c-Fos Recombinant Antibody [PSH07-51] - Mouse IgG1 (Chimeric) - BSA and Azide free

HA610275

Product Type: Recombinant Chimeric Antibody, primary antibodies

Species reactivity: Human, Mouse, Rat, Cynomolgus monkey, Pig

Applications: IHC-Fr, IHC-P, IF-Cell, WB

Molecular Wt: Predicted band size: 41 kDa

Clone number: PSH07-51

Description: Protein c-Fos is a proto-oncogene that is the human homolog of the retroviral oncogene v-

fos. It is encoded in humans by the FOS gene. It was first discovered in rat fibroblasts as the transforming gene of the FBJ MSV (Finkel–Biskis–Jinkins murine osteogenic sarcoma virus). It is a part of a bigger Fos family of transcription factors which includes c-Fos, FosB, Fra-1 and Fra-2. It has been mapped to chromosome region $14q21 \rightarrow q31$. c-Fos encodes a 62 kDa protein, which forms heterodimer with c-jun (part of Jun family of transcription factors), resulting in the formation of AP-1 (Activator Protein-1) complex which binds DNA at AP-1 specific sites at the promoter and enhancer regions of target genes and converts extracellular signals into changes of gene expression. It plays an important role in many

cellular functions and has been found to be overexpressed in a variety of cancers.

Immunogen: Recombinant protein within human Protein c-Fos aa 1-380.

Positive control: Mouse brain tissue, rat brain tissue.

Subcellular location: Nucleus, Endoplasmic reticulum, Cytoplasm, cytosol.

Database links: SwissProt: P01100 Human | P01101 Mouse | P12841 Rat

Recommended Dilutions:

IHC-Fr 1:1,000-1:5,000

IHC-P 1:5,000 IF-Cell 1:1.000

WB 1:5,000-1:10,000

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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Technical:0086-571-89986345

Service mail:support@huabio.cn



Images

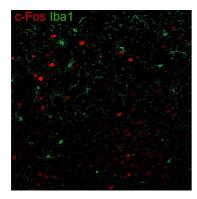


Fig1: Application: IHC-Fr

Species: Mouse

Site: Cerebral cortex (restraint stress induced)

Sample: Frozen section

Antibody concentration: 1/1,000 (c-Fos, HA610275, Mouse, red);

1/1,000 (Iba1, ET1705-78, Rabbit, green)

Antigen retrieval: Not required

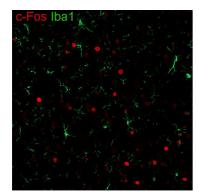


Fig2: Application: IHC-Fr

Species: Rat

Site: Cerebral cortex (restraint stress induced)

Sample: Frozen section

Antibody concentration: 1/1,000 (c-Fos, HA610275, Mouse, red);

1/1,000 (Iba1, ET1705-78, Rabbit, green)

Antigen retrieval: Not required

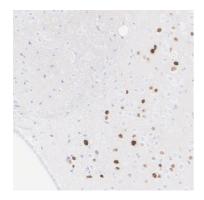


Fig3: Immunohistochemical analysis of paraffin-embedded mouse brain tissue with Mouse anti-c-Fos antibody (HA610275) at 1/5,000 dilution.

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 (HA610275) at 1/5,000 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.

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Fig4: Immunohistochemical analysis of paraffin-embedded rat brain tissue with Mouse anti-c-Fos antibody (HA610275) at 1/5,000 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 ddH $_2$ O and PBS, and then probed with the primary antibody (HA610275) at 1/5,000 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.

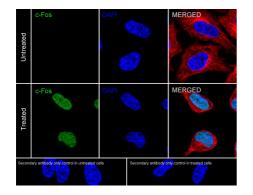


Fig5: Immunocytochemistry analysis of HeLa cells untreated / HeLa cells starved 16 hours then treated with 200nM TPA for 4 hours labeling c-Fos with Mouse anti-c-Fos antibody (HA610275) at 1/1,000 dilution.

Cells were fixed in 100% precooled methanol for 5 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 Mouse anti-c-Fos antibody (HA610275) at 1/1,000 dilution in 1% BSA in PBST overnight at 4 ℃. Goat Anti-Mouse IgG H&L (iFluor™ 488, HA1125) 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 (ET1602-4, red) was stained at 1/100 dilution overnight at $+4^{\circ}$ C. Goat Anti-Rabbit IgG H&L (iFluor † 594, HA1122) were used as the secondary antibody at 1/1,000 dilution.

Secondary withoutly unity control in until likely cells

Secondary withoutly unity control in treated cells.

Fig6: Immunocytochemistry analysis of C6 cells untreated / C6 cells starved 16 hours then treated with 200nM TPA for 4 hours labeling c-Fos with Mouse anti-c-Fos antibody (HA610275) at 1/1,000 dilution.

Cells were fixed in 100% precooled methanol for 5 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 Mouse anti-c-Fos antibody (HA610275) at 1/1,000 dilution in 1% BSA in PBST overnight at 4 $^{\circ}$ C. Goat Anti-Mouse IgG H&L (iFluor ** 488, HA1125) 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 (ET1602-4, red) was stained at 1/100 dilution overnight at $+4^{\circ}$ C. Goat Anti-Rabbit IgG H&L (iFluor 594, HA1122) were used as the secondary antibody at 1/1,000 dilution.

Fig7: Western blot analysis of c-Fos on different lysates with Mouse anti-c-Fos antibody (HA610275) at 1/5,000 dilution.

Lane 1: HeLa cell lysate

Lane 2: HeLa serum starved for 16 hours then add 200 nM TPA

for 4 hours cell lysate

Lane 3: RAW264.7 cell lysate

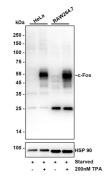
Lane 4: RAW264.7 serum starved for 16 hours then add 200 nM

TPA for 4 hours cell lysate

Lysates/proteins at 15 µg/Lane.

Predicted band size: 41 kDa Observed band size: 41-55 kDa

Exposure time: 59 seconds; ECL: K1801;



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

Background References

- 1. Matsuoka K et al. Metabolic rewiring controlled by c-Fos governs cartilage integrity in osteoarthritis. Ann Rheum Dis. 2023 Sep
- 2. Osada N et al. c-FOS is an integral component of the IKZF1 transactivator complex and mediates lenalidomide resistance in multiple myeloma. Clin Transl Med. 2023 Aug