

Product Data Sheet

Anti-GRP78 Antibody

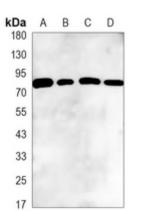
Catalog #	Source	Reactivity	Applications
CQA8406	Mouse	H, R	WB, IH, IF/IC
Description	٦	Mouse monoclonal antiboc	y to GRP78
Immunogen	F	Purified recombinant huma	n BiP/GRP78 (C-terminus) protein fragments expressed in
	E	E.coli.	
Purification	T	The antibody was purified b	by immunogen affinity chromatography.
Specificity	F	Recognizes endogenous lev	els of GRP78 protein.
Clonality	٦	Monoclonal	
Conjugation			
Form	L	Liquid in PBS containing 50 ⁴	% glycerol, 0.5% BSA and 0.02% sodium azide, pH 7.3.
Dilution	١	WB (1/500 - 1/1000), IH (1/5	0 - 1/100), IF/IC (1/50 - 1/100)
Gene Symbol	ŀ	HSPA5	
Alternative N	ames (GRP78; 78 kDa glucose-regi	ulated protein; GRP-78; Endoplasmic reticulum lumenal
	(Ca(2+)-binding protein grp7	'8; Heat shock 70 kDa protein 5; Immunoglobulin heavy
	C	chain-binding protein; BiP	
Entrez Gene	3	3309 (Human); 25617 (Rat)	
SwissProt	F	P11021 (Human); P06761 (I	Rat)
Storage/Stabi	ility S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid
	f	freeze/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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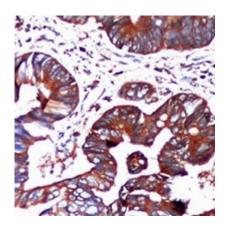




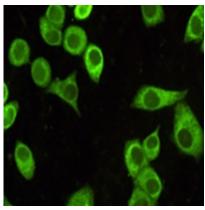
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Western blot analysis of GRP78 expression in Hela (A), C6 (B), Lncap (C), MDAMB468 (D) whole cell lysates. (Predicted band size: 72 kD; Observed band size: 78 kD)



Immunohistochemical analysis of GRP78 staining in human colon 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 GRP78 staining in HeLa 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark.

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