

Product Data Sheet

Anti-MRPL44 Antibody

Catalog #	Source	Reactivity	Applications		
CPA6562	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Ra	Rabbit polyclonal antibody to MRPL44			
Immunogen	KL	H-conjugated synthetic p	eptide encompassing a sequence within the center		
	re	gion of human MRPL44. T	he exact sequence is proprietary.		
Purification	Th	e antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous lev	els of MRPL44 protein.		
Clonality	Ро	lyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	d 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/1000), IH (1/50) - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	М	RPL44			
Alternative Na	ames 39	S ribosomal protein L44 r	nitochondrial; L44mt; MRP-L44		
Entrez Gene		65080 (Human); 69163 (Mouse)			
SwissProt	Q	Q9H9J2 (Human); Q9CY73 (Mouse)			
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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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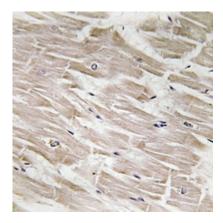
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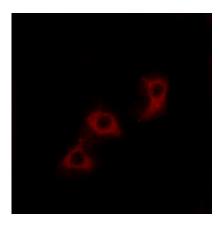
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Western blot analysis of MRPL44 expression in mouse liver (A), rat kidney (B) whole cell lysates. (Predicted band size: 37 kD; Observed band size: 37 kD)



Immunohistochemical analysis of MRPL44 staining in human heart 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 MRPL44 staining in HuvEc 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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