

Product Data Sheet

Anti-BCR Antibody

Catalog #	Source	Reactivity	Applications		
CPA4350	Rabbit	H <i>,</i> M, R	SE, WB, IH		
Description	Ra	abbit polyclonal antibody	to BCR		
Immunogen	KL	H-conjugated synthetic p	eptide encompassing a sequence within the N-term		
	re	gion of human BCR. The e	xact sequence is proprietary.		
Purification	Th	ne antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous lev	els of BCR protein.		
Clonality	lonality Polyclonal				
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.			
Dilution	SE	(Use a concentration of (0.5 ug/ml), WB (1/500 - 1/1000), IH (1/100 - 1/200)		
Gene Symbol	BC	CR			
Alternative Na	ames BC	CR1; D22S11; Breakpoint o	luster region protein; Renal carcinoma antigen		
	N	Y-REN-26			
Entrez Gene	61	13 (Human); 110279 (Mou	se)		
SwissProt	P1	11274 (Human); Q6PAJ1 (N	Aouse)		
Storage/Stabi	lity Sh	nipped at 4°C. Upon delive	ry 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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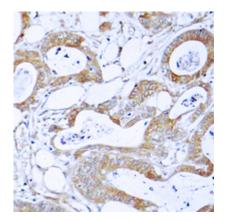


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kDa A B C 180 130 95 Western blot analysis of BCR expression in mouse brain (A), EC9706 (B), U87MG (C) whole cell lysates. (Predicted band size: 142 kD; Observed band size: 160 kD)



Immunohistochemical analysis of BCR staining in human colon 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.

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