

Product Data Sheet

Anti-A-RAF (Phospho-Y302) Antibody

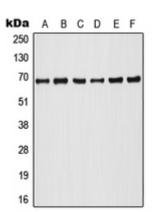
Catalog #	Source	Reactivity	Applications			
CPA1072	Rabbit	H, M, R, B, P, S	WB, IH, IF/IC			
Description	Rab	Rabbit polyclonal antibody to A-RAF (Phospho-Y302)				
Immunogen	KLH	-conjugated synthetic pho	sphopeptide corresponding to residues surrounding			
	Y302	2 of human A-RAF protein	. The exact sequence is proprietary.			
Purification	The	antibody was purified by	immunogen affinity chromatography.			
Specificity	Reco	ognizes endogenous levels	s of A-RAF protein only when phosphorylated at Y302.			
Clonality	Polyclonal					
Conjugation						
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	and	0.01% sodium azide.				
Dilution	WB	WB (1/500 - 1/1000), IH (1/50 - 1/100), IF/IC (1/50 - 1/200)				
Gene Symbol	ARA	F				
Alternative Na	ames ARA	F1; PKS; PKS2; Serine/thre	eonine-protein kinase A-Raf; Proto-oncogene A-Raf;			
	Prot	o-oncogene A-Raf-1; Prot	o-oncogene Pks			
Entrez Gene	369	369 (Human); 11836 (Mouse); 64363 (Rat)				
SwissProt	P10	398 (Human); P04627 (Mc	ouse); P14056 (Rat)			
Storage/Stabi	lity Ship	ped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid			
	free	ze/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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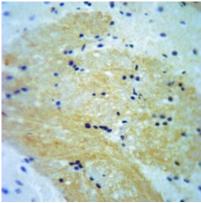




For research purposes only, not for human use

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Western blot analysis of A-RAF (Phospho-Y302) expression in HeLa PMA-treated (A), Raji (B), HT29 (C), NIH3T3 (D), SP2/O PMA-treated (E), PC12 PMA-treated (F) whole cell lysates. (Predicted band size: 67 kD; Observed band size: 68 kD)



Immunohistochemical analysis of A-RAF (Phospho-Y302) 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.

Immunofluorescent analysis of A-RAF (Phospho-Y302) staining in HepG2 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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