

# **Product Data Sheet**

### Recombinant Anti-LC3-I/II Rabbit mAb

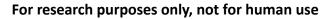
Catalog #	Source	e Reactivity	Applications		
CMA4223	Rabbit	H, M, R	WB, IF/IC		
Description		Recombinant rabbit monoclonal antibody to LC3-I/II			
Immunogen		KLH-conjugated synthetic pe	otide encompassing a sequence within human LC3-I/II.		
		The exact sequence is proprie	etary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	s of LC3-I/II protein		
Clonality		Monoclonal			
Conjugation					
Form		Liquid in PBS containing 50%	glycerol, 0.2% BSA and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/1000), IF/IC (1/5	60 - 1/200)		
Gene Symbol		MAP1LC3A			
Alternative Names		Microtubule-associated proteins 1A/1B light chain 3A; Autophagy-related protein			
		LC3 A; Autophagy-related ub	quitin-like modifier LC3 A; MAP1 light chain 3-like		
		protein 1; MAP1A/MAP1B lig	ht chain 3 A; MAP1A/MAP1B LC3 A;		
		Microtubule-associated prote	ein 1 lig		
Entrez Gene		84557 (Human); 66734 (Mou	se); 362245 (Rat)		
SwissProt		Q9H492 (Human); Q91VR7 (I	Mouse); Q6XVN8 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y 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

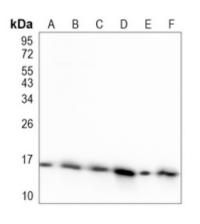
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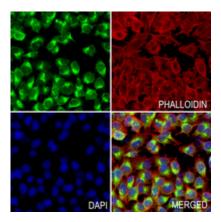




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Western blot analysis of LC3-I/II expression in HEK293T (A), Hela (B), THP1 (C), mouse brain (D), mouse liver (E), rat brain (F) whole cell lysates. (Predicted band size: 14 kD; Observed band size: 16 kD)



Immunofluorescent analysis of LC3-I/II staining in A375 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 an AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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