

Product Data Sheet

Anti-BAD Antibody

Catalog #	Source	Reactivity	Applications		
CPA5910	Rabbit	H, M, R, B	WB, IH		
Description	F	Rabbit polyclonal antibody t	o BAD		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
	r	region of human BAD. The e	xact sequence is proprietary.		
Purification	٦	The antibody was purified b	y immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	els of BAD protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ć	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	- 1/200)		
Gene Symbol	E	BAD			
Alternative Na	imes l	BBC6; BCL2L8; Bcl2 antagon	st of cell death; BAD; Bcl-2-binding component 6;		
	E	Bcl-2-like protein 8; Bcl2-L-8	; Bcl-XL/Bcl-2-associated death promoter		
Entrez Gene	ŗ	572 (Human); 12015 (Mouse	e); 64639 (Rat)		
SwissProt	(Q92934 (Human); Q61337 (I	Mouse); O35147 (Rat)		
Storage/Stabil	ity S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	f	freeze/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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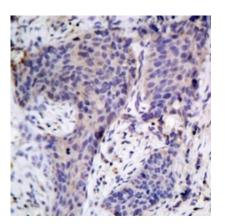
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Western blot analysis of BAD expression in A375 (A), HEK293T (B), Hela (C), AML12 (D) whole cell lysates. (Predicted band size: 18 kD; Observed band size: 23 kD)



Immunohistochemical analysis of BAD 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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