

Product Data Sheet

Anti-GRP94 Antibody

Catalog #	Source	Reactivity	Applications		
CPA2195	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Ra	abbit polyclonal antibody	o GRP94		
Immunogen	KI	LH-conjugated synthetic po	eptide encompassing a sequence within the C-term		
	re	egion of human GRP94. Th	e exact sequence is proprietary.		
Purification	Tł	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous lev	els of GRP94 protein.		
Clonality	Clonality Polyclonal				
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IH (1/10	0 - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	H	SP90B1			
Alternative Na	ames G	RP94; TRA1; Endoplasmin	94 kDa glucose-regulated protein; GRP-94; Heat shock		
	рі	rotein 90 kDa beta membe	r 1; Tumor rejection antigen 1; gp96 homolog		
Entrez Gene		7184 (Human)			
SwissProt	P:	14625 (Human)			
Storage/Stabi	lity Sł	hipped at 4°C. Upon delive	ry 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

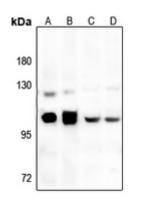
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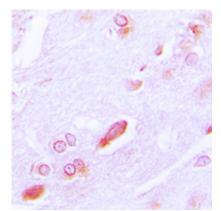


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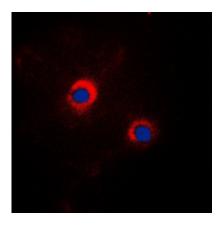
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Western blot analysis of GRP94 expression in K562 (A), A549 (B), SP20 (C), PMVEC (D) whole cell lysates. (Predicted band size: 92 kD; Observed band size: 100 kD)



Immunohistochemical analysis of GRP94 staining in human brain 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 GRP94 staining in NIH3T3 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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