

#### Phospho-GSK3(S21) Antibody Blocking peptide Synthetic peptide Catalog # BP3521a

## Specification

# Phospho-GSK3(S21) Antibody Blocking peptide - Product Information

Primary Accession

#### <u>P49841</u>

## Phospho-GSK3(S21) Antibody Blocking peptide - Additional Information

Gene ID 2932

**Other Names** Glycogen synthase kinase-3 beta, GSK-3 beta, Serine/threonine-protein kinase GSK3B, GSK3B

## Target/Specificity

The synthetic peptide sequence used to generate the antibody <a href=/product/products/AP3521a>AP3521a</a> was selected from the region of human Phospho-GSK3-S21. 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-GSK3(S21) Antibody Blocking peptide - Protein Information

Name GSK3B (HGNC:4617)

#### Function

Constitutively active protein kinase that acts as a negative regulator in the hormonal control of glucose homeostasis, Wnt signaling and regulation of transcription factors and microtubules, by phosphorylating and inactivating glycogen synthase (GYS1 or GYS2), EIF2B, CTNNB1/beta-catenin, APC, AXIN1, DPYSL2/CRMP2, JUN, NFATC1/NFATC, MAPT/TAU and MACF1 (PubMed:<a href="http://www.uniprot.org/citations/11430833" target="\_blank">11430833</a>, PubMed:<a href="http://www.uniprot.org/citations/11430833" target="\_blank">12554650</a>, PubMed:<a href="http://www.uniprot.org/citations/12554650" target="\_blank">14690523</a>, PubMed:<a href="http://www.uniprot.org/citations/14690523" target="\_blank">14690523</a>, PubMed:<a href="http://www.uniprot.org/citations/16484495" target="\_blank">16484495</a>, PubMed:<a href="http://www.uniprot.org/citations/1846781" target="\_blank">1846781</a>, PubMed:<a href="http://www.uniprot.org/citations/1846781" target="\_blank">1846781</a>, PubMed:<a href="http://www.uniprot.org/citations/1846781" target="\_blank">1846781</a>, PubMed:<a href="http://www.uniprot.org/citations/1846781" target="\_blank">1846781</a>, PubMed:<a href="http://www.uniprot.org/citations/20937854" target="\_blank">20937854</a>, PubMed:<a href="http://www.uniprot.org/citations/20937854" target="\_blank">20937854</a>, PubMed:<a href="http://www.uniprot.org/citations/20937854" target="\_blank">20937854</a>, PubMed:<a href="http://www.uniprot.org/citations/20937854" target="\_blank">20937854</a>, PubMed:<a href="http://www.uniprot.org/citations/9072970" target="\_blank">9072970</a>). Requires primed phosphorylation of the majority of its substrates (PubMed:<a



href="http://www.uniprot.org/citations/11430833" target="\_blank">11430833</a>, PubMed:<a href="http://www.uniprot.org/citations/16484495" target="\_blank">16484495</a>). In skeletal muscle, contributes to insulin regulation of glycogen synthesis by phosphorylating and inhibiting GYS1 activity and hence glycogen synthesis (PubMed:<a

href="http://www.uniprot.org/citations/8397507" target="\_blank">8397507</a>). May also mediate the development of insulin resistance by regulating activation of transcription factors (PubMed:<a href="http://www.uniprot.org/citations/8397507" target="\_blank">8397507</a>). Regulates protein synthesis by controlling the activity of initiation factor 2B (EIF2BE/EIF2B5) in the same manner as glycogen synthase (PubMed:<a href="http://www.uniprot.org/citations/8397507" target="\_blank">8397507</a>). In Wnt signaling, GSK3B forms a multimeric complex with APC, AXIN1 and CTNNB1/beta-catenin and phosphorylates the N-terminus of CTNNB1 leading to its degradation mediated by ubiquitin/proteasomes (PubMed:<a

href="http://www.uniprot.org/citations/12554650" target="\_blank">12554650</a>). Phosphorylates JUN at sites proximal to its DNA-binding domain, thereby reducing its affinity for DNA (PubMed:<a href="http://www.uniprot.org/citations/1846781"

target="\_blank">1846781</a>). Phosphorylates NFATC1/NFATC on conserved serine residues promoting NFATC1/NFATC nuclear export, shutting off NFATC1/NFATC gene regulation, and thereby opposing the action of calcineurin (PubMed:<a

href="http://www.uniprot.org/citations/9072970" target="\_blank">9072970</a>). Phosphorylates MAPT/TAU on 'Thr-548', decreasing significantly MAPT/TAU ability to bind and stabilize microtubules (PubMed:<a href="http://www.uniprot.org/citations/14690523"

target="\_blank">14690523</a>). MAPT/TAU is the principal component of neurofibrillary tangles in Alzheimer disease (PubMed:<a href="http://www.uniprot.org/citations/14690523" target=" blank">14690523</a>). Plays an important role in ERBB2-dependent stabilization of

microtubules at the cell cortex (PubMed:<a href="http://www.uniprot.org/citations/20937854" target="\_blank">20937854</a>). Phosphorylates MACF1, inhibiting its binding to microtubules which is critical for its role in bulge stem cell migration and skin wound repair (By similarity). Probably regulates NF-kappa-B (NFKB1) at the transcriptional level and is required for the

NF-kappa-B-mediated anti- apoptotic response to TNF-alpha (TNF/TNFA) (By similarity). Negatively regulates replication in pancreatic beta-cells, resulting in apoptosis, loss of beta-cells and diabetes (By similarity). Through phosphorylation of the anti-apoptotic protein MCL1, may control cell apoptosis in response to growth factors deprivation (By similarity). Phosphorylates MUC1 in breast cancer cells, decreasing the interaction of MUC1 with CTNNB1/beta-catenin (PubMed:<a

href="http://www.uniprot.org/citations/9819408" target="\_blank">9819408</a>). Is necessary for the establishment of neuronal polarity and axon outgrowth (PubMed:<a

href="http://www.uniprot.org/citations/20067585" target="\_blank">20067585</a>). Phosphorylates MARK2, leading to inhibition of its activity (By similarity). Phosphorylates SIK1 at

'Thr-182', leading to sustainment of its activity (PubMed:<a

href="http://www.uniprot.org/citations/18348280" target="\_blank">18348280</a>).

Phosphorylates ZC3HAV1 which enhances its antiviral activity (PubMed: <a

href="http://www.uniprot.org/citations/22514281" target="\_blank">22514281</a>). Phosphorylates SNAI1, leading to its ubiquitination and proteasomal degradation (PubMed:<a href="http://www.uniprot.org/citations/15448698" target="\_blank">15448698</a>, PubMed:<a href="http://www.uniprot.org/citations/15647282" target="\_blank">15647282</a>, PubMed:<a href="http://www.uniprot.org/citations/15647282" target="\_blank">15647282</a>, PubMed:<a href="http://www.uniprot.org/citations/25827072" target="\_blank">25827072</a>, PubMed:<a href="http://www.uniprot.org/citations/25827072" target="\_blank">29059170</a>, PubMed:<a href="http://www.uniprot.org/citations/29059170" target="\_blank">29059170</a>,

Phosphorylates SFPQ at 'Thr-687' upon T-cell activation (PubMed:<a href="http://www.uniprot.org/citations/20932480" target=" blank">20932480</a>).

Phosphorylates NR1D1 st 'Ser-55' and 'Ser-59' and stabilizes it by protecting it from proteasomal degradation. Regulates the circadian clock via phosphorylation of the major clock components including BMAL1, CLOCK and PER2 (PubMed:<a href="http://www.uniprot.org/citations/19946213" target="\_blank">19946213</a>, PubMed:<a href="http://www.uniprot.org/citations/28903391" target="\_blank">28903391</a>). Phosphorylates FBXL2 at 'Thr-404' and primes it for ubiquitination by the SCF(FBXO3) complex and proteasomal degradation (By similarity). Phosphorylates CLOCK AT 'Ser-427' and targets it for proteasomal degradation (PubMed:<a href="http://www.uniprot.org/citations/19946213" target="\_blank">19946213</a>). Phosphorylates the ser-427' and targets it for proteasomal degradation (By similarity). Phosphorylates CLOCK AT 'Ser-427' and targets it for proteasomal degradation (PubMed:<a href="http://www.uniprot.org/citations/19946213" target="\_blank">19946213</a>). Phosphorylates BMAL1 at 'Ser-17' and 'Ser-21' and primes it for ubiquitination and proteasomal



degradation (PubMed: <a href="http://www.uniprot.org/citations/28903391" target=" blank">28903391</a>). Phosphorylates OGT at 'Ser-3' or 'Ser-4' which positively regulates its activity. Phosphorylates MYCN in neuroblastoma cells which may promote its degradation (PubMed:<a href="http://www.uniprot.org/citations/24391509" target=" blank">24391509</a>). Regulates the circadian rhythmicity of hippocampal long-term potentiation and BMAL1 and PER2 expression (By similarity). Acts as a regulator of autophagy by mediating phosphorylation of KAT5/TIP60 under starvation conditions, activating KAT5/TIP60 acetyltransferase activity and promoting acetylation of key autophagy regulators, such as ULK1 and RUBCNL/Pacer (PubMed: <a href="http://www.uniprot.org/citations/30704899" target=" blank">30704899</a>). Negatively regulates extrinsic apoptotic signaling pathway via death domain receptors. Promotes the formation of an anti-apoptotic complex, made of DDX3X, BRIC2 and GSK3B, at death receptors, including TNFRSF10B. The anti-apoptotic function is most effective with weak apoptotic signals and can be overcome by stronger stimulation (PubMed:<a href="http://www.uniprot.org/citations/18846110" target=" blank">18846110</a>). Phosphorylates E2F1, promoting the interaction between E2F1 and USP11, stabilizing E2F1 and promoting its activity (PubMed:<a href="http://www.uniprot.org/citations/17050006" target=" blank">17050006</a>, PubMed:<a href="http://www.uniprot.org/citations/28992046" target=" blank">28992046</a>). Phosphorylates mTORC2 complex component RICTOR at 'Ser-1235' in response to endoplasmic stress, inhibiting mTORC2 (PubMed:<a href="http://www.uniprot.org/citations/21343617" target=" blank">21343617</a>). Phosphorylates mTORC2 complex component RICTOR at 'Thr-1695' which facilitates FBXW7-mediated ubiguitination and subsequent degradation of RICTOR (PubMed:<a href="http://www.uniprot.org/citations/25897075" target="\_blank">25897075</a>). Phosphorylates FXR1, promoting FXR1 ubiguitination by the SCF(FBXO4) complex and FXR1 degradation by the proteasome (By similarity). Phosphorylates interleukin-22 receptor subunit IL22RA1, preventing its proteasomal degradation (By similarity).

## **Cellular Location**

Cytoplasm. Nucleus. Cell membrane. Note=The phosphorylated form shows localization to cytoplasm and cell membrane (PubMed:20937854) The MEMO1-RHOA-DIAPH1 signaling pathway controls localization of the phosphorylated form to the cell membrane (PubMed:20937854)

**Tissue Location** 

Expressed in testis, thymus, prostate and ovary and weakly expressed in lung, brain and kidney. Colocalizes with EIF2AK2/PKR and TAU in the Alzheimer disease (AD) brain

### Phospho-GSK3(S21) Antibody Blocking peptide - Protocols

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

Blocking Peptides

# Phospho-GSK3(S21) Antibody Blocking peptide - Images

### Phospho-GSK3(S21) Antibody Blocking peptide - Background

Glycogen synthase kinase-3 (GSK3) is a proline-directed serine-threonine kinase that was initially identified as a phosphorylating and inactivating glycogen synthase. Two isoforms, alpha (GSK3A; MIM 606784) and beta, show a high degree of amino acid homology (Stambolic and Woodgett, 1994 [PubMed 7980435]).GSK3B is involved in energy metabolism, neuronal cell development, and body pattern formation (Plyte et al., 1992 [PubMed 1333807]).[supplied by OMIM].

### Phospho-GSK3(S21) Antibody Blocking peptide - References

Izumi,N.,J. Biol. Chem. 283 (19), 12981-12991 (2008)Deng,H.,J. Biol. Chem. 283 (15), 10198-10207 (2008)Ma,C.,J. Biol. Chem. 283 (14), 9248-9256 (2008)