

Product Data Sheet

Anti-EBP1 Antibody

Catalog #	Source	Reactivity	Applications		
CQA3073	Rabbit	H, M, R	WB, IH, IF/IC, IP		
Description	Ra	bbit polyclonal antibody	to EBP1		
Immunogen	Re	combinant full length pro	otein of human EBP1		
Purification	Th	e antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels of EBP1 protein.			
Clonality	Ро	lyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	d 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/2000), IH (1/50) - 1/200), IF/IC (1/50 - 1/200), IP (1/50 - 1/100)		
Gene Symbol	PA	2G4			
Alternative N	ames EB	P1; Proliferation-associat	ed protein 2G4; Cell cycle protein p38-2G4 homolog;		
	hG	64-1; ErbB3-binding prote	in 1		
Entrez Gene	50	36 (Human); 18813 (Mou	ise)		
SwissProt	QS)UQ80 (Human); P50580	(Mouse)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	ry 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

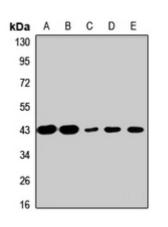
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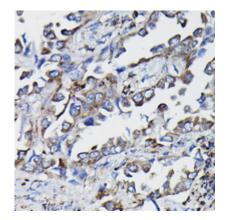


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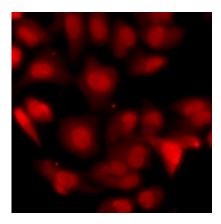
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Western blot analysis of EBP1 expression in Hela (A), K562 (B), PC12 (C), NIH3T3 (D), mouse liver (E) whole cell lysates. (Predicted band size: 38; 43 kD; Observed band size: 43 kD)



Immunohistochemical analysis of EBP1 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 EBP1 staining in U2OS 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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