Anti-Cofilin Antibody [PSH0-33] - BSA and Azide free HA750612



Product Type: Recombinant Rabbit monoclonal IgG, primary antibodies

Species reactivity: Human, Mouse, Rat, Monkey

Applications: WB, IF-Cell, FC

Molecular Wt: Predicted band size: 19 kDa

Clone number: PSH0-33

Description: Cofilin 1 (non-muscle; n-cofilin), also known as CFL1, is a human gene, part of the

ADF/cofilin family. Cofilin is a widely distributed intracellular actin-modulating protein that binds and depolymerizes filamentous F-actin and inhibits the polymerization of monomeric G-actin in a pH-dependent manner. It is involved in the translocation of actin-cofilin complex from cytoplasm to nucleus. One group reports that reelin signaling leads to serine3-phosphorylation of cofilin-1, and this interaction may play a role in the reelin-related regulation of neuronal migration. Cofilin 1 has been shown to interact with HSPH1 and

LIMK1.

Immunogen: Synthetic peptide within human Cofilin aa 117-166 / 166.

Positive control: Hela cell lysate, HEK-293 cell lysate, MCF-7 cell lysate, MDA-MB-468 cell lysate, K-562 cell

lysate, SH-SY5Y cell lysate, HUVEC cell lysate, Jurkat cell lysate, A431 cell lysate, COS-1 cell lysate, VERO cell lysate, NIH/3T3 cell lysate, PC-12 cell lysate, Neuro-2a cell lysate,

NIH/3T3, Hela.

Subcellular location: Nucleus matrix, Cytoplasm, Cytoskeleton, Cell projection.

Database links: SwissProt: P23528 Human | P18760 Mouse | P45592 Rat

Recommended Dilutions:

WB 1:1,000 **IF-Cell** 1:100

FC 1:200-1:1,000

Storage Buffer: PBS (pH7.4).

Storage Instruction: Store at +4 $^{\circ}$ C after thawing. Aliquot store at -20 $^{\circ}$ C. Avoid repeated freeze / thaw cycles.

Purity: Protein A affinity purified.

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Images

 Fig1: Western blot analysis of Cofilin on different lysates with Rabbit anti-Cofilin antibody (HA750612) at 1/1,000 dilution.

Lane 1: Hela cell lysate (30 µg/Lane)
Lane 2: HEK-293 cell lysate (30 µg/Lane)
Lane 3: MCF-7 cell lysate (30 µg/Lane)

Lane 4: MDA-MB-468 cell lysate (24 $\mu g/Lane$)

Lane 5: K-562 cell lysate (16 µg/Lane)
Lane 6: SH-SY5Y cell lysate (22 µg/Lane)
Lane 7: HUVEC cell lysate (30 µg/Lane)
Lane 8: Jurkat cell lysate (30 µg/Lane)
Lane 9: A431 cell lysate (20 µg/Lane)

Lane 10: COS-1 cell lysate (30 µg/Lane) Lane 11: VERO cell lysate (30 µg/Lane) Lane 12: NIH/3T3 cell lysate (22 µg/Lane)

Lane 13: PC-12 cell lysate (30 µg/Lane)
Lane 14: Neuro-2a cell lysate (19 µg/Lane)

Lane 14. Neuro-za cen lysale (19 µg/L

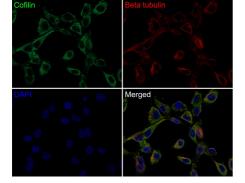
Predicted band size: 19 kDa Observed band size: 19 kDa

Exposure time: 20 seconds;

15% SDS-PAGE gel.

Proteins were transferred to a PVDF membrane and blocked with 5% NFDM/TBST for 1 hour at room temperature. The primary antibody (HA750612) at 1/1,000 dilution was used in 5% NFDM/TBST at room temperature for 2 hours. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1:200,000 dilution was used for 1 hour at room temperature.

Fig2: Immunocytochemistry analysis of NIH/3T3 cells labeling Cofilin with Rabbit anti-Cofilin antibody (HA750612) at 1/100 dilution.



Cells were fixed in 4% paraformaldehyde for 10 minutes at 37 $^{\circ}$ C, permeabilized with 0.05% Triton X-100 in PBS for 20 minutes, and then blocked with 2% negative goat serum for 30 minutes at room temperature. Cells were then incubated with Rabbit anti-Cofilin antibody (HA750612) at 1/100 dilution in 2% negative goat serum overnight at 4 $^{\circ}$ C. Goat Anti-Rabbit IgG H&L (iFluor M 488, HA1121) was used as the secondary antibody at 1/1,000 dilution. Nuclear DNA was labelled in blue with DAPI.

Beta tubulin (M1305-2, red) was stained at 1/100 dilution overnight at $+4^{\circ}$ C. Goat Anti-Mouse IgG H&L (iFluor 594, HA1126) was used as the secondary antibody at 1/1,000 dilution.

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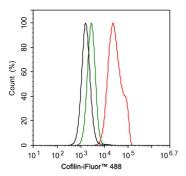


Fig3: Flow cytometric analysis of Hela cells labeling Cofilin.

Cells were fixed and permeabilized. Then stained with the primary antibody (HA750612, 1ug/ml) (red) compared with Rabbit IgG Isotype Control (green). After incubation of the primary antibody at +4 $^{\circ}$ C for an hour, the cells were stained with a iFluor 488 conjugate-Goat anti-Rabbit IgG Secondary antibody (HA1121) at 1/1,000 dilution for 30 minutes at +4 $^{\circ}$ C. Unlabelled sample was used as a control (cells without incubation with primary antibody; black).

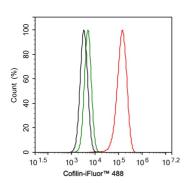


Fig4: Flow cytometric analysis of NIH/3T3 cells labeling Cofilin.

Cells were fixed and permeabilized. Then stained with the primary antibody (HA750612, 1ug/ml) (red) compared with Rabbit IgG Isotype Control (green). After incubation of the primary antibody at +4°C for an hour, the cells were stained with a iFluor™ 488 conjugate-Goat anti-Rabbit IgG Secondary antibody (HA1121) at 1/1,000 dilution for 30 minutes at +4°C. Unlabelled sample was used as a control (cells without incubation with primary antibody; black).

Note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

Background References

- 1. Namme JN et al. Cofilin Signaling in the CNS Physiology and Neurodegeneration. Int J Mol Sci. 2021 Oct
- 2. Bamburg JR et al. Cofilin and Actin Dynamics: Multiple Modes of Regulation and Their Impacts in Neuronal Development and Degeneration. Cells. 2021 Oct