

Product Data Sheet

Anti-RPL39 Antibody

Catalog #	Source	Reactivity	Applications
CPA4213	Rabbit	H, M, R, B, C	WB, IH, IF/IC
Description	Rab	bit polyclonal antibody	to RPL39
Immunogen	KLH	-conjugated synthetic p	eptide encompassing a sequence within the N-term
	regi	on of human RPL39. The	e exact sequence is proprietary.
Purification	The	antibody was purified b	by immunogen affinity chromatography.
Specificity	Reco	ognizes endogenous lev	els of RPL39 protein.
Clonality	Poly	vclonal	
Form	Liqu	iid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1/500 - 1/1000), IH (1/10	00 - 1/200), IF/IC (1/100 - 1/500)
Gene Symbol	RPL	39	
Alternative Na	ames 60S	ribosomal protein L39	
Entrez Gene	617	0 (Human); 67248 (Mou	ise); 25347 (Rat)
SwissProt	P62	891 (Human); P62892 (I	Mouse); P62893 (Rat)
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ry 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

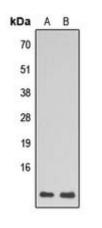
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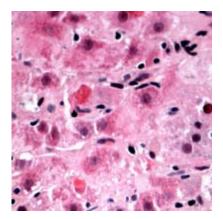
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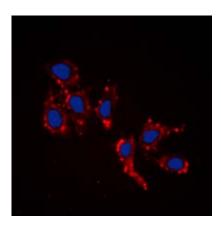
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Western blot analysis of RPL39 expression in HL60 (A), HepG2 (B) whole cell lysates.



Immunohistochemical analysis of RPL39 staining in human liver 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of RPL39 staining in HepG2 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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