

Product Data Sheet

Anti-RIN1 Antibody

Catalog #	Source	Reactivity	Applications
CPA2408	Rabbit	H, M, R	WB, IH, IF/IC
Description	Rab	bit polyclonal antibody	to RIN1
Immunogen	KLH	I-conjugated synthetic p	eptide encompassing a sequence within the C-term
	regi	ion of human RIN1. The	exact sequence is proprietary.
Purification	The	antibody was purified	by immunogen affinity chromatography.
Specificity	Rec	ognizes endogenous lev	vels of RIN1 protein.
Clonality	Poly	yclonal	
Conjugation			
Form	Liqu	uid in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1/500 - 1/1000), IH (1/5	0 - 1/100), IF/IC (1/50 - 1/200)
Gene Symbol	RIN	1	
Alternative Na	ames Ras	and Rab interactor 1; R	as inhibitor JC99; Ras interaction/interference protein 1
Entrez Gene	961	.0 (Human); 225870 (M	ouse); 207119 (Rat)
SwissProt	Q13	3671 (Human); Q921Q7	(Mouse); P97680 (Rat)
Storage/Stabi	lity Ship	oped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid
	free	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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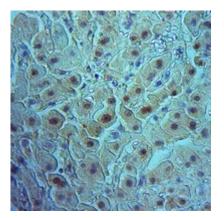
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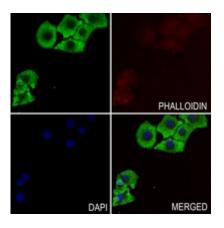
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Western blot analysis of RIN1 expression in U87MG (A), Panc1 (B) whole cell lysates. (Predicted band size: 84 kD; Observed band size: 90 kD)



Immunohistochemical analysis of RIN1 staining in human liver 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 RIN1 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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