

Product Data Sheet

Anti-p53 (Phospho-S37) Antibody

Catalog #	Source	Reactivity	Applications		
-		-			
CPA3345	Rabbit	Н	WB, IH		
Description	to p53 (Phospho-S37)				
Immunogen		KLH-conjugated synthetic p	hosphopeptide corresponding to residues surrounding		
		S37 of human p53 protein.	The exact sequence is proprietary.		
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous levels of p53 protein only when phosphorylated at S37.			
Clonality	onality Polyclonal				
Conjugation					
Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% g					
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/1	00 - 1/200)		
Gene Symbol		ТР53			
Alternative Na	ames	P53; Cellular tumor antiger	p53; Antigen NY-CO-13; Phosphoprotein p53; Tumor		
		suppressor p53			
Entrez Gene		7157 (Human)			
SwissProt		P04637 (Human)			
Storage/Stabi	ility	Shipped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
		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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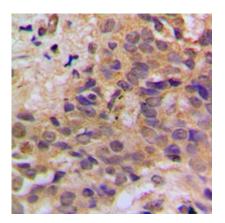
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For research purposes only, not for human use

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Western blot analysis of p53 (Phospho-S37) expression in COLO205 (A), Jurkat UV-treated (B) whole cell lysates. (Predicted band size: 43 kD; Observed band size: 53 kD)



Immunohistochemical analysis of p53 (Phospho-S37) staining in human breast 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.

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