

[KO Validated] N-terminal alpha-synuclein mAb [MJFF-B-3A4]

Catalog No.: A27117 KO Validated

Basic Information

Observed MW

18kDa/

Calculated MW

14kDa

Category

Primary antibody

Applications

WB,IF/ICC,ELISA

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC5166

Background

Alpha-synuclein is a member of the synuclein family, which also includes beta- and gamma-synuclein. Alpha-synuclein is a neuronal protein that plays many roles in synaptic activity such as membrane trafficking of synaptic vesicles and neurotransmitter release. Multiplications in SNCA have been implicated in genetic forms of Parkinson disease and pathological post-translational modifications of alpha-synuclein are a hallmark of Parkinson's disease. These modifications include phosphorylation, nitration, acetylation, truncation, and aggregation.

Recommended Dilutions

WB 1:5000 - 1:30000

IF/ICC 1:200 - 1:500

ELISA Recommended starting concentration is 1 µg/mL.

Please optimize the concentration based on your specific assay requirements.

Immunogen Information

Gene ID Swiss Prot 6622 P37840

Immunogen

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

Synonyms

NACP; PARK1; α-Synuclein; SNCA

Contact

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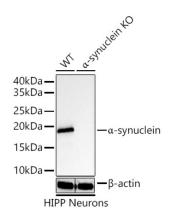
Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

Storage

Store at -20 $^{\circ}\text{C}.$ Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% Sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



Western blot analysis of lysates from wild type (WT) and N-terminal alpha-synuclein knockout (KO) HIPP Neurons cells using [KO Validated] N-terminal alpha-synuclein mAb [MJFF-B-3A4] (A27117) at 1:5000 dilution .

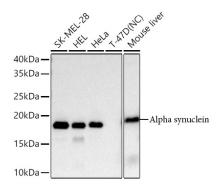
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.



Western blot analysis of various lysates using [KO Validated] N-terminal alpha-synuclein mAb [MJFF-B-3A4] (A27117) at 1:5000 dilution .

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

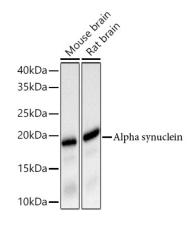
Lysates/proteins: 25 μg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Negative control (NC): T-47D

Exposure time: 60s.



Western blot analysis of various lysates using [KO Validated] N-terminal alpha-synuclein mAb [MJFF-B-3A4] (A27117) at 1:5000 dilution .

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

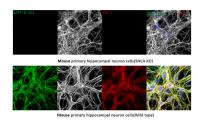
Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 3s.

Validation Data



Confocal imaging of Mouse primary hippocampal neuron cells(Wild type) and Mouse primary hippocampal neuron cells(SNCA KO) using N-terminal alphasynuclein mAb [MJFF-B-3A4] (A27117, dilution 1:500) followed by a further incubation with AF488 conjugated donkey Anti-Rabbit (H+L) (A-21206, dilution 1:500) (Green). The cells were counterstained with total aSyn antibody and MAP2 antibody followed by incubation with AF647 conjugated Donkey Anti-mouse IgG (H+L) Ab (A-31571) (Red) and AF568 conjugated Donkey Anti-chicken IgG (H+L) Ab (703-545-155, dilution 1:400) (Grey).DAPI was used for nuclear staining (Blue). Objective: 40x.