

Product Data Sheet

Anti-MRPL15 Antibody

Catalog #	Source	Reactivity	Applications		
CPA2637	Rabbit	H, M, Mk	WB, IH, IF/IC		
Description	Rat	obit polyclonal antibody	to MRPL15		
Immunogen	KLF	I-conjugated synthetic p	eptide encompassing a sequence within the center		
	reg	region of human MRPL15. The exact sequence is proprietary.			
Purification	The	e antibody was purified l	oy immunogen affinity chromatography.		
Specificity	Rec	cognizes endogenous lev	els of MRPL15 protein.		
Clonality	Pol	yclonal			
Conjugation					
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	d 0.01% sodium azide.			
Dilution	WB	3 (1/500 - 1/1000), IH (1/5	0 - 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	MR	RPL15			
Alternative Na	ames 395	S ribosomal protein L15	mitochondrial; L15mt; MRP-L15		
Entrez Gene	290	088 (Human); 27395 (Mo	buse)		
SwissProt	Q91	P015 (Human); Q9CPR5	(Mouse)		
Storage/Stabi	lity Shi	pped at 4°C. Upon delive	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

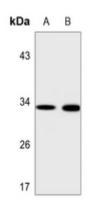
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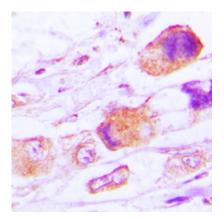
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For research purposes only, not for human use

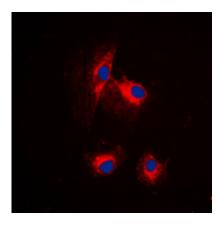
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Western blot analysis of MRPL15 expression in MCF7 (A), K562 (B) whole cell lysates. (Predicted band size: 33 kD; Observed band size: 33 kD)



Immunohistochemical analysis of MRPL15 staining in human lung 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 MRPL15 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 humidified 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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