

Product Data Sheet

Anti-PRAF2 Antibody

Catalog #	Source	Reactivity	Applications
CPA2542	Rabbit	H, M, R, B, Mk, P	WB, IH, IF/IC
Description	Ral	bbit polyclonal antibody to	PRAF2
Immunogen	KLł	H-conjugated synthetic pept	ide encompassing a sequence within the C-term
	reg	ion of human PRAF2. The e	xact sequence is proprietary.
Purification	The	e antibody was purified by i	mmunogen affinity chromatography.
Specificity	Red	cognizes endogenous levels	of PRAF2 protein.
Clonality	Pol	lyclonal	
Conjugation			
Form	Liq	uid in 0.42% Potassium pho	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	d 0.01% sodium azide.	
Dilution	WE	3 (1/500 - 1/1000), IH (1/50 -	1/100), IF/IC (1/50 - 1/200)
Gene Symbol	PR	AF2	
Alternative Na	ames PR/	A1 family protein 2	
Entrez Gene	112	230 (Human); 54637 (Mous	e)
SwissProt	06	0831 (Human); Q9JIG8 (Mo	use)
Storage/Stabi	lity Shi	pped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid
	fre	eze/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

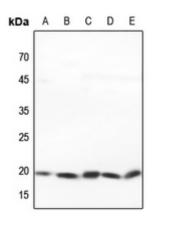
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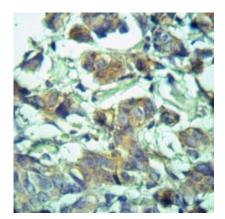


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Western blot analysis of PRAF2 expression in HEK293T (A), Hela (B), mouse brain (C), rat brain (D), rat lung (E) whole cell lysates. (Predicted band size: 19 kD; Observed band size: 19 kD)



Immunohistochemical analysis of PRAF2 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 PRAF2 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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