

Product Data Sheet

Anti-14-3-3 eta Antibody

Catalog #	Source	Reactivity	Applications	
CPA3353	Rabbit	H, M, R, B, Mk	WB, IH, IF/IC	
Description	R	Rabbit polyclonal antibody to	14-3-3 eta	
Immunogen	К	LH-conjugated synthetic per	otide encompassing a sequence within the center	
	re	egion of human 14-3-3 eta. ⁻	The exact sequence is proprietary.	
Purification	Т	he antibody was purified by	immunogen affinity chromatography.	
Specificity	R	Recognizes endogenous level	s of 14-3-3 eta protein.	
Clonality	Р	Polyclonal		
Conjugation				
Form	Li	iquid in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
	а	nd 0.01% sodium azide.		
Dilution	V	VB (1/500 - 1/1000), IH (1/100) - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	Y	WHAH		
Alternative N	a <mark>mes</mark> Y	'WHA1; 14-3-3 protein eta; F	rotein AS1	
Entrez Gene	7	7533 (Human); 22629 (Mouse); 25576 (Rat)		
SwissProt	C	204917 (Human); P68510 (M	ouse); P68511 (Rat)	
Storage/Stabi	lity S	hipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid	
	fr	reeze/thaw cycles.		

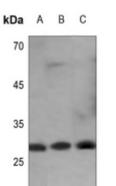
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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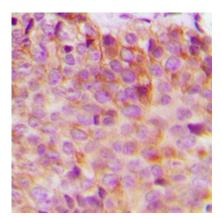
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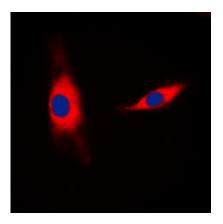
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Western blot analysis of 14-3-3 eta expression in Hela (A), mouse lung (B), rat lung (C) whole cell lysates. (Predicted band size: 28 kD; Observed band size: 28 kD)



Immunohistochemical analysis of 14-3-3 eta 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 14-3-3 eta staining in MCF7 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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