

Product Data Sheet

Anti-RPL28 Antibody

Catalog #	Source	Reactivity	Applications			
CPA2022	Rabbit	H, M, R, Mk	WB, IH			
Description	Rabl	Rabbit polyclonal antibody to RPL28				
Immunogen	KLH-	conjugated synthetic pe	otide encompassing a sequence within the center			
	regio	on of human RPL28. The	exact sequence is proprietary.			
Purification	The	antibody was purified by	immunogen affinity chromatography.			
Specificity	Reco	ognizes endogenous leve	s of RPL28 protein.			
Clonality	Poly	clonal				
Conjugation						
Form	Liqu	id in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	0.01% sodium azide.				
Dilution	WB	(1/500 - 1/1000), IH (1/50	- 1/100)			
Gene Symbol	RPL2	28				
Alternative Na	ames 60S	ribosomal protein L28				
Entrez Gene	6158	3 (Human)				
SwissProt	P467	779 (Human)				
Storage/Stabi	lity Ship	ped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid			
	freez	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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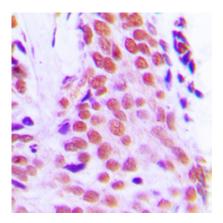
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KDa <u>A</u> <u>B</u> <u>C</u> <u>D</u> <u>E</u> <u>F</u> <u>G</u> 72 55 34 26 17 For research purposes only, not for human use

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Western blot analysis of RPL28 expression in HEK293T (A), Hela (B), H446 (C), mouse liver (D), mouse kidney (E), rat liver (F), rat kidney (G) whole cell lysates. (Predicted band size: 15 kD; Observed band size: 16 kD)



Immunohistochemical analysis of RPL28 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.

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