BOSTER BIOLOGICAL TECHNOLOGY Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

Web: www.boster.com Phone: 027-67845390/1/2 Email: boster@boster.com

antibody and ELISA

Basic Information		
Product Name	Anti-ATF6 Antibody	
Gene Name	ATF6	
Source	Rabbit	
Clonality	Polyclonal	
lsotype	IgG	
Species Reactivity	human, mouse, rat	
Tested Application	WB, ICC/IF, FCM, ELISA	
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
Immunogen	E. coli-derived human ATF6 recombinant protein (Position: A423-Q670). Human ATF6 shares 91.9% and 93.1% amino acid (aa) sequence identity with mouse and rat ATF6, respectively.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	90-100 kDa	
Dilution Ratios	Western blot (WB): Immunocytochemistry/Immunofluorescence (ICC/IF) Flow Cytometry (Fixed): Enzyme linked immunosorbent assay (ELISA):	1:500-2000 1:1:50-400 1:50-200 1:100-1000

## **Storage**

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

## **Background Information**

ATF6, a member of the leucine zipper protein family, is an endoplasmic reticulum (ER) stress-regulated transmembrane transcription factor that activates the transcription of ER molecules. This gene is mapped to chromosome 1q23.3. ATF6 can constitutively induce the promoter of glucose-regulated protein (grp) genes through activation of the endoplasmic reticulum (ER) stress element (ERSE).

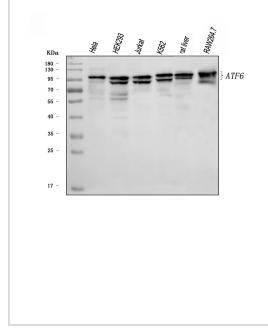
## Reference

Anti-ATF6 Antibody被引用在2文献中。

antibody and ELISA experts BOSTER BIOLOGICAL TECHNOLOGY Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

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## **Selected Validation Data**

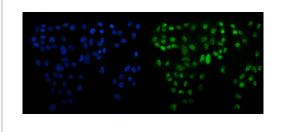


Western blot analysis of anti- ATF6 antibody (A00655-3). The sample well of each lane was loaded with 30ug of sample under reducing conditions.

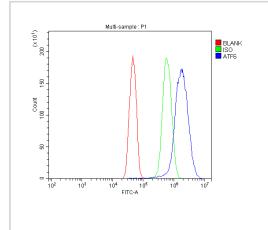
- Lane 1: human Hela whole cell lysates,
- Lane 2: human HEK293 whole cell lysates,
- Lane 3: human Jurkat whole cell lysates,
- Lane 4: human K562 whole cell lysates,
- Lane 5: rat liver tissue lysates,

Lane 6: mouse RAW264.7 whole cell lysates.

Use rabbit anti- ATF6 1:1000, probed with a goat anti-rabbit IgG-HRP secondary antibody. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog#EK1002). A specific band was detected for ATF6 at approximately 90-100KD. The expected band size for ATF6 is at 75KD.



IF analysis of ATF6 using anti-ATF6 antibody (A00655-3). ATF6 was detected in an immunocytochemical section of A431 cells. The section was incubated with rabbit anti-ATF6 Antibody (A00655-3) at a dilution of 1:100. DyLight®488 Conjugated Goat Anti-Rabbit IgG (Green) (Catalog # BA1127) was used as secondary antibody. The section was counterstained with DAPI (Catalog # AR1176) (Blue).



Flow Cytometry analysis of U251 cells using anti-ATF6 antibody (A00655-3).

Overlay histogram showing U251 cells stained with A00655-3 (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-ATF6 Antibody (A00655-3) at 1:100 dilution for 30 min at 20°C. DyLight®488 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 sample without incubation with primary



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antibody and secondary antibody (Red line) was used as a blank control.