

Product Data Sheet

Anti-MRPS36 Antibody

Catalog #	Source	Reactivity	Applications			
CPA5971	Rabbit	H, M, R	WB, IH			
Description	R	Rabbit polyclonal antibody to MRPS36				
Immunogen	К	ptide encompassing a sequence within the N-term				
	re	region of human MRPS36. The exact sequence is proprietary.				
Purification	T	The antibody was purified by immunogen affinity chromatography.				
Specificity	R	els of MRPS36 protein.				
Clonality	P	olyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	a	nd 0.01% sodium azide.				
Dilution	W	VB (1/500 - 1/1000), IH (1/50	- 1/200)			
Gene Symbol	Ν	1RPS36				
Alternative Names		28S ribosomal protein S36 mitochondrial; MRP-S36; S36mt				
Entrez Gene		92259 (Human); 66128 (Mouse)				
SwissProt P82909 (Human); Q9CQX8		82909 (Human); Q9CQX8 (Mouse)			
Storage/Stabi	lity Sl	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid				
	fr	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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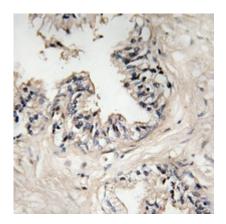
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kDa <u>A</u> <u>B</u> <u>C</u> 55 43 34 26 17 10 Western blot analysis of MRPS36 expression in rat brain (A), mouse kidney (B), rat kidney (C) whole cell lysates. (Predicted band size: 11 kD; Observed band size: 17 kD)



Immunohistochemical analysis of MRPS36 staining in human prostate 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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