

**Phospho-MDM2(S395) Antibody Blocking peptide**  
**Synthetic peptide**  
**Catalog # BP3579a****Specification****Phospho-MDM2(S395) Antibody Blocking peptide**  
**- Product Information**Primary Accession [Q00987](#)**Phospho-MDM2(S395) Antibody Blocking peptide**  
**- Additional Information****Gene ID** 4193**Other Names**E3 ubiquitin-protein ligase Mdm2, 632-,  
Double minute 2 protein, Hdm2,  
Oncoprotein Mdm2, p53-binding protein  
Mdm2, MDM2**Target/Specificity**

The synthetic peptide sequence used to generate the antibody [AP3579a](/products/AP3579a) was selected from the region of human Phospho-MDM2-pS395. 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-MDM2(S395) Antibody Blocking peptide**  
**- Protein Information****Name** MDM2**Phospho-MDM2(S395) Antibody Blocking peptide - Background**

MDM2 is a target of the transcription factor tumor protein p53. This protein is a nuclear phosphoprotein that binds and inhibits transactivation by tumor protein p53, as part of an autoregulatory negative feedback loop. Overexpression of MDM2 can result in excessive inactivation of tumor protein p53, diminishing its tumor suppressor function. This protein has E3 ubiquitin ligase activity, which targets tumor protein p53 for proteasomal degradation. MDM2 also affects the cell cycle, apoptosis, and tumorigenesis through interactions with other proteins, including retinoblastoma 1 and ribosomal protein L5.

**Phospho-MDM2(S395) Antibody Blocking peptide - References**

Lu X, et al. (2007) The Wip1 Phosphatase acts as a gatekeeper in the p53-Mdm2 autoregulatory loop. *Cancer Cell* 12, 342-54  
Balass M, et al. (2002) Characterization of two peptide epitopes on Mdm2 oncoprotein that affect p53 degradation. *Peptides* 23, 1719-25  
Maya R, et al. (2001) ATM-dependent phosphorylation of Mdm2 on serine 395: role in p53 activation by DNA damage. *Genes Dev* 15, 1067-77

Also component of the TRIM28/KAP1-ERBB4-MDM2 complex which links growth factor and DNA damage response pathways. Mediates ubiquitination and subsequent proteasome degradation of DYRK2 in nucleus. Ubiquitinates IGF1R and SNAI1 and promotes them to proteasomal degradation (PubMed:<a href="http://www.uniprot.org/citations/12821780" target="\_blank">12821780</a>, PubMed:<a href="http://www.uniprot.org/citations/15053880" target="\_blank">15053880</a>, PubMed:<a href="http://www.uniprot.org/citations/15195100" target="\_blank">15195100</a>, PubMed:<a href="http://www.uniprot.org/citations/15632057" target="\_blank">15632057</a>, PubMed:<a href="http://www.uniprot.org/citations/16337594" target="\_blank">16337594</a>, PubMed:<a href="http://www.uniprot.org/citations/17290220" target="\_blank">17290220</a>, PubMed:<a href="http://www.uniprot.org/citations/19098711" target="\_blank">19098711</a>, PubMed:<a href="http://www.uniprot.org/citations/19219073" target="\_blank">19219073</a>, PubMed:<a href="http://www.uniprot.org/citations/19837670" target="\_blank">19837670</a>, PubMed:<a href="http://www.uniprot.org/citations/19965871" target="\_blank">19965871</a>, PubMed:<a href="http://www.uniprot.org/citations/20173098" target="\_blank">20173098</a>)

tations/20385133"  
target="\_blank">20385133</a>,  
PubMed:<a href="http://www.uniprot.org/ci  
tations/20858735"  
target="\_blank">20858735</a>,  
PubMed:<a href="http://www.uniprot.org/ci  
tations/22128911"  
target="\_blank">22128911</a>).  
Ubiquitinates DCX, leading to DCX  
degradation and reduction of the dendritic  
spine density of olfactory bulb granule cells  
(By similarity). Ubiquitinates DLG4, leading  
to proteasomal degradation of DLG4 which  
is required for AMPA receptor endocytosis  
(By similarity). Negatively regulates  
NDUFS1, leading to decreased  
mitochondrial respiration, marked oxidative  
stress, and commitment to the  
mitochondrial pathway of apoptosis  
(PubMed:<a href="http://www.uniprot.org/c  
itations/30879903"  
target="\_blank">30879903</a>). Binds  
NDUFS1 leading to its cytosolic retention  
rather than mitochondrial localization  
resulting in decreased supercomplex  
assembly (interactions between complex I  
and complex III), decreased complex I  
activity, ROS production, and apoptosis  
(PubMed:<a href="http://www.uniprot.org/c  
itations/30879903"  
target="\_blank">30879903</a>).

#### **Cellular Location**

Nucleus, nucleoplasm. Cytoplasm. Nucleus,  
nucleolus. Nucleus. Note=Expressed  
predominantly in the nucleoplasm.  
Interaction with ARF(P14) results in the  
localization of both proteins to the  
nucleolus. The nucleolar localization signals  
in both ARF(P14) and MDM2 may be  
necessary to allow efficient nucleolar  
localization of both proteins. Colocalizes  
with RASSF1 isoform A in the nucleus

#### **Tissue Location**

Ubiquitous. Isoform Mdm2-A, isoform  
Mdm2-B, isoform Mdm2-C, isoform Mdm2-D,  
isoform Mdm2-E, isoform Mdm2-F and  
isoform Mdm2-G are observed in a range of  
cancers but absent in normal tissues

### **Phospho-MDM2(S395) Antibody Blocking peptide - Protocols**

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

- [Blocking Peptides](#)