ATG7 Rabbit mAb

Catalog No.: A21895 Recombinant 1 Publications



Basic Information

Observed MW

75kDa

Calculated MW

78kDa

Category

Primary antibody

Applications

ELISA, WB, IHC-P, IP

Cross-Reactivity

Human

CloneNo number

ARC2631

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 p53-dependent 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 1:500 - 1:1000

IHC-P 1:50 - 1:200

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

whole cells

Immunogen Information

Gene ID Swiss Prot 10533 095352

Immunogen

A synthetic peptide corresponding to a sequence within amino acids 600-703 of human ATG7 (O95352).

Synonyms

GSA7; APG7L; SCAR31; APG7-LIKE; ATG7

Contact

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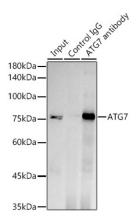
Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

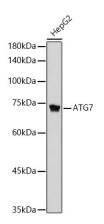
Storage

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

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



Immunoprecipitation analysis of 300 μ g extracts of HepG2 cells using 3 μ g ATG7 antibody (A21895). Western blot was performed from the immunoprecipitate using ATG7 antibody (A21895) at a dilution of 1:500.



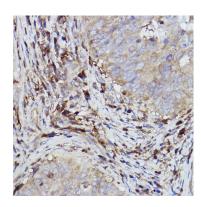
Western blot analysis of lysates from HepG2 cells, using ATG7 Rabbit mAb (A21895) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit lgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 180s.



Immunohistochemistry analysis of paraffinembedded human colon carcinoma using ATG7 Rabbit mAb (A21895) at dilution of 1:100 (40x lens).Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol.