

p53 Tumor Suppressor Protein Antibody - With BSA and Azide
Mouse Monoclonal Antibody [Clone TRP/817]
Catalog # AH12450

Specification

p53 Tumor Suppressor Protein Antibody - With BSA and Azide - Product Information

Application	WB, IHC, IF, FC
Primary Accession	P04637
Other Accession	7157 , 654481
Reactivity	Human
Host	Mouse
Clonality	Monoclonal
Isotype	Mouse / IgG2b / kappa
Calculated MW	53kDa KDa

p53 Tumor Suppressor Protein Antibody - With BSA and Azide - Additional Information

Gene ID 7157

Other Names

Cellular tumor antigen p53, Antigen NY-CO-13, Phosphoprotein p53, Tumor suppressor p53, TP53, P53

Application Note

WB~~1:1000
IHC~~1:100~500
IF~~1:50~200
FC~~1:10~50

Storage

Store at 2 to 8°C. Antibody is stable for 24 months.

Precautions

p53 Tumor Suppressor Protein Antibody - With BSA and Azide is for research use only and not for use in diagnostic or therapeutic procedures.

p53 Tumor Suppressor Protein Antibody - With BSA and Azide - Protein Information

Name TP53

Synonyms P53

Function

Multifunctional transcription factor that induces cell cycle arrest, DNA repair or apoptosis upon binding to its target DNA sequence (PubMed: [11025664](http://www.uniprot.org/citations/11025664) target="_blank">11025664, PubMed: [12524540](http://www.uniprot.org/citations/12524540) target="_blank">12524540, PubMed: [12810724](http://www.uniprot.org/citations/12810724) target="_blank">12810724, PubMed: [15186775](http://www.uniprot.org/citations/15186775) target="_blank">15186775, PubMed: [15340061](http://www.uniprot.org/citations/15340061) target="_blank">15340061)

[target="_blank">15340061](#), PubMed: [target="_blank">17317671](http://www.uniprot.org/citations/17317671), PubMed: [target="_blank">17349958](http://www.uniprot.org/citations/17349958), PubMed: [target="_blank">19556538](http://www.uniprot.org/citations/19556538), PubMed: [target="_blank">20673990](http://www.uniprot.org/citations/20673990), PubMed: [target="_blank">20959462](http://www.uniprot.org/citations/20959462), PubMed: [target="_blank">22726440](http://www.uniprot.org/citations/22726440), PubMed: [target="_blank">24051492](http://www.uniprot.org/citations/24051492), PubMed: [target="_blank">24652652](http://www.uniprot.org/citations/24652652), PubMed: [target="_blank">35618207](http://www.uniprot.org/citations/35618207), PubMed: [target="_blank">36634798](http://www.uniprot.org/citations/36634798), PubMed: [target="_blank">38653238](http://www.uniprot.org/citations/38653238), PubMed: [target="_blank">9840937](http://www.uniprot.org/citations/9840937)). Acts as a tumor suppressor in many tumor types; induces growth arrest or apoptosis depending on the physiological circumstances and cell type (PubMed: [target="_blank">11025664](http://www.uniprot.org/citations/11025664), PubMed: [target="_blank">12524540](http://www.uniprot.org/citations/12524540), PubMed: [target="_blank">12810724](http://www.uniprot.org/citations/12810724), PubMed: [target="_blank">15186775](http://www.uniprot.org/citations/15186775), PubMed: [target="_blank">15340061](http://www.uniprot.org/citations/15340061), PubMed: [target="_blank">17189187](http://www.uniprot.org/citations/17189187), PubMed: [target="_blank">17317671](http://www.uniprot.org/citations/17317671), PubMed: [target="_blank">17349958](http://www.uniprot.org/citations/17349958), PubMed: [target="_blank">19556538](http://www.uniprot.org/citations/19556538), PubMed: [target="_blank">20673990](http://www.uniprot.org/citations/20673990), PubMed: [target="_blank">20959462](http://www.uniprot.org/citations/20959462), PubMed: [target="_blank">22726440](http://www.uniprot.org/citations/22726440), PubMed: [target="_blank">24051492](http://www.uniprot.org/citations/24051492), PubMed: [target="_blank">24652652](http://www.uniprot.org/citations/24652652), PubMed: [target="_blank">38653238](http://www.uniprot.org/citations/38653238), PubMed: [target="_blank">9840937](http://www.uniprot.org/citations/9840937)). Negatively regulates cell division by controlling expression of a set of genes required for this process (PubMed: [target="_blank">11025664](http://www.uniprot.org/citations/11025664), PubMed: [target="_blank">12524540](http://www.uniprot.org/citations/12524540), PubMed: [target="_blank">12810724](http://www.uniprot.org/citations/12810724), PubMed: [target="_blank">15186775](http://www.uniprot.org/citations/15186775), PubMed: [target="_blank">15340061](http://www.uniprot.org/citations/15340061), PubMed: [target="_blank">17317671](http://www.uniprot.org/citations/17317671), PubMed: [target="_blank">17349958](http://www.uniprot.org/citations/17349958), PubMed: [target="_blank">19556538](http://www.uniprot.org/citations/19556538), PubMed: [target="_blank">20673990](http://www.uniprot.org/citations/20673990), PubMed: [target="_blank">20959462](http://www.uniprot.org/citations/20959462), PubMed: [target="_blank">22726440](http://www.uniprot.org/citations/22726440), PubMed: [target="_blank">24051492](http://www.uniprot.org/citations/24051492), PubMed: [target="_blank">24652652](http://www.uniprot.org/citations/24652652), PubMed: [target="_blank">9840937](http://www.uniprot.org/citations/9840937)). One of the activated genes is an inhibitor of cyclin-dependent kinases. Apoptosis induction seems to be mediated either by stimulation of BAX and FAS antigen expression, or by repression of Bcl-2 expression (PubMed: [target="_blank">12524540](http://www.uniprot.org/citations/12524540), PubMed: [target="_blank">17189187](http://www.uniprot.org/citations/17189187)). Its pro-apoptotic activity is activated via its interaction with PPP1R13B/ASPP1 or TP53BP2/ASPP2 (PubMed: [target="_blank">12524540](http://www.uniprot.org/citations/12524540)). However, this activity is inhibited when the interaction with PPP1R13B/ASPP1 or TP53BP2/ASPP2 is displaced by PPP1R13L/iASPP (PubMed: [target="_blank">12524540](http://www.uniprot.org/citations/12524540)). In cooperation with mitochondrial PPIF is involved in activating oxidative stress-induced necrosis; the function is largely independent of transcription. Induces the transcription of long intergenic non-coding RNA p21 (lincRNA-p21) and lincRNA-Mkn1.

LincRNA-p21 participates in TP53-dependent transcriptional repression leading to apoptosis and seems to have an effect on cell-cycle regulation. Implicated in Notch signaling cross-over. Prevents CDK7 kinase activity when associated to CAK complex in response to DNA damage, thus stopping cell cycle progression. Isoform 2 enhances the transactivation activity of isoform 1 from some but not all TP53-inducible promoters. Isoform 4 suppresses transactivation activity and impairs growth suppression mediated by isoform 1. Isoform 7 inhibits isoform 1-mediated apoptosis. Regulates the circadian clock by repressing CLOCK-BMAL1-mediated transcriptional activation of PER2 (PubMed:24051492).

Cellular Location

Cytoplasm. Nucleus. Nucleus, PML body. Endoplasmic reticulum. Mitochondrion matrix. Cytoplasm, cytoskeleton, microtubule organizing center, centrosome Note=Recruited into PML bodies together with CHEK2 (PubMed:12810724) Translocates to mitochondria upon oxidative stress (PubMed:22726440) Translocates to mitochondria in response to mitomycin C treatment (PubMed:27323408). Competitive inhibition of TP53 interaction with HSPA9/MOT-2 by UBXLN2A results in increased protein abundance and subsequent translocation of TP53 to the nucleus (PubMed:24625977) [Isoform 2]: Nucleus. Cytoplasm. Note=Localized mainly in the nucleus with minor staining in the cytoplasm [Isoform 4]: Nucleus. Cytoplasm. Note=Predominantly nuclear but translocates to the cytoplasm following cell stress [Isoform 8]: Nucleus. Cytoplasm. Note=Localized in both nucleus and cytoplasm in most cells. In some cells, forms foci in the nucleus that are different from nucleoli

Tissue Location

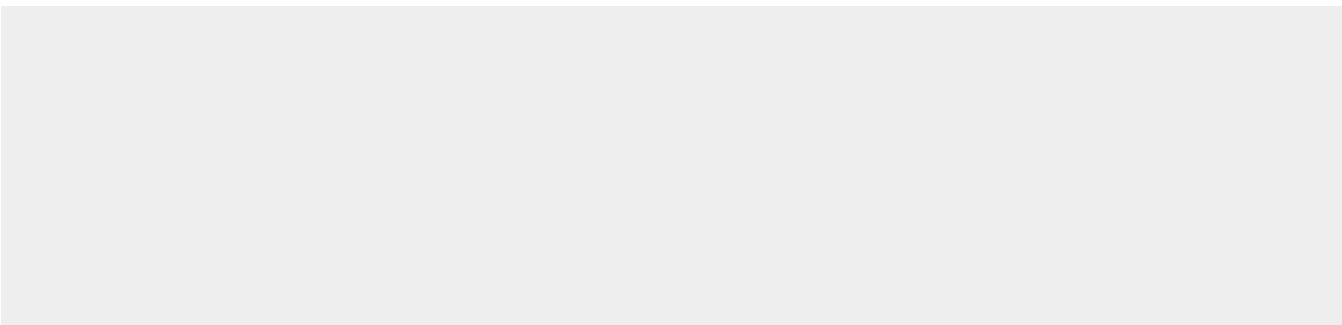
Ubiquitous. Isoforms are expressed in a wide range of normal tissues but in a tissue-dependent manner. Isoform 2 is expressed in most normal tissues but is not detected in brain, lung, prostate, muscle, fetal brain, spinal cord and fetal liver. Isoform 3 is expressed in most normal tissues but is not detected in lung, spleen, testis, fetal brain, spinal cord and fetal liver. Isoform 7 is expressed in most normal tissues but is not detected in prostate, uterus, skeletal muscle and breast. Isoform 8 is detected only in colon, bone marrow, testis, fetal brain and intestine. Isoform 9 is expressed in most normal tissues but is not detected in brain, heart, lung, fetal liver, salivary gland, breast or intestine

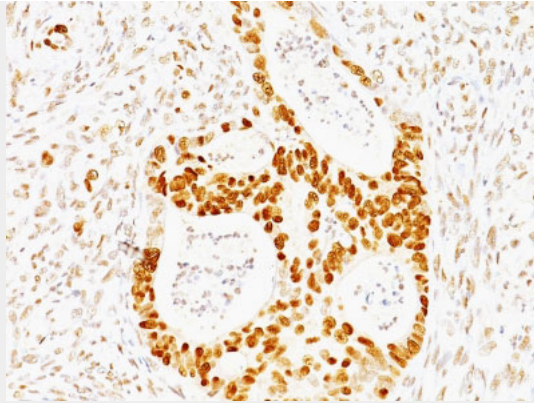
p53 Tumor Suppressor Protein Antibody - With BSA and Azide - Protocols

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

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

p53 Tumor Suppressor Protein Antibody - With BSA and Azide - Images





Formalin-fixed, paraffin-embedded human Colon Carcinoma stained with p53 Monoclonal Antibody (TRP/817)

p53 Tumor Suppressor Protein Antibody - With BSA and Azide - Background

Recognizes a 53kDa protein, which is identified as p53 suppressor gene product. It reacts with the mutant as well as the wild form of p53 protein. p53 is a tumor suppressor gene expressed in a wide variety of tissue types and is involved in regulating cell growth, replication, and apoptosis. It binds to MDM2, SV40 T antigen and human papilloma virus E6 protein. Positive nuclear staining with p53 antibody has been reported to be a negative prognostic factor in breast carcinoma, lung carcinoma, colorectal, and urothelial carcinoma. Anti-p53 positivity has also been used to differentiate uterine serous carcinoma from endometrioid carcinoma as well as to detect intratubular germ cell neoplasia. Mutations involving p53 are found in a wide variety of malignant tumors, including breast, ovarian, bladder, colon, lung, and melanoma.

p53 Tumor Suppressor Protein Antibody - With BSA and Azide - References

Soussi, T., et al. 2000. p53 website and analysis of p53 gene mutations in human cancer: forging a link between epidemiology and carcinogenesis. Hum. Mutat. 15: 105-113. |