

Phospho-p53 (Ser20) Ab

Cat.#: AF3073 Size: 100ul,200ul	Concn.: 1mg/ml Source: Rabbit	Mol.Wt.: 53kDa Clonality: Polyclonal
Application:	WB 1:500-1:2000 IHC 1:200, IF/ICC 1:100-1:500	
Reactivity:	Human,Mouse,Rat	
Purification:	The Ab is from purified rabbit serum by affinity purification via sequential chromatography on phospho- and non-phospho-peptide affinity columns.	
Specificity:	Phospho-p53 (Ser20) Ab detects endogenous levels of p53 only when phosphorylated at Serine 20	
Immunogen:	A synthesized peptide derived from human p53 around the phosphorylation site of Serine 20	
Uniprot:	P04637	
Description:	Tumor protein p53, a nuclear protein, 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.	
Subcellular Location:	Cytoplasm; Cytoplasm. Nucleus Endoplasmic reticulum. Interact nuclear localization. Recruited in CHEK2; Nucleus. Cytoplasm. Loc cytoplasm in most cells. In som nucleus that are different from Localized in the nucleus in most cytoplasm in some cells; Nucleus mainly in the nucleus with minor Nucleus. Cytoplasm. Predomina the cytoplasm when expressed Cytoplasm. Predominantly nucl cytoplasm following cell stress.	tion with BANP promotes into PML bodies together with icalized in both nucleus and e cells, forms foci in the nucleoli; Nucleus. Cytoplasm. t cells but found in the us. Cytoplasm. Localized or staining in the cytoplasm; antly nuclear but localizes to with isoform 4 and Nucleus. ear but translocates to the
Tissue Specificity:	Ubiquitous. Isoforms are express normal tissues but in a tissue-d is expressed in most normal tis brain, lung, prostate, muscle, for fetal liver. Isoform 3 is express is not detected in lung, spleen, and fetal liver. Isoform 7 is exp but is not detected in prostate, breast. Isoform 8 is detected or	ependent manner. Isoform 2 sues but is not detected in etal brain, spinal cord and ed in most normal tissues but testis, fetal brain, spinal cord ressed in most normal tissues uterus, skeletal muscle and

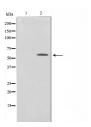


testis, fetal brain and intestine. Isoform 9 is expressed in most normal tissues but is not detected in brain, heart, lung, fetal liver, salivary gland, breast or intestine.

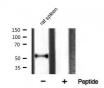
Similarity: The nuclear export signal acts as a transcriptional repression domain. The TADI and TADII motifs (residues 17 to 25 and 48 to 56) correspond both to 9aaTAD motifs which are transactivation domains present in a large number of yeast and animal transcription factors.Belongs to the p53 family.

Storage Condition and R Buffer: N

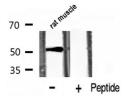
Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.Store at -20 °C.Stable for 12 months from date of receipt



Western blot analysis of p53 phosphorylation expression in UV treated COS7 whole cell lysates,The lane on the left is treated with the antigen-specific peptide.



Western blot analysis of p53 phosphorylation expression in rat spleen lysates, The lane on the right is treated with the antigen-specific peptide.

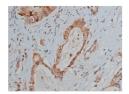


Western blot analysis of p53 phosphorylation expression in rat muscle lysates, The lane on the right is treated with the antigen-specific peptide.



AF3073 at 1/50 staining human lung cancer tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.





AF3073 at 1/50 staining human lung cancer tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.



AF3073 staining MDA-MB-435 by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted Ab in 5% w/v milk , 1X TBS, 0.1% Tween@20 at 4°C with gentle shaking, overnight.

For Research Use Only. Not for use in diagnostic and therapeutic procedures. Not for resale without express authorization.