane components, suggesting that tau functions as a linker protein between both (PubMed:<a href="http://www.uniprot.org/citations/21985311" target="\_blank">21985311" target="\_blank">21985311</a>, PubMed:<a href="http://www.uniprot.org/citations/32961270" target="\_blank">32961270</a>). Axonal polarity is predetermined by TAU/MAPT localization (in the neuronal cell) in the domain of the cell body defined by the centrosome. The short isoforms allow plasticity of the cytoskeleton whereas the longer isoforms may preferentially play a role in its



#### stabilization.

#### **Cellular Location**

Cytoplasm, cytosol. Cell membrane; Peripheral membrane protein; Cytoplasmic side. Cytoplasm, cytoskeleton. Cell projection, axon. Cell projection, dendrite. Secreted Note=Mostly found in the axons of neurons, in the cytosol and in association with plasma membrane components (PubMed:10747907). Can be secreted; the secretion is dependent on protein unfolding and facilitated by the cargo receptor TMED10; it results in protein translocation from the cytoplasm into the ERGIC (endoplasmic reticulum- Golgi intermediate compartment) followed by vesicle entry and secretion (PubMed:32272059).

#### **Tissue Location**

Expressed in neurons. Isoform PNS-tau is expressed in the peripheral nervous system while the others are expressed in the central nervous system

## Phospho-MAPT(S720) Antibody Blocking peptide - Protocols

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

# • Blocking Peptides

Phospho-MAPT(S720) Antibody Blocking peptide - Images

# Phospho-MAPT(S720) Antibody Blocking peptide - Background

MAPT promotes microtubule assembly and stability, and might be involved in the establishment and maintenance of neuronal polarity. The C-terminus binds axonal microtubules while the N-terminus binds neural plasma membrane components, suggesting that tau functions as a linker protein between both. Axonal polarity is predetermined by tau localization (in the neuronal cell) in the domain of the cell body defined by the centrosome. The short isoforms allow plasticity of the cytoskeleton whereas the longer isoforms may preferentially play a role in its stabilization. MAPT gene mutations have been associated with several neurodegenerative disorders such as Alzheimer's disease, Pick's disease, frontotemporal dementia, cortico-basal degeneration and progressive supranuclear palsy.

# Phospho-MAPT(S720) Antibody Blocking peptide - References

Chun, W., J. Biol. Chem. 282 (32), 23410-23417 (2007) Hanger, D.P., J. Biol. Chem. 282 (32), 23645-23654 (2007)