



# Phospho-TP53 (S9) Recombinant Monoclonal Antibody

<b>Product Code</b>	CSB-RA024077A09phHU
<b>Abbreviation</b>	Cellular tumor antigen p53
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	P04637
<b>Immunogen</b>	A synthesized peptide derived from Human Phospho-TP53 (S9)
<b>Species Reactivity</b>	Human
<b>Tested Applications</b>	ELISA, IF; Recommended dilution: IF:1:20-1:200
<b>Relevance</b>	Acts as a tumor suppressor in many tumor types; induces growth arrest or apoptosis depending on the physiological circumstances and cell type. Involved in cell cycle regulation as a trans-activator that acts to negatively regulate cell division by controlling a set of genes required for this process. 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. 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 seem to have to 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-ARNTL/BMAL1-mediated transcriptional activation of PER2 (PubMed:24051492).
<b>Form</b>	Liquid
<b>Conjugate</b>	Non-conjugated
<b>Storage Buffer</b>	Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
<b>Purification Method</b>	Affinity-chromatography
<b>Isotype</b>	Rabbit IgG
<b>Clonality</b>	Monoclonal
<b>Alias</b>	Cellular tumor antigen p53, Antigen NY-CO-13, Phosphoprotein p53, Tumor suppressor p53, TP53, P53
<b>Immunogen Species</b>	Homo sapiens (Human)

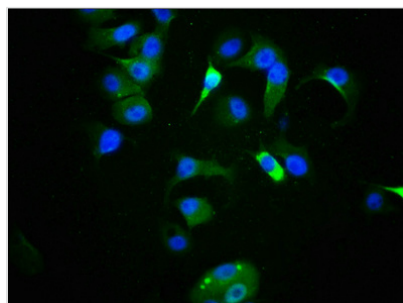


**Research Area** Cell Biology

**Gene Names** TP53

**Clone No.** 2E1

**Image**



Immunofluorescence staining of MCF-7 cells with CSB-RA024077A09phHU at 1:100, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4?. The secondary antibody was Alexa Fluor 488-conjugated AffiniPure Goat Anti-Rabbit IgG (H+L).

**Description**

In the development of the phospho-TP53 (S9) recombinant monoclonal antibody, the initial phase involves the retrieval of genes encoding the TP53 antibody from rabbits immunized with a synthetic peptide originating from the human TP53 protein phosphorylated at S9. Following this, these genes are adeptly integrated into expression vectors. Subsequently, these genetically modified vectors are introduced into mammalian suspension cells, where they are thoughtfully cultivated to encourage the production and secretion of the antibodies. Following this growth phase, an intricate purification process employing affinity chromatography is executed to meticulously isolate the phospho-TP53 (S9) recombinant monoclonal antibody from the surrounding cell culture supernatant. Lastly, the functionality of the antibody is meticulously scrutinized through ELISA and IF tests, conclusively affirming its capacity to engage with the human TP53 protein phosphorylated at S9.

Phosphorylation of p53 at S9 serves as a crucial regulatory mechanism to maintain genomic integrity and prevent the development of cancer by coordinating DNA repair, cell cycle control, and cell fate decisions in response to stress and damage. Dysregulation of p53 phosphorylation can lead to uncontrolled cell proliferation and is often observed in cancer cells.