

Product Data Sheet

Anti-CRKL Antibody

Catalog #	Source	Reactivity	Applications
CPA5787	Rabbit	H, M, R	WB, IH, IF/IC
Description	Rat	obit polyclonal antibody	to CRKL
Immunogen	KLH	I-conjugated synthetic p	eptide encompassing a sequence within the center
	reg	ion of human CRKL. The	exact sequence is proprietary.
Purification	The	e antibody was purified	by immunogen affinity chromatography.
Specificity	Rec	cognizes endogenous lev	vels of CRKL protein.
Clonality	Pol	yclonal	
Conjugation			
Form	Liq	uid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	d 0.01% sodium azide.	
Dilution	WE	3 (1/500 - 1/1000), IH (1/5	0 - 1/100), IF/IC (1/100 - 1/500)
Gene Symbol	CRI	٢L	
Alternative Na	ames Crk	-like protein	
Entrez Gene	139	99 (Human); 12929 (Mo	use); 100911248, 287942 (Rat)
SwissProt	P46	6109 (Human); P47941 (Mouse); Q5U2U2 (Rat)
Storage/Stabi	lity Shi	pped at 4°C. Upon deliv	ery 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

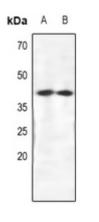
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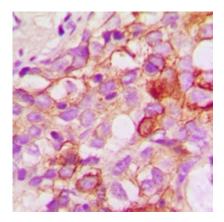
Cohesion

For research purposes only, not for human use

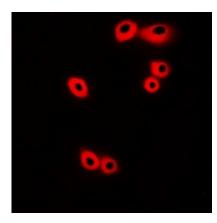
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Western blot analysis of CRKL expression in HEK293T (A), SGC7901 (B) whole cell lysates. (Predicted band size: 33 kD; Observed band size: 39 kD)



Immunohistochemical analysis of CRKL 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 CRKL 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 Alexa Fluor 647-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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