

Product Data Sheet

Anti-CRK Antibody

Catalog #	Source	Reactivity	Applications
CPA3185	Rabbit	H, M, R	WB, IH, IF/IC
Description	Ra	bbit polyclonal antibody	to CRK
Immunogen	KL	H-conjugated synthetic p	eptide encompassing a sequence within the center
	re	gion of human CRK. The e	exact sequence is proprietary.
Purification	Th	e antibody was purified b	by immunogen affinity chromatography.
Specificity	Re	cognizes endogenous lev	els of CRK protein.
Clonality	Ро	lyclonal	
Conjugation			
Form	Lic	quid in 0.42% Potassium p	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	an	d 0.01% sodium azide.	
Dilution	W	B (1/500 - 1/1000), IH (1/5	0 - 1/100), IF/IC (1/50 - 1/200)
Gene Symbol	CR	K	
Alternative Na	ames Ad	lapter molecule crk; Prot	o-oncogene c-Crk; p38
Entrez Gene	13	98 (Human); 12928 (Mou	ıse); 54245 (Rat)
SwissProt	P4	6108 (Human); Q64010 (Mouse); Q63768 (Rat)
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid
	fre	eeze/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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Western blot analysis of CRK expression in mouse liver (A), rat liver (B) whole cell lysates. (Predicted band size: 33 kD; Observed band size: 42 kD)



Immunohistochemical analysis of CRK 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 CRK staining in HuvEc 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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