

## Recombinant Mouse IgG1 protein [PSH04-91] - Isotype Control

# HA601336



<b>Product Type:</b>	Recombinant Mouse monoclonal IgG1, primary antibodies
<b>Applications:</b>	WB, FC, IP, CHIP
<b>Clone number:</b>	PSH04-91

**Description:** Isotype control antibodies are used to estimate the nonspecific binding of target primary antibodies due to Fc receptor binding or other protein-protein interactions. An isotype control antibody should have the same immunoglobulin type and be used at the same concentration as the test antibody.

**Immunogen:** Small molecule.

#### Recommended Dilutions:

<b>WB</b>	1:1,000
<b>FC</b>	1:1,000
<b>IP</b>	1-2 $\mu$ g/sample
<b>CHIP</b>	Use 0.5~2 $\mu$ g for 25 $\mu$ g of chromatin.

**Storage Buffer:** 1\*PBS (pH7.4), 0.1% BSA, 40% Glycerol, 0.2% Proclean 950.

**Storage Instruction:** Shipped at 4 $^{\circ}$ C. Store at +4 $^{\circ}$ C short term (1-2 weeks). It is recommended to aliquot into single-use upon delivery. Store at -20 $^{\circ}$ C long term.

**Purity:** Protein A affinity purified.

Hangzhou Huaan Biotechnology Co., Ltd.

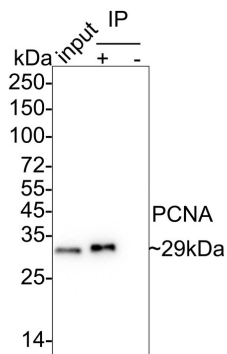
Orders:0086-571-88062880

Technical:0086-571-89986345

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## Images



**Fig1:** PCNA was immunoprecipitated from 0.2 mg HeLa cell lysate with HA601172 at 2  $\mu\text{g}/10 \mu\text{l}$  beads. Western blot was performed from the immunoprecipitate using HA601172 at 1/10,000 dilution. Anti-Mouse IgG for IP Nano-secondary antibody (NBI02H) at 1/5,000 dilution was used for 1 hour at room temperature.

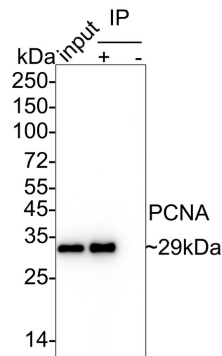
Lane 1: HeLa cell lysate (input)

Lane 2: HA601172 IP in HeLa cell lysate

Lane 3: Recombinant Mouse IgG1 (HA601336) instead of HA601172 in HeLa cell lysate

Blocking/Dilution buffer: 5% NFDM/TBST

Exposure time: 1 minute 59 seconds; ECL: K1801



**Fig2:** PCNA was immunoprecipitated from 0.2 mg NIH/3T3 cell lysate with HA601172 at 2  $\mu\text{g}/10 \mu\text{l}$  beads. Western blot was performed from the immunoprecipitate using HA601172 at 1/10,000 dilution. Anti-Mouse IgG for IP Nano-secondary antibody (NBI02H) at 1/5,000 dilution was used for 1 hour at room temperature.

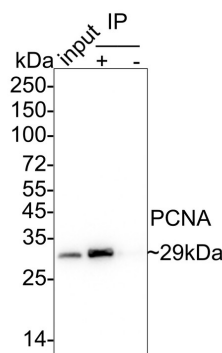
Lane 1: NIH/3T3 cell lysate (input)

Lane 2: HA601172 IP in NIH/3T3 cell lysate

Lane 3: Recombinant Mouse IgG1 (HA601336) instead of HA601172 in NIH/3T3 cell lysate

Blocking/Dilution buffer: 5% NFDM/TBST

Exposure time: 5 seconds; ECL: K1802



**Fig3:** PCNA was immunoprecipitated from 0.2 mg PC-12 cell lysate with HA601172 at 2  $\mu\text{g}/10 \mu\text{l}$  beads. Western blot was performed from the immunoprecipitate using HA601172 at 1/10,000 dilution. Anti-Mouse IgG for IP Nano-secondary antibody (NBI02H) at 1/5,000 dilution was used for 1 hour at room temperature.

Lane 1: PC-12 cell lysate (input)

Lane 2: HA601172 IP in PC-12 cell lysate

Lane 3: Recombinant Mouse IgG1 (HA601336) instead of HA601172 in PC-12 cell lysate

Blocking/Dilution buffer: 5% NFDM/TBST

Exposure time: 43 seconds; ECL: K1801

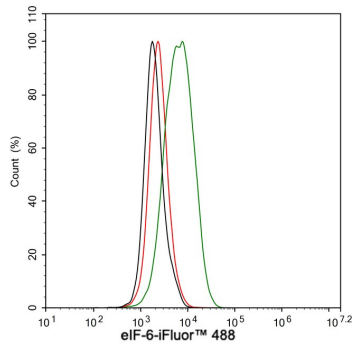
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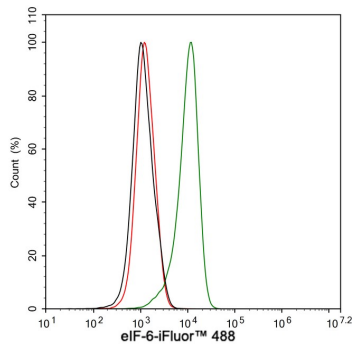
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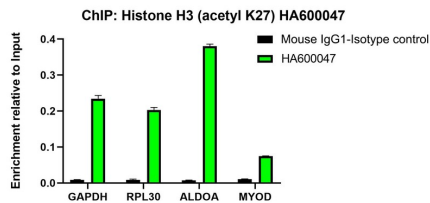
**Fig4:** Flow cytometric analysis of HeLa cells labeling eIF-6.

Cells were fixed and permeabilized. Then stained with the primary antibody (HA601293, 1µg/mL) (green) compared with Recombinant Mouse IgG1 (HA601336, 1µg/mL) (red). After incubation of the primary antibody at +4°C for an hour, the cells were stained with a iFluor™ 488 conjugate-Goat anti-Mouse IgG Secondary antibody (HA1125) 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).



**Fig5:** Flow cytometric analysis of C6 cells labeling eIF-6.

Cells were fixed and permeabilized. Then stained with the primary antibody (HA601293, 1µg/mL) (green) compared with Recombinant Mouse IgG1 (HA601336, 1µg/mL) (red). After incubation of the primary antibody at +4°C for an hour, the cells were stained with a iFluor™ 488 conjugate-Goat anti-Mouse IgG Secondary antibody (HA1125) 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).



**Fig6:** Chromatin immunoprecipitations were performed with cross-linked chromatin from HeLa cells treated with 500ng/mL TSA for 4 hours with Histone H3 (acetyl K27) (HA600047) or Recombinant Mouse IgG1 (HA601336) according to the ChIP protocol. The enriched DNA was quantified by real-time PCR using indicated primers. The amount of immunoprecipitated DNA in each sample is represented as signal relative to the total amount of input chromatin, which is equivalent to one.

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