# MAP1LC3A Rabbit mAb

Catalog No: #48865

Package Size: #48865-1 50ul #48865-2 100ul



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Description	
Product Name	MAP1LC3A Rabbit mAb
Host Species	Recombinant Rabbit
Clonality	Monoclonal antibody
Clone No.	ST47-03
Purification	ProA affinity purified
Applications	WB, ICC/IF, IHC, IP, FC
Species Reactivity	Hu, Ms, Rt
Immunogen Description	recombinant protein
Other Names	ATG8E antibody Autophagy-related protein LC3 A antibody Autophagy-related ubiquitin-like modifier LC3 A
	antibody LC3 antibody LC3A antibody MAP1 light chain 3 like protein 1 antibody MAP1 light chain 3-like
	protein 1 antibody MAP1A/1B light chain 3 A antibody MAP1A/MAP1B LC3 A antibody MAP1A/MAP1B light
	chain 3 A antibody MAP1ALC3 antibody MAP1BLC3 antibody Map1lc3a antibody Microtubule associated
	proteins 1A/1B light chain 3 antibody Microtubule-associated protein 1 light chain 3 alpha antibody
	Microtubule-associated proteins 1A and 1B, light chain 3 antibody Microtubule-associated proteins 1A/1B light
	chain 3A antibody MLP3A_HUMAN antibody
Accession No.	Swiss-Prot#:Q9H492
Uniprot	Q9H492
GenelD	84557;
Calculated MW	14 kDa
Formulation	1*TBS (pH7.4), 1%BSA, 40%Glycerol. Preservative: 0.05% Sodium Azide.
Storage	Store at -20°C

## **Application Details**

WB: 1:1,000-1:2,000 IHC: 1:50-1:200 ICC: 1:50-1:200FC: 1:50-1:100

## Images



Western blot analysis of MAP1LC3A on different lysates using anti-MAP1LC3A antibody at 1/1,000 dilution. Positive control: Lane 1: SHG-44 Lane 2: Mouse brain Lane 3: Mouse liver Lane 4: Mouse skeletal muscle



Immunohistochemical analysis of paraffin-embedded human liver tissue using anti-MAP1LC3A antibody. Counter stained with hematoxylin.

Immunohistochemical analysis of paraffin-embedded mouse liver tissue using anti-MAP1LC3A antibody. Counter stained with hematoxylin.

![](_page_1_Picture_3.jpeg)

Immunohistochemical analysis of paraffin-embedded mouse brain tissue using anti-MAP1LC3A antibody. Counter stained with hematoxylin.

![](_page_1_Picture_5.jpeg)

ICC staining MAP1LC3A in Hela cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.

![](_page_1_Picture_7.jpeg)

ICC staining MAP1LC3A in PC12 cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.

![](_page_2_Figure_0.jpeg)

ICC staining MAP1LC3A in HUVEC cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.

![](_page_2_Figure_2.jpeg)

Flow cytometric analysis of SH-SY-5Y cells with MAP1LC3A antibody at 1/50 dilution (blue) compared with an unlabelled control (cells without incubation with primary antibody; red). Alexa Fluor 488-conjugated goat anti rabbit IgG was used as the secondary antibody.

#### Background

Microtubules, the primary component of the cytoskeletal network, interact with proteins called microtubule-associated proteins (MAPs). The microtubule-associated proteins can be divided into two groups, structural and dynamic. The structural microtubule-associated proteins, MAP-1A, MAP-1B, MAP-2A, MAP-2B and MAP-2C, stimulate tubulin assembly, enhance micro-tubule stability and influence the spatial distribution of microtubules within cells. Both MAP-1 and, to a greater extent, MAP-2 have been implicated as agents of microtubule depolymerization by suppressing the dynamic instability of the microtubules. The suppression of microtubule dynamic instability by the MAP proteins is thought to be associated with phosphorylation of the MAPs.

#### References

Note: This product is for in vitro research use only