

Product Data Sheet

Anti-ATG5 Antibody

Catalog #	Source	Reactivity	Applications		
CQA2332	Rabbit	H, M, R	WB, IH, IF/IC		
Description		Rabbit polyclonal antibody	to ATG5		
Immunogen		Recombinant full length pro	otein of human ATG5		
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous levels of ATG5 protein.			
Clonality		Polyclonal			
Conjugation					
Form	rm Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycero				
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/2000), IH (1/50	0 - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol		ATG5			
Alternative Names		APG5L; ASP; Autophagy protein 5; APG5-like; Apoptosis-specific protein			
Entrez Gene		9474 (Human); 11793 (Mouse)			
SwissProt		Q9H1Y0 (Human); Q99J83 (Mouse); Q3MQ06 (Rat)			
Storage/Stabi	ility	Shipped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
		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

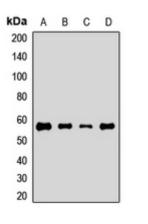
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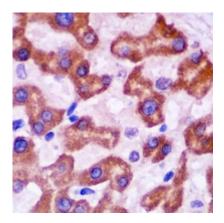


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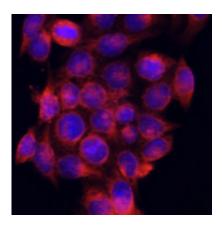
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Western blot analysis of ATG5 expression in Hela (A), NIH3T3 (B), PC12 (C), rat bain (D) whole cell lysates. (Predicted band size: 22; 32 kD; Observed band size: 55 kD)



Immunohistochemical analysis of ATG5 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.



Immunofluorescent analysis of ATG5 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 humidified chamber. Cells were washed with PBST and incubated with a AF594-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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