

Product Data Sheet

Anti-CREB Antibody

Catalog #	Source	Reactivity	Applications		
CPA1269	Rabbit	H, M, R, B, C, P, Z	WB, IH, ChIP		
Description	F	Rabbit polyclonal antibody to	CREB		
Immunogen	K	KLH-conjugated synthetic pept	ide encompassing a sequence within the center		
	r	egion of human CREB. The ex	act sequence is proprietary.		
Purification	Т	The antibody was purified by i	mmunogen affinity chromatography.		
Specificity	F	Recognizes endogenous levels	of CREB protein.		
Clonality	P	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	and 0.01% sodium azide.			
Dilution	V	NB (1/500 - 1/1000), IH (1/50 -	1/100), ChIP (1/10 - 1/100)		
Gene Symbol	C	CREB1			
Alternative Na	ames C	Cyclic AMP-responsive elemer	t-binding protein 1; CREB-1; cAMP-responsive		
	e	element-binding protein 1			
Entrez Gene	1	1385 (Human); 12912 (Mouse); 81646 (Rat)		
SwissProt	P	P16220 (Human); Q01147 (Mo	ouse); P15337 (Rat)		
Storage/Stabi	lity S	Shipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	f	reeze/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

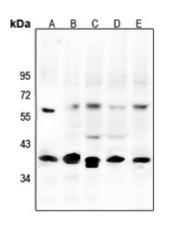
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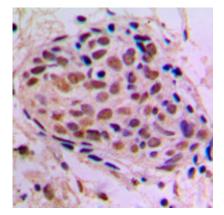


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Western blot analysis of CREB expression in C6 (A), AML12 (B), HCT116 (C), Panc1 (D), A549 (E) whole cell lysates. (Predicted band size: 35 kD; Observed band size: 43 kD)



Immunohistochemical analysis of CREB 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.

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