

## **Product Data Sheet**

### **Anti-4EBP1** Antibody

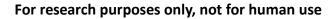
Catalog #	Source	Reactivity	Applications		
CPA4399	Rabbit	H, M, R, Mk	WB, IH		
Description	Ra	bbit polyclonal antibody t	o 4EBP1		
Immunogen	KLI	H-conjugated synthetic pe	ptide encompassing a sequence within the center		
	reg	region of human 4EBP1. The exact sequence is proprietary.			
Purification	The	e antibody was purified by	immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous leve	ls of 4EBP1 protein.		
Clonality	Ро	lyclonal			
Conjugation					
Form	Liq	juid in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	an	d 0.01% sodium azide.			
Dilution	WE	B (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol	EIF	4EBP1			
Alternative Na	ames EIF	4EBP1; Eukaryotic transla	tion initiation factor 4E-binding protein 1; 4E-BP1;		
	elF	4E-binding protein 1; Pho	sphorylated heat- and acid-stable protein regulated by		
	ins	sulin 1; PHAS-I			
Entrez Gene	19	78 (Human); 13685 (Mous	e); 116636 (Rat)		
SwissProt	Q1	.3541 (Human); Q60876 (I	Nouse); Q62622 (Rat)		
Storage/Stabi	lity Shi	ipped at 4°C. Upon deliver	y 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

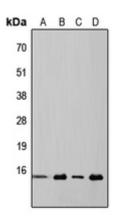
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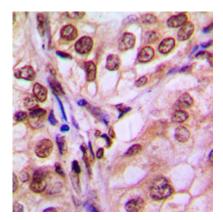




# **Product Data Sheet**



Western blot analysis of 4EBP1 expression in HeLa (A), MCF7 (B), mouse kidney (C), rat liver (D) whole cell lysates. (Predicted band size: 12 kD; Observed band size: 15 kD)



Immunohistochemical analysis of 4EBP1 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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