

# Phospho-STAT1-Y701 Antibody Blocking peptide

Synthetic peptide Catalog # BP3259a

### **Specification**

### Phospho-STAT1-Y701 Antibody Blocking peptide - Product Information

**Primary Accession** 

P42224

## Phospho-STAT1-Y701 Antibody Blocking peptide - Additional Information

### **Gene ID 6772**

#### **Other Names**

Signal transducer and activator of transcription 1-alpha/beta, Transcription factor ISGF-3 components p91/p84, STAT1

## **Target/Specificity**

The synthetic peptide sequence used to generate the antibody <a href=/product/products/AP3259a>AP3259a</a> was selected from the 696-705 <CR>region of human Phospho-STAT1-Y701. 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-STAT1-Y701 Antibody Blocking peptide - Protein Information

### Name STAT1

## **Function**

Signal transducer and transcription activator that mediates cellular responses to interferons (IFNs), cytokine KITLG/SCF and other cytokines and other growth factors (PubMed:<a href="http://www.uniprot.org/citations/12764129" target="\_blank">12764129</a>, PubMed:<a href="http://www.uniprot.org/citations/12855578" target="\_blank">12855578</a>, PubMed:<a href="http://www.uniprot.org/citations/15322115" target="\_blank">15322115</a>, PubMed:<a href="http://www.uniprot.org/citations/23940278" target="\_blank">23940278</a>, PubMed:<a href="http://www.uniprot.org/citations/34508746" target="\_blank">34508746</a>, PubMed:<a href="http://www.uniprot.org/citations/35568036" target="\_blank">35568036</a>, PubMed:<a href="http://www.uniprot.org/citations/9724754" target="\_blank">9724754</a>). Following type I IFN (IFN-alpha and IFN-beta) binding to cell surface receptors, signaling via protein kinases leads to activation of Jak kinases (TYK2 and JAK1) and to tyrosine phosphorylation of STAT1 and STAT2.



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The phosphorylated STATs dimerize and associate with ISGF3G/IRF-9 to form a complex termed ISGF3 transcription factor, that enters the nucleus (PubMed:<a

href="http://www.uniprot.org/citations/28753426" target="\_blank">28753426</a>, PubMed:<a href="http://www.uniprot.org/citations/35568036" target="\_blank">35568036</a>). ISGF3 binds to the IFN stimulated response element (ISRE) to activate the transcription of IFN-stimulated genes (ISG), which drive the cell in an antiviral state (PubMed:<a

href="http://www.uniprot.org/citations/28753426" target=" blank">28753426</a>, PubMed:<a href="http://www.uniprot.org/citations/35568036" target="blank">35568036</a>). In response to type II IFN (IFN-gamma), STAT1 is tyrosine- and serine-phosphorylated (PubMed: <a href="http://www.uniprot.org/citations/26479788" target=" blank">26479788</a>). It then forms a homodimer termed IFN-gamma-activated factor (GAF), migrates into the nucleus and binds to the IFN gamma activated sequence (GAS) to drive the expression of the target genes, inducing a cellular antiviral state (PubMed: <a href="http://www.uniprot.org/citations/8156998" target=" blank">8156998</a>). Becomes activated in response to KITLG/SCF and KIT signaling (PubMed: <a href="http://www.uniprot.org/citations/15526160" target="blank">15526160</a>). May mediate cellular responses to activated FGFR1, FGFR2, FGFR3 and FGFR4 (PubMed:<a href="http://www.uniprot.org/citations/19088846" target=" blank">19088846</a>). Following bacterial lipopolysaccharide (LPS)-induced TLR4 endocytosis, phosphorylated at Thr-749 by IKBKB which promotes binding of STAT1 to the 5'-TTTGAGGC-3' sequence in the ARID5A promoter, resulting in transcriptional activation of ARID5A and subsequent ARID5A-mediated stabilization of IL6 (PubMed:<a href="http://www.uniprot.org/citations/32209697" target=" blank">32209697</a>). Phosphorylation at Thr-749 also promotes binding of STAT1 to

target="\_blank">32209697</a>). Phosphorylation at Thr-749 also promotes binding of STAT1 to the 5'-TTTGAGTC-3' sequence in the IL12B promoter and activation of IL12B transcription (PubMed:<a href="http://www.uniprot.org/citations/32209697" target="\_blank">32209697</a>). Involved in food tolerance in small intestine: associates with the Gasdermin-D, p13 cleavage product (13 kDa GSDMD) and promotes transcription of CIITA, inducing type 1 regulatory T (Tr1) cells in upper small intestine (By similarity).

### **Cellular Location**

Cytoplasm. Nucleus Note=Translocated into the nucleus upon tyrosine phosphorylation and dimerization, in response to IFN-gamma and signaling by activated FGFR1, FGFR2, FGFR3 or FGFR4 (PubMed:15322115). Monomethylation at Lys- 525 is required for phosphorylation at Tyr-701 and translocation into the nucleus (PubMed:28753426). Translocates into the nucleus in response to interferon-beta stimulation (PubMed:26479788)

## Phospho-STAT1-Y701 Antibody Blocking peptide - Protocols

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

### • Blocking Peptides

Phospho-STAT1-Y701 Antibody Blocking peptide - Images

## Phospho-STAT1-Y701 Antibody Blocking peptide - Background

STAT1 is a member of the STAT protein family. In response to cytokines and growth factors, STAT family members are phosphorylated by the receptor associated kinases, and then form homo- or heterodimers that translocate to the cell nucleus where they act as transcription activators. This protein can be activated by various ligands including interferon-alpha, interferon-gamma, EGF, PDGF and IL6. This protein mediates the expression of a variety of genes, which is thought to be important for cell viability in response to different cell stimuli and pathogens.

### Phospho-STAT1-Y701 Antibody Blocking peptide - References

Garcin, D., et al., J. Virol. 78(16):8799-8811 (2004).Melen, K., et al., J. Med. Virol. 73(4):536-547 (2004).Klampfer, L., et al., J. Biol. Chem. 279(29):30358-30368 (2004).Marg, A., et al., J. Cell Biol.





165(6):823-833 (2004).Chim, C.S., et al., Blood 103(12):4630-4635 (2004).