# [KD Validated] ATG7 Rabbit mAb

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Catalog No.: A19604 Recombinant 11 Publications

# **Basic Information**

### **Observed MW**

78kDa/78kd

### **Calculated MW**

78kDa

### Category

Primary antibody

### **Applications**

WB,IHC-P,IP,ELISA

### **Cross-Reactivity**

Human, Mouse, Rat

#### CloneNo number

ARC0083

# **Background**

This gene encodes an E1-like activating enzyme that is essential for autophagy and cytoplasmic to vacuole transport. The encoded protein is also thought to modulate p53dependent cell cycle pathways during prolonged metabolic stress. It has been associated with multiple functions, including axon membrane trafficking, axonal homeostasis, mitophagy, adipose differentiation, and hematopoietic stem cell maintenance. Alternative splicing results in multiple transcript variants.

# **Recommended Dilutions**

**WB** 1:1000 - 1:6000

1:200 - 1:800 **IHC-P** 

0.5μg-4μg antibody for ΙP 200µg-400µg extracts of

whole cells

Recommended starting **ELISA** 

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

# **Immunogen Information**

**Gene ID Swiss Prot** 10533 095352

### **Immunogen**

Synthetic peptide. This information is considered to be commercially sensitive.

### **Synonyms**

GSA7; [KD Validated] ATG7

# **Contact**

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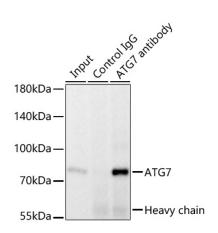
### **Product Information**

Source Isotype **Purification** Rabbit IgG Affinity purification

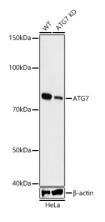
#### Storage

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

Buffer: PBS with 0.02% sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



Immunoprecipitation of ATG7 from 300  $\mu$ g extracts of HeLa cells was performed using 2  $\mu$ g of [KD Validated] ATG7 Rabbit mAb (A19604). Rabbit IgG isotype control (AC005) was used to precipitate the Control IgG sample. IP samples were eluted with 1X reducing Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using [KD Validated] ATG7 Rabbit mAb (A19604) at a dilution of 1:1000.



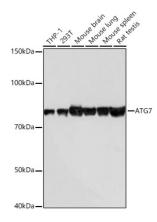
Western blot analysis of lysates from wild type(WT) and ATG7 knockdown (KD) HeLa cells using [KD Validated] ATG7 Rabbit mAb (A19604) at 1:1000 dilution incubated overnight at  $4^{\circ}$ C. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 60s.



Western blot analysis of various lysates using [KD Validated] ATG7 Rabbit mAb (A19604) at 1:1000 dilution incubated overnight at  $4^{\circ}$ C.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

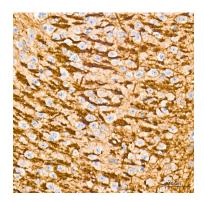
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

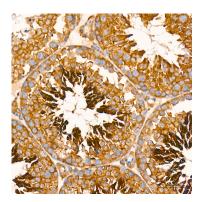
Exposure time: 10s.



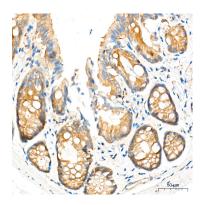
Immunohistochemistry analysis of paraffinembedded Human lung cancer tissue using [KD Validated] ATG7 Rabbit mAb (A19604) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



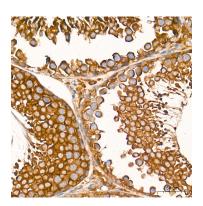
Immunohistochemistry analysis of paraffinembedded Mouse brain tissue using [KD Validated] ATG7 Rabbit mAb (A19604) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse testis tissue using [KD Validated] ATG7 Rabbit mAb (A19604) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Rat colon tissue using [KD Validated] ATG7 Rabbit mAb (A19604) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Rat testis tissue using [KD Validated] ATG7 Rabbit mAb (A19604) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.