

**Phospho-YAP(S127) Blocking Peptide**  
**Synthetic peptide**  
**Catalog # BP3769a****Specification****Phospho-YAP(S127) Blocking Peptide - Product Information**

Primary Accession [P46937](#)  
Other Accession [Q2EJA0](#), [P46938](#),  
[P46936](#),  
[NP\\_001123617.1](#)

**Phospho-YAP(S127) Blocking Peptide - Additional Information**

**Gene ID** 10413

**Other Names**

Transcriptional coactivator YAP1,  
Yes-associated protein 1, Protein yorkie  
homolog, Yes-associated protein YAP65  
homolog, YAP1, YAP65

**Target/Specificity**

The synthetic peptide sequence is selected  
from aa 119-133 of HUMAN YAP1

**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-YAP(S127) Blocking Peptide - Protein Information**

**Name** YAP1

**Synonyms** YAP65

**Phospho-YAP(S127) Blocking Peptide - Background**

Transcriptional regulator which can act both  
as a coactivator and a corepressor and is the  
critical downstream regulatory target in the  
Hippo signaling pathway that plays a pivotal  
role in organ size control and tumor  
suppression by restricting proliferation and  
promoting apoptosis. The core of this pathway  
is composed of a kinase cascade wherein  
MST1/MST2, in complex with its regulatory  
protein SAV1, phosphorylates and activates  
LATS1/2 in complex with its regulatory protein  
MOB1, which in turn phosphorylates and  
inactivates YAP1 oncoprotein and WWTR1/TAZ.  
Plays a key role to control cell proliferation in  
response to cell contact. Phosphorylation of  
YAP1 by LATS1/2 inhibits its translocation into  
the nucleus to regulate cellular genes  
important for cell proliferation, cell death, and  
cell migration. The presence of TEAD  
transcription factors are required for it to  
stimulate gene expression, cell growth,  
anchorage-independent growth, and epithelial  
mesenchymal transition (EMT) induction.  
Isoform 2 and isoform 3 can activate the  
C-terminal fragment (CTF) of ERBB4 (isoform  
3).

**Function**

Transcriptional regulator which can act both as a coactivator and a corepressor and is the critical downstream regulatory target in the Hippo signaling pathway that plays a pivotal role in organ size control and tumor suppression by restricting proliferation and promoting apoptosis (PubMed:<a href="http://www.uniprot.org/citations/17974916" target="\_blank">17974916</a>, PubMed:<a href="http://www.uniprot.org/citations/18280240" target="\_blank">18280240</a>, PubMed:<a href="http://www.uniprot.org/citations/18579750" target="\_blank">18579750</a>, PubMed:<a href="http://www.uniprot.org/citations/21364637" target="\_blank">21364637</a>, PubMed:<a href="http://www.uniprot.org/citations/30447097" target="\_blank">30447097</a>). The core of this pathway is composed of a kinase cascade wherein STK3/MST2 and STK4/MST1, in complex with its regulatory protein SAV1, phosphorylates and activates LATS1/2 in complex with its regulatory protein MOB1, which in turn phosphorylates and inactivates YAP1 oncoprotein and WWTR1/TAZ (PubMed:<a href="http://www.uniprot.org/citations/18158288" target="\_blank">18158288</a>). Plays a key role in tissue tension and 3D tissue shape by regulating cortical actomyosin network formation. Acts via ARHGAP18, a Rho GTPase activating protein that suppresses F-actin polymerization (PubMed:<a href="http://www.uniprot.org/citations/25778702" target="\_blank">25778702</a>). Plays a key role in controlling cell proliferation in response to cell contact. Phosphorylation of YAP1 by LATS1/2 inhibits its translocation into the nucleus to regulate cellular genes important for cell proliferation, cell death, and cell migration (PubMed:<a href="http://www.uniprot.org/citations/18158288" target="\_blank">18158288</a>). The presence of TEAD transcription factors are required for it to stimulate gene expression, cell growth, anchorage- independent growth, and epithelial mesenchymal transition (EMT) induction (PubMed:<a href="http://www.uniprot.org/citations/18579750" target="\_blank">18579750</a>). Suppresses ciliogenesis via acting as a transcriptional corepressor of the TEAD4

target genes AURKA and PLK1 (PubMed:<a href="http://www.uniprot.org/citations/25849865" target="\_blank">25849865</a>). In conjunction with WWTR1, involved in the regulation of TGFB1-dependent SMAD2 and SMAD3 nuclear accumulation (By similarity).

#### **Cellular Location**

Cytoplasm. Nucleus. Note=Both phosphorylation and cell density can regulate its subcellular localization (PubMed:18158288, PubMed:20048001). Phosphorylation sequesters it in the cytoplasm by inhibiting its translocation into the nucleus (PubMed:18158288, PubMed:20048001). At low density, predominantly nuclear and is translocated to the cytoplasm at high density (PubMed:18158288, PubMed:20048001, PubMed:25849865). PTPN14 induces translocation from the nucleus to the cytoplasm (PubMed:22525271). Localized mainly to the nucleus in the early stages of embryo development with expression becoming evident in the cytoplasm at the blastocyst and epiblast stages (By similarity).  
{ECO:0000250|UniProtKB:P46938,  
ECO:0000269|PubMed:18158288,  
ECO:0000269|PubMed:20048001,  
ECO:0000269|PubMed:22525271,  
ECO:0000269|PubMed:25849865}

#### **Tissue Location**

Increased expression seen in some liver and prostate cancers. Isoforms lacking the transactivation domain found in striatal neurons of patients with Huntington disease (at protein level).

#### **Phospho-YAP(S127) Blocking Peptide - Protocols**

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

- [Blocking Peptides](#)