

Product Data Sheet

Anti-p53 Antibody

Catalog #	Source	Reactivity	Applications
CPA4294	Rabbit	H, M, R	WB, IH, IF/IC
Description	F	Rabbit polyclonal antibody t	o p53
Immunogen	ł	KLH-conjugated synthetic pe	ptide encompassing a sequence within the C-term
	r	region of human p53. The e	act sequence is proprietary.
Purification	٦	The antibody was purified b	y immunogen affinity chromatography.
Specificity	ſ	Recognizes endogenous leve	els of p53 protein.
Clonality	F	Polyclonal	
Form	l	Liquid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	ā	and 0.01% sodium azide.	
Dilution	١	WB (1/500 - 1/1000), IH (1/10	0 - 1/200), IF/IC (1/100 - 1/500)
Gene Symbol	٦	TP53	
Alternative N	ames l	P53; Cellular tumor antigen	p53; Antigen NY-CO-13; Phosphoprotein p53; Tumor
	S	suppressor p53	
Entrez Gene	7	7157 (Human); 22059 (Mou	se); 24842 (Rat)
SwissProt	F	P04637 (Human); P02340 (N	1ouse); P10361 (Rat)
Storage/Stabi	lity S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid
	f	freeze/thaw cycles.	

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

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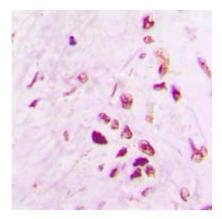


kDa	Α	в	С	D
200				
140				
100				
80				
60				
50	-	-	-	-
40				
30				100
20				

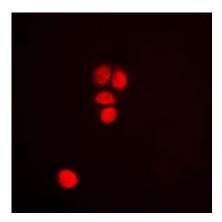
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Western blot analysis of p53 expression in HeLa (A), SHSY5Y (B), Raw264.7 (C), PC12 (D) whole cell lysates.



Immunohistochemical analysis of p53 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of p53 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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