

Product Data Sheet

Anti-GP73 Antibody

Catalog #	Source	Reactivity	Applications
CQA1561	Rabbit	H, M, R	WB, IH, IF/IC
Description	Rab	bit polyclonal antibody	to GP73
Immunogen	Reco	ombinant full length pro	otein of human GP73
Purification	The	antibody was purified I	by immunogen affinity chromatography.
Specificity	Reco	ognizes endogenous lev	els of GP73 protein.
Clonality	Poly	/clonal	
Conjugation			
Form	Liqu	id in 0.42% Potassium إ	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1/500 - 1/2000), IH (1/5	0 - 1/200), IF/IC (1/50 - 1/100)
Gene Symbol	GOL	-M1	
Alternative Na	ames C9o	rf155; GOLPH2; Golgi m	embrane protein 1; Golgi membrane protein GP73; Golgi
	pho	sphoprotein 2	
Entrez Gene	512	80 (Human)	
SwissProt	Q8N	NBJ4 (Human)	
Storage/Stabi	lity Ship	oped 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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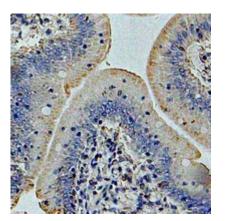
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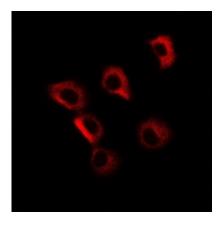
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Western blot analysis of GP73 expression in SKOV3 (A), SHSY5Y (B) whole cell lysates. (Predicted band size: 44; 45 kD; Observed band size: 75 kD)



Immunohistochemical analysis of GP73 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 GP73 staining in MCF7 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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