

# **Product Data Sheet**

### Anti-BRK (Phospho-Y447) Antibody

Catalog #	Source	Reactivity	Applications		
CPA5746	Rabbit	, H, R	WB, IF/IC		
Description	Rabl	Rabbit polyclonal antibody to BRK (Phospho-Y447)			
Immunogen	KLH	-conjugated synthetic	phosphopeptide corresponding to residues surrounding		
	Y447	7 of human BRK protei	n. The exact sequence is proprietary.		
Purification	The	antibody was purified	by immunogen affinity chromatography.		
Specificity	Reco	ognizes endogenous le	vels of BRK protein only when phosphorylated at Y447.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqu	id 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), IF/IC	1/100 - 1/500)		
Gene Symbol	PTK	6			
Alternative Na	ames BRK	; Protein-tyrosine kina	se 6; Breast tumor kinase; Tyrosine-protein kinase BRK		
Entrez Gene	5753	3 (Human)			
SwissProt	Q13	882 (Human)			
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	ery 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

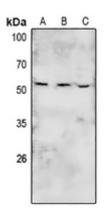
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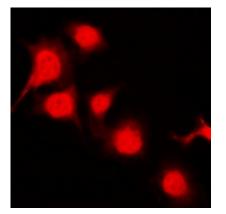


For research purposes only, not for human use

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Western blot analysis of BRK (Phospho-Y447) expression in HEK293T (A), rat kidney (B), rat lung (C) whole cell lysates. (Predicted band size: 51 kD; Observed band size: 52 kD)



Immunofluorescent analysis of BRK (Phospho-Y447) staining in NIH3T3 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 Alexa Fluor 647-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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