

ATF6 Rabbit mAb

Catalog No.: A25317 **Recombinant** **2 Publications**

Basic Information

Observed MW

90kDa/75kDa/50kDa/50kDa

Calculated MW

75kDa

Category

Primary antibody

Applications

WB,IHC-P,IP,(ELISA),ChIP

Cross-Reactivity

Human, Mouse

CloneNo number

ARC3252

Background

This gene encodes a transcription factor that activates target genes for the unfolded protein response (UPR) during endoplasmic reticulum (ER) stress. Although it is a transcription factor, this protein is unusual in that it is synthesized as a transmembrane protein that is embedded in the ER. It functions as an ER stress sensor/transducer, and following ER stress-induced proteolysis, it functions as a nuclear transcription factor via a cis-acting ER stress response element (ERSE) that is present in the promoters of genes encoding ER chaperones. This protein has been identified as a survival factor for quiescent but not proliferative squamous carcinoma cells. There have been conflicting reports about the association of polymorphisms in this gene with diabetes in different populations, but another polymorphism has been associated with increased plasma cholesterol levels. This gene is also thought to be a potential therapeutic target for cystic fibrosis.

Recommended Dilutions

WB 1:2000 - 1:6000

IHC-P 1:50 - 1:200

IP 0.5µg-4µg antibody for
400µg-600µg extracts of
whole cells

ELISA Recommended starting
concentration is 1 µg/mL.
Please optimize the
concentration based on
your specific assay
requirements.

ChIP 3µg antibody for
10µg-15µg of Chromatin

Contact

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Immunogen Information

Gene ID

22926

Swiss Prot

P18850

Immunogen

Recombinant protein

Synonyms

ACHM7; ATF6A; ATF6

Product Information

Source

Rabbit

Isotype

IgG

Purification

Affinity purification

Storage

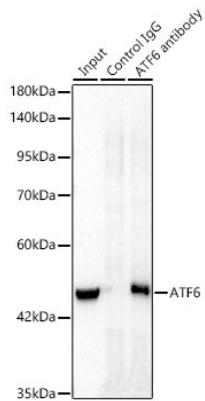
Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.05% proclin300,0.05% BSA,50% glycerol,pH7.3.

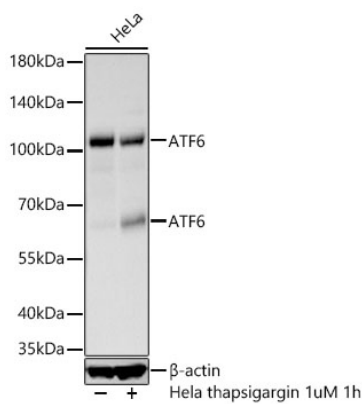


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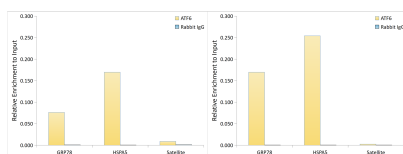
Validation Data



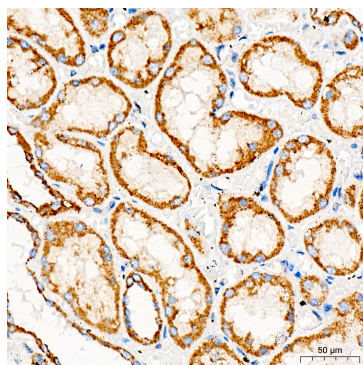
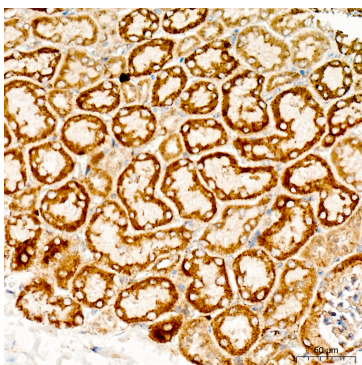
Immunoprecipitation of ATF6 from 500 µg extracts of HeLa cells was performed using 3 µg of ATF6 Rabbit mAb (A25317). Rabbit IgG isotype control(AC042) was used to precipitate the Control IgG sample. IP samples were eluted with 1X Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using ATF6 Rabbit mAb (A25317) at a dilution of 1:1000.



Western blot analysis of lysates from HeLa cells using ATF6 Rabbit mAb (A25317) at 1:5000 dilution incubated overnight at 4°C. HeLa cells were treated by tunicamycin (2 µg/ml) for 1 hours. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 30 µg per lane. Blocking buffer: 3 % nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020) Exposure time: 90 s.



Chromatin immunoprecipitation was performed with 20 µg of cross-linked chromatin from HeLa cells (left) and HeLa cells treated by thapsigargin (1uM, 1h)(right), using 2 µg of ATF6 Rabbit mAb (A25317) and Rabbit IgG isotype control (AC042). The enrichment of immunoprecipitated DNA at different genomic loci was examined by quantitative PCR. The histogram compares the ratio of the immunoprecipitated DNA to the input at given loci.



Validation Data

Immunohistochemistry analysis of paraffin-embedded Mouse kidney tissue using ATF6 Rabbit mAb (A25317) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.

Immunohistochemistry analysis of paraffin-embedded Human kidney tissue using ATF6 Rabbit mAb (A25317) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.