

# **Product Data Sheet**

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

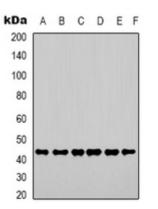
Catalog #	Source	Reactivity	Applications
CPA9245	Rabbit	H, M, R, Rb, C, Mk, S	i, X WB, IH, IF/IC
Description	Ral	obit polyclonal antibody to	) Beta-actin
Immunogen	Red	combinant protein corresp	oonding to human Beta-actin.
Purification	The	e antibody was purified by	immunogen affinity chromatography.
Specificity	Red	cognizes endogenous level	ls of Beta-actin protein.
Clonality	Pol	yclonal	
Conjugation			
Form	Liq	uid in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	d 0.01% sodium azide.	
Dilution	WE	3 (1/1000 - 1/3000), IH (1/20	00 - 1/500), IF/IC (1/100 - 1/200)
Gene Symbol	AC	ТВ	
Alternative N	ames Act	in cytoplasmic 1; Beta-acti	in
Entrez Gene	60	(Human); 11461 (Mouse);	81822 (Rat)
SwissProt	P60	0709 (Human); P60710 (M	ouse); P60711 (Rat)
Storage/Stabi	ility Shi	pped at 4°C. Upon delivery	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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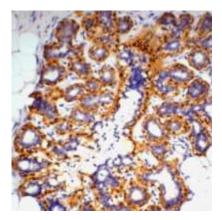




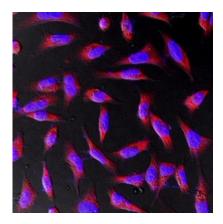
For research purposes only, not for human use

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Western blot analysis of Beta-actin expression in HepG2 (A), 293T (B), mouse kidney (C), rat liver (D), rabbit testis (E), sheep lung (F) whole cell lysates. (Predicted band size: 41 kD; Observed band size: 42 kD)



Immunohistochemical analysis of Beta-actin staining in human ovary 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 Hela 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 hidified 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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