

## **Product Data Sheet**

### **Anti-Beta-actin Antibody**

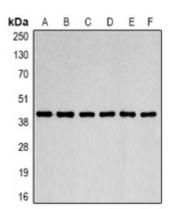
Catalog #	Source	Reactivity		Applications	
CPA9100	Mouse	H, M, R, C, D, Mk, Rb,	, Insect	WB, IH, IF/IC	
Description	Mo	use monoclonal antibody t	to Beta-actin		
Immunogen	Rec	ombinant protein correspo	onding to hur	man Beta-actin.	
Purification					
Specificity	Rec	ognizes endogenous levels	s of Beta-acti	n protein.	
Clonality	Mo	Monoclonal			
Conjugation					
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 0.2% BSA,			
	30%	s glycerol, and 0.01% sodiu	ım azide.		
Dilution	WB	(1/3000 - 1/10000), IH (1/50	0 - 1/100), IF/	IC (1/50 - 1/100)	
Gene Symbol	ACT	В			
Alternative N	ames Acti	n cytoplasmic 1; Beta-acti	n		
Entrez Gene 60 (Hu		ıman); 11461 (Mouse); 81822 (Rat)			
SwissProt	P60	709 (Human); P60710 (Mc	ouse); P60712	1 (Rat)	
Storage/Stabi	i <b>lity</b> Ship	ped at 4°C. Upon delivery	aliquot and	store at -20°C for one year. Avoid	
	free	ze/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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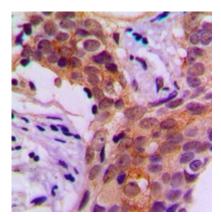




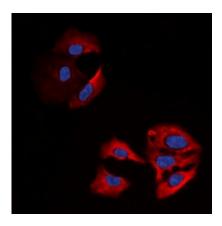
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Western blot analysis of Beta-actin expression in Jurkat (A), MCF7 (B), NIH3T3 (C), mouse brain (D), rat brain (E), COS7 (F) whole cell lysates. (Predicted band size: 41 kD; Observed band size: 42 kD)



Immunohistochemical analysis of Beta-actin staining in human prostate 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 Beta-actin staining in A549 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. DAPI was used to stain the cell nuclei (blue).

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