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# [KD Validated] LC3B Rabbit mAb

Catalog No.: A19665 Recombinant 192 Publications



## **Basic Information**

**Observed MW** 14kDa/16kDa

**Calculated MW** 15kDa

Category Primary antibody

Applications WB,IHC-P,IF/ICC,IP,ELISA

**Cross-Reactivity** Human, Mouse, Rat

**CloneNo number** ARC0144

# Background

The product of this gene is a subunit of neuronal microtubule-associated MAP1A and MAP1B proteins, which are involved in microtubule assembly and important for neurogenesis. Studies on the rat homolog implicate a role for this gene in autophagy, a process that involves the bulk degradation of cytoplasmic component.

## **Recommended Dilutions**

WB	1:1000 - 1:4000
IHC-P	1:100 - 1:500
IF/ICC	1:200 - 1:800
IP	0.5µg-4µg antibody for 200µg-400µg extracts of whole cells
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

Contact

6	400-999-6126
$\times$	cn.market@abclonal.com.cn
€	www.abclonal.com.cn

# Immunogen Information

#### Gene ID 81631

Swiss Prot Q9GZQ8

#### Immunogen

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

Synonyms

LC3B; ATG8F; MAP1LC3B-a; MAP1A/1BLC3; 3B

# **Product Information**

Source Rabbit

Isotype lgG

**Purification** Affinity purification

#### Storage

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS containing 50% glycerol and 0.05% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.

### Validation Data



Western blot analysis of lysates from wild type(WT) and LC3B knockdown (KD) 293T cells, using [KD Validated] LC3B Rabbit mAb (A19665) at 1:1000 dilution. wild type(WT) and LC3B knockdown (KD) 293T cells were treated with Chloroquine (50  $\mu$ M) at 37°C for 20 hours. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25 $\mu$ g per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 30s.



Immunoprecipitation analysis of 300 µg extracts from 293T cells using 3 µg [KD Validated] LC3B Rabbit mAb (A19665). Western blot was performed from the immunoprecipitate using LC3B antibody (A19665) at a dilution of 1:1000.



Western blot analysis of various lysates, using [KD Validated] LC3B Rabbit mAb (A19665) at 1:1000 dilution. 293T, C6 and NIH/3T3 cells were treated with Chloroquine (50  $\mu$ M) at 37 °C for 20 hours. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25 $\mu$ g per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 5s.

## Validation Data



Immunohistochemistry analysis of paraffinembedded Human brain using [KD Validated] LC3B Rabbit mAb (A19665) at dilution of 1:100 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Rat brain using [KD Validated] LC3B Rabbit mAb (A19665) at dilution of 1:100 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Confocal imaging of HeLa cells (treated with Chloroquine) and HeLa cells (untreated) using [KD Validated] LC3B Rabbit mAb (A19665, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.



Confocal imaging of C6 cells (treated with Chloroquine) and C6 cells (untreated) using [KD Validated] LC3B Rabbit mAb (A19665, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.



Confocal imaging of NIH/3T3 cells (treated with Chloroquine) and NIH/3T3 cells (untreated) using [KD Validated] LC3B Rabbit mAb (A19665, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.