

SwissProt

Storage/Stability

Product Data Sheet

Anti-p53 (Phospho-S392) Antibody

P04637 (Human)

freeze/thaw cycles.

Catalog #	Source	Reactivity	Applications
CPA4291	Rabbit	Н	WB, IH, IF/IC
Description		Rabbit polyclonal antibod	y to p53 (Phospho-S392)
Immunogen		KLH-conjugated synthetic	phosphopeptide corresponding to residues surrounding
		S392 of human p53 protei	n. The exact sequence is proprietary.
Purification		The antibody was purified	by immunogen affinity chromatography.
Specificity		Recognizes endogenous le	evels of p53 protein only when phosphorylated at S392.
Clonality		Polyclonal	
Conjugation			
Form		Liquid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
		and 0.01% sodium azide.	
Dilution		WB (1/500 - 1/1000), IH (1/	100 - 1/200), IF/IC (1/100 - 1/500)
Gene Symbol		TP53	
Alternative Na	ames	P53; Cellular tumor antige	n p53; Antigen NY-CO-13; Phosphoprotein p53; Tumor
		suppressor p53	
Entrez Gene		7157 (Human)	

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

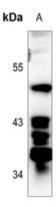
Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

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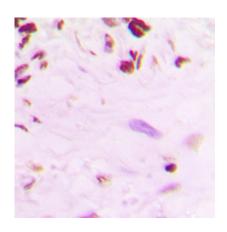
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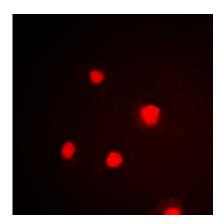
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Western blot analysis of p53 (Phospho-S392) expression in HEK293T (A) whole cell lysates. (Predicted band size: 43 kD; Observed band size: 53 kD)



Immunohistochemical analysis of p53 (Phospho-S392) staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of p53 (Phospho-S392) staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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