# **Anti-Atg16L1 Antibody [JU30-62]**

#### ET7106-65



Product Type: Recombinant Rabbit monoclonal IgG, primary antibodies

Species reactivity: Human, Mouse, Rat

Applications: WB, IHC-P

Molecular Wt: Predicted band size: 68 kDa

Clone number: JU30-62

**Description:** Plays an essential role in autophagy: interacts with ATG12-ATG5 to mediate the conjugation

of phosphatidylethanolamine (PE) to LC3 (MAP1LC3A, MAP1LC3B or MAP1LC3C), to produce a membrane-bound activated form of LC3 named LC3-II. Thereby, controls the elongation of the nascent autophagosomal membrane. Regulates mitochondrial antiviral signaling (MAVS)-dependent type I interferon (IFN-I) production. Negatively regulates NOD1- and NOD2-driven inflammatory cytokine response. Plays a role in regulating

morphology and function of Paneth cell.

Immunogen: Recombinant protein within N-terminal Human Atg16L1.

Positive control: HeLa cell lysate, Jurkat cell lysate, RAW264.7 cell lysate, mouse colon tissue, PC-12 cell

lysates.

Subcellular location: Cytoplasm.

Database links: SwissProt: Q676U5 Human | Q8C0J2 Mouse

**Recommended Dilutions:** 

**WB** 1:500-1:2,000

**IHC-P** 1:500

**Storage Buffer:** 1\*TBS (pH7.4), 0.05% BSA, 40% Glycerol. Preservative: 0.05% Sodium Azide.

Storage Instruction: Store at +4°C after thawing. Aliquot store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

**Purity:** Protein A affinity purified.

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#### **Images**

kDa ve jillar 250-150-100-72-55-42-35-25-14Fig1: Western blot analysis of Atg16L1 on different lysates with Rabbit anti-Atg16L1 antibody (ET7106-65) at 1/1,000 dilution.

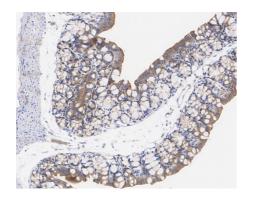
Lane 1: HeLa cell lysate Lane 2: Jurkat cell lysate Lane 3: RAW264.7 cell lysate

Lysates/proteins at 20 µg/Lane.

Predicted band size: 68 kDa Observed band size: 68/66 kDa

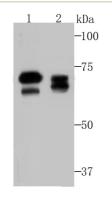
Exposure time: 2 minutes;

4-20% SDS-PAGE gel.



**Fig2:** Immunohistochemical analysis of paraffin-embedded mouse colon tissue with Rabbit anti-Atg16L1 antibody (ET7106-65) at 1/500 dilution.

The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH<sub>2</sub>O and PBS, and then probed with the primary antibody (ET7106-65) at 1/500 dilution for 1 hour at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.



**Fig3:** Western blot analysis of Atg16L1 on Hela and PC-12 cell lysates. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody (ET7106-65, 1/500) was used in 5% BSA at room temperature for 2 hours. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1:5,000 dilution was used for 1 hour at room temperature.

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