Product datasheet

Anti-ATG4B Antibody

Catalog Number: *A02885-1*

BOSTER
antibody and ELISA experts
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Basic Information

Product Name	Anti-ATG4B Antibody
Gene Name	ATG4B
Source	Rabbit
Clonality	Polyclonal
Isotype	IgG
Species Reactivity	human, rat
Tested Application	WB, FCM, ELISA
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.
Immunogen	E.coli-derived human ATG4B recombinant protein (Position: M1-L392).
Concentration	500 ug/ml
Purification	Immunogen affinity purified.
Observed MW	48 kDa
Dilution Ratios	Western blot (WB): 1:500-2000 Flow Cytometry (Fixed): 1:50-200 Enzyme linked immunosorbent assay (ELISA):1:100-1000

Storage

12 months from date of receipt，-20℃ as supplied.

Background Information

Cysteine protease ATG4B is an enzyme that in humans is encoded by the ATG4B gene. Autophagy is the process by which endogenous proteins and damaged organelles are destroyed intracellularly. Autophagy is postulated to be essential for cell homeostasis and cell remodeling during differentiation, metamorphosis, non-apoptotic cell death, and aging. Reduced levels of autophagy have been described in some malignant tumors, and a role for autophagy in controlling the unregulated cell growth linked to cancer has been proposed. This gene encodes a member of the autophagin protein family. The encoded protein is also designated as a member of the C-54 family of cysteine proteases. Alternate transcriptional splice variants, encoding different isoforms, have been characterized.

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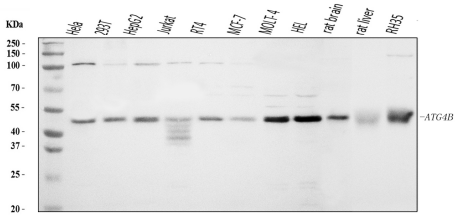
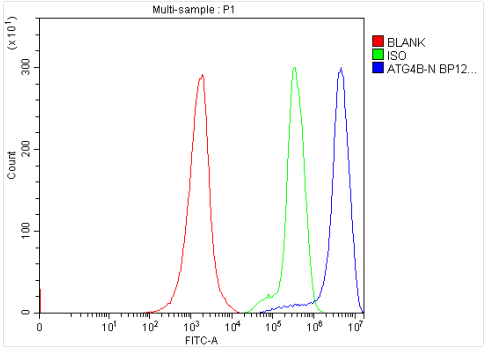
Selected Validation Data



Western blot analysis of ATG4B using anti-ATG4B antibody (A02885-1). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.
Lane 1: Hela whole cell lysates,
Lane 2: 293T whole cell lysates,
Lane 3: HepG2 whole cell lysates,
Lane 4: Jurkat whole cell lysates,
Lane 5: RT4 whole cell lysates,
Lane 6: MCF-7 whole cell lysates,
Lane 7: MOLF-4 whole cell lysates,
Lane 8: HEL whole cell lysates,
Lane 9: rat brain tissue lysates,
Lane 10: rat liver tissue lysates,
Lane 11: RH35 whole cell lysates.
After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-ATG4B antigen affinity purified polyclonal antibody (A02885-1) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for ATG4B at approximately 48 kDa. The expected band size for ATG4B is at 44 kDa.



Flow Cytometry analysis of Jurkat cells using anti-ATG4B antibody (A02885-1).
Overlay histogram showing Jurkat cells stained with A02885-1 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-ATG4B Antibody (A02885-1) at 1:100 dilution for 30 min at 20°C. Fluoro488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled

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sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.

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