

Anti-6X His tag Antibody [PSH07-10] - BSA and Azide free

HA751132



Product Type:	Recombinant Rabbit monoclonal IgG, primary antibodies
Species reactivity:	Species independent
Applications:	WB, IF-Cell, IHC-P, FC, IP, ELISA
Clone number:	PSH07-10

Description: His-Tag Antibody detects recombinant proteins containing the 6xHis epitope tag. The antibody recognizes the His-tag fused to the amino- or carboxy- termini of targeted proteins in transfected or transformed cells.

Immunogen: Synthetic peptide immune sequence is HHHHHHHHC.

Recommended Dilutions:

WB	1:5,000
IF-Cell	1:5,000
IHC-P	1:5,000
FC	1:200-500
IP	1-2µg/sample
ELISA	1:5,000

Storage Buffer: PBS (pH7.4).

Storage Instruction: Store at +4℃ after thawing. Aliquot store at -20℃. Avoid repeated freeze / thaw cycles.

Purity: Protein A affinity purified.

Hangzhou Huaan Biotechnology Co., Ltd.

Orders:0086-571-88062880

Technical:0086-571-89986345

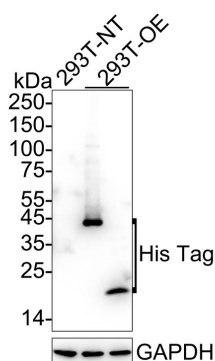
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Applications:WB=Western blot IHC-P=Immunohistochemistry (paraffin) IF-Cell=Immunofluorescence (Cell) IF-Tissue=Immunofluorescence (Tissue) FC=Flow cytometry IP=Immunoprecipitation

Images

Fig1: Western blot analysis of 6X His tag on different lysates with Rabbit anti-6X His tag antibody (HA751132) at 1/5,000 dilution.



Lane 1: 293T transfected with His-tagged empty control cell lysate
Lane 2: 293T transfected with His-tagged ACAT2 (N-terminal) cell lysate

Lane 3: 293T transfected with His-tagged Histone H3.1 (C-terminal) cell lysate

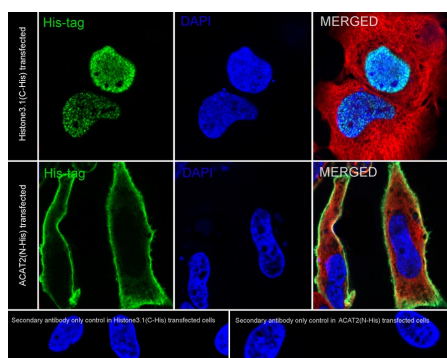
Lysates/proteins at 10 µg/Lane.

Exposure time: 2 seconds; ECL: K1801;

4-20% 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 (HA751132) at 1/5,000 dilution was used in 5% NFDM/TBST at 4°C overnight. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1/50,000 dilution was used for 1 hour at room temperature.

Fig2: Immunocytochemistry analysis of HeLa cells labeling 6X His tag with Rabbit anti-6X His tag antibody (HA751132) at 1/5,000 dilution.



HeLa cells, transfected with His-tagged Histone H3.1 (C-terminal) or ACAT2 (N-terminal) expression vector, respectively, were fixed in 4% paraformaldehyde for 10 minutes at room temperature, permeabilized with 0.1% Triton X-100 in PBS for 15 minutes at room temperature, then blocked with 1% BSA in 10% negative goat serum for 1 hour at room temperature. Cells were then incubated with Rabbit anti-6X His tag antibody (HA751132) at 1/5,000 dilution in 1% BSA in PBST overnight at 4 °C. Goat Anti-Rabbit IgG H&L (iFluor™ 488, HA1121) was used as the secondary antibody at 1/1,000 dilution. PBS instead of the primary antibody was used as the secondary antibody only control. Nuclear DNA was labelled in blue with DAPI.

Beta tubulin (M1305-2, red) was stained at 1/100 dilution overnight at +4°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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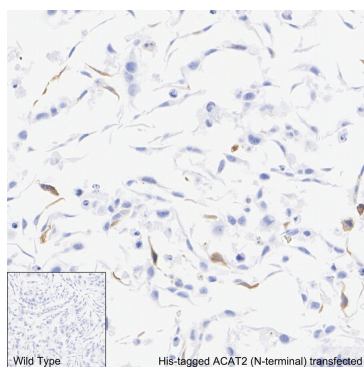


Fig3: Immunohistochemical analysis of paraffin-embedded HeLa transfected with His-tagged ACAT2 (N-terminal) cells with Rabbit anti-6X His tag antibody (HA751132) at 1/10,000 dilution.

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 ddH₂O and PBS, and then probed with the primary antibody (HA751132) at 1/10,000 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.

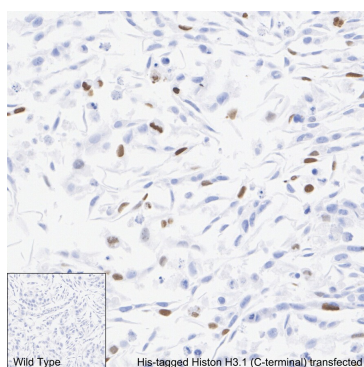


Fig4: Immunohistochemical analysis of paraffin-embedded HeLa transfected with His-tagged Histone H3.1 (C-terminal) cells with Rabbit anti-6X His tag antibody (HA751132) at 1/10,000 dilution.

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 ddH₂O and PBS, and then probed with the primary antibody (HA751132) at 1/10,000 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.

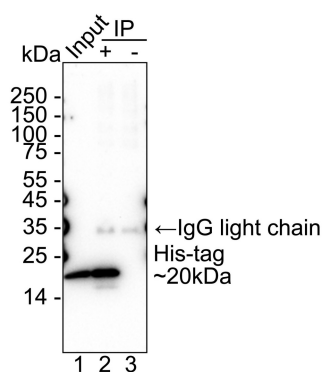


Fig5: 6X His tag was immunoprecipitated from 0.2 mg 293T transfected with His-tagged Histone H3.1 (C-terminal) cell lysate with HA751132 at 2 µg/10 µl beads. Western blot was performed from the immunoprecipitate using HA751132 at 1/500 dilution. Anti-Rabbit IgG for IP Nano-secondary antibody (NBI01H) at 1/5,000 dilution was used for 1 hour at room temperature.

Lane 1: 293T transfected with His-tagged Histone H3.1 (C-terminal) cell lysate (input)

Lane 2: HA751132 IP in 293T transfected with His-tagged Histone H3.1 (C-terminal) cell lysate

Lane 3: Rabbit IgG instead of HA751132 in 293T transfected with His-tagged Histone H3.1 (C-terminal) cell lysate

Blocking/Dilution buffer: 5% NFDM/TBST
Exposure time: 8 seconds; ECL: K1801

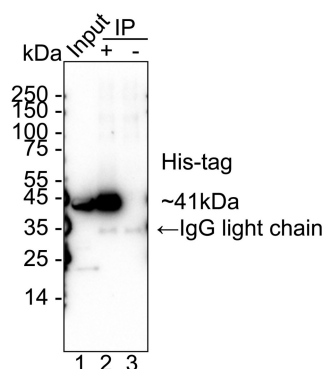


Fig6: 6X His tag was immunoprecipitated from 0.2 mg 293T transfected with His-tagged ACAT2 (N-terminal) cell lysate with HA751132 at 2 µg/10 µl beads. Western blot was performed from the immunoprecipitate using HA751132 at 1/500 dilution. Anti-Rabbit IgG for IP Nano-secondary antibody (NBI01H) at 1/5,000 dilution was used for 1 hour at room temperature.

Lane 1: 293T transfected with His-tagged ACAT2 (N-terminal) cell lysate (input)

Lane 2: HA751132 IP in 293T transfected with His-tagged ACAT2 (N-terminal) cell lysate

Lane 3: Rabbit IgG instead of HA751132 in 293T transfected with His-tagged ACAT2 (N-terminal) cell lysate

Blocking/Dilution buffer: 5% NFDM/TBST

Exposure time: 8 seconds; ECL: K1801

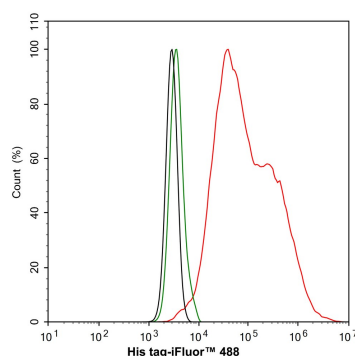


Fig7: Flow cytometric analysis of HeLa transfected with His-tagged ACAT2 (N-terminal) cells labeling 6X His tag.

Cells were fixed and permeabilized. Then stained with the primary antibody (HA751132, 1/200) (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".

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