

### **Product Data Sheet**

# **Anti-LC3B Antibody**

Catalog # Source Reactivity Applications

CPA9415 Rabbit H, M, R WB, IH, IF/IC

**Description** Rabbit polyclonal antibody to LC3B

Immunogen Recombinant protein corresponding to human LC3B.

**Purification** The antibody was purified by immunogen affinity chromatography.

**Specificity** Recognizes endogenous levels of LC3B protein.

**Clonality** Polyclonal

Conjugation

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

Dilution WB (1/1000 - 1/2000), IH (1/200 - 1/500), IF/IC (1/100 - 1/200)

Gene Symbol MAP1LC3B

Alternative Names MAP1ALC3; Microtubule-associated proteins 1A/1B light chain 3B;

Autophagy-related protein LC3 B; Autophagy-related ubiquitin-like modifier LC3 B;

MAP1 light chain 3-like protein 2; MAP1A/MAP1B light chain 3 B; MAP1A/MAP1B

LC3 B; Microtubule-associated protein 1 light chain 3 beta

**Entrez Gene** 81631 (Human); 67443 (Mouse); 64862 (Rat)

SwissProt Q9GZQ8 (Human); Q9CQV6 (Mouse); Q62625 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

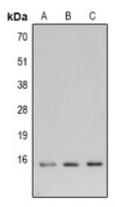
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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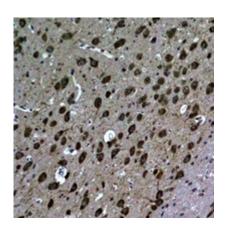
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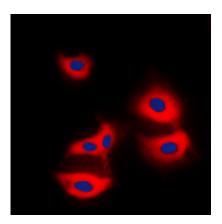
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Western blot analysis of LC3B expression in Hela (A), NIH3T3 (B), rat brain (C) whole cell lysates. (Predicted band size: 14 kD; Observed band size: 14; 16 kD)



Immunohistochemical analysis of LC3B staining in rat brain 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 LC3B 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 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. DAPI was used to stain the cell nuclei (blue).

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