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# Phospho-p53 (T55) Antibody

Rabbit mAb Catalog # AP90494

### **Product Information**

**Application** WB **Primary Accession** P04637 Reactivity Human Clonality Monoclonal

**Other Names** Antigen NY-CO-13; TP53; Tumor suppressor p53; Cellular tumor antigen p53;

Phosphoprotein p53

Isotype Rabbit IgG Host Rabbit Calculated MW 43653

#### **Additional Information**

Dilution WB 1:500~1:2000 **Purification** Affinity-chromatography

**Immunogen** A synthesized peptide derived from human Phospho-p53 (T55)

**Description** Plays an essential role in the regulation of cell cycle, specifically in the

transition from G0 to G1. It is found in very low levels in normal cells, however, in a variety of transformed cell lines, it is expressed in high

amounts, and believed to contribute to transformation and malignancy. p53 is

a DNA-binding protein containing DNA-binding, oligomerization and transcription activation domains. It is postulated to bind as a tetramer to a p53-binding site and activate expression of downstream genes that inhibit growth and/or invasion, and thus function as a tumor suppressor.

**Storage Condition and Buffer** Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium

azide and 50% glycerol. Store at +4°C short term. Store at -20°C long term.

Avoid freeze / thaw cycle.

#### **Protein Information**

TP53 Name

**Synonyms** P53

**Function** Multifunctional transcription factor that induces cell cycle arrest, DNA repair

or apoptosis upon binding to its target DNA sequence (PubMed: 11025664,

PubMed: 12524540, PubMed: 12810724, PubMed: 15186775, PubMed: 15340061, PubMed: 17317671, PubMed: 17349958, PubMed:19556538, PubMed:20673990, PubMed:20959462, PubMed:22726440, PubMed:24051492, PubMed:24652652, PubMed:35618207, PubMed:36634798, PubMed:38653238,

PubMed: 9840937). Acts as a tumor suppressor in many tumor types; induces

growth arrest or apoptosis depending on the physiological circumstances and cell type (PubMed: 11025664, PubMed: 12524540, PubMed: 12810724, PubMed: 15186775, PubMed: 15340061, PubMed: 17189187, PubMed:17317671, PubMed:17349958, PubMed:19556538, PubMed: 20673990, PubMed: 20959462, PubMed: 22726440, PubMed: 24051492, PubMed: 24652652, PubMed: 38653238, PubMed: 9840937). Negatively regulates cell division by controlling expression of a set of genes required for this process (PubMed: 11025664, PubMed: 12524540, PubMed: 12810724, PubMed: 15186775, PubMed: 15340061, PubMed: 17317671, PubMed: 17349958, PubMed: 19556538, PubMed: 20673990, PubMed: 20959462, PubMed:22726440, PubMed:24051492, PubMed:24652652, PubMed: 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: 12524540, PubMed: 17189187). Its pro-apoptotic activity is activated via its interaction with PPP1R13B/ASPP1 or TP53BP2/ASPP2 (PubMed: 12524540). However, this activity is inhibited when the interaction with PPP1R13B/ASPP1 or TP53BP2/ASPP2 is displaced by PPP1R13L/iASPP (PubMed: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-Mkln1. 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).

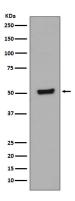
#### **Cellular Location**

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 UBXN2A 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

## **Images**



Western blot analysis of Phospho-p53 (T55) in HEK293 cell lysates treated with Calyculin A and Okadaic Acid.

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