

Product Data Sheet

Anti-RPL39 Antibody

Catalog #	Source	Reactivity	Applications	
CPA4213	Rabbit	H, M, R, B, C	WB, IH, IF/IC	
Description		Rabbit polyclonal antibody to	RPL39	
Immunogen		KLH-conjugated synthetic pep	tide encompassing a sequence within the N-term	
		region of human RPL39. The e	exact sequence is proprietary.	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous levels	s of RPL39 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium pho	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/1000), IH (1/100	- 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol		RPL39		
Alternative N	ames	60S ribosomal protein L39		
Entrez Gene		6170 (Human); 67248 (Mouse); 25347 (Rat)		
SwissProt		P62891 (Human); P62892 (Mo	ouse); P62893 (Rat)	
Storage/Stabi	ility	Shipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid	
		freeze/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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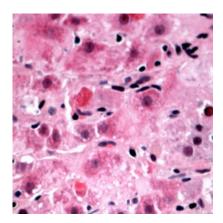
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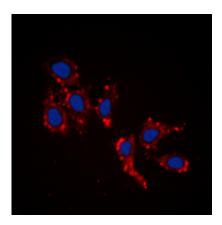
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Western blot analysis of RPL39 expression in HL60 (A), HepG2 (B) whole cell lysates. (Predicted band size: 6 kD; Observed band size: 6 kD)



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 conjugated 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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