# abcam

## Product datasheet

## Anti-c-Fos antibody [EPR20769] ab214672

Recombinant RabMAb

★★★★★ 3 Abreviews 11 References 6 Images

#### Overview

**Product name** Anti-c-Fos antibody [EPR20769]

Rabbit monoclonal [EPR20769] to c-Fos **Description** 

**Host species** Rahhit

**Tested applications** Suitable for: IP, ICC/IF, WB, IHC-P

Species reactivity Reacts with: Mouse, Human

**Predicted to work with:** Common marmoset

This product was produced with the following immunogens: **Immunogen** 

Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Recombinant fragment within Human c-Fos aa 200-300. The exact immunogen sequence used to generate this antibody is proprietary information. If additional detail on the immunogen is needed to determine the suitability of the antibody for your needs, please contact our Scientific Support

team to discuss your requirements.

Database link: P01100

Run BLAST with Run BLAST with

Positive control WB: RAW264.7, Jurkat (PMA treated) and HeLa (20% FBS 2 hours) whole cell lysates. ICC/IF:

HeLa cells.

**General notes** This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

**Properties** 

**Form** Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

**Purity** Protein A purified

Clonality Monoclonal Clone number EPR20769

Isotype ΙgG

## **Applications**

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab214672 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IP		1/40.
ICC/IF		1/500.
WB		1/1000. Detects a band of approximately 55-60 kDa (predicted molecular weight: 40 kDa).
IHC-P	<b>★★★★★</b> (3)	Use at an assay dependent concentration.

## **Target**

#### **Function**

Nuclear phosphoprotein which forms a tight but non-covalently linked complex with the JUN/AP-1 transcription factor. In the heterodimer, FOS and JUN/AP-1 basic regions each seems to interact with symmetrical DNA half sites. On TGF-beta activation, forms a multimeric

SMAD3/SMAD4/JUN/FOS complex at the AP1/SMAD-binding site to regulate TGF-betamediated signaling. Has a critical function in regulating the development of cells destined to form and maintain the skeleton. It is thought to have an important role in signal transduction, cell

proliferation and differentiation.

Sequence similarities

Belongs to the bZIP family. Fos subfamily.

Contains 1 bZIP domain.

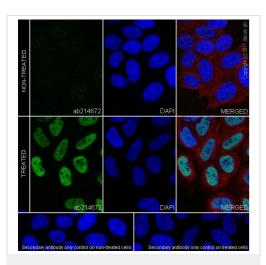
Post-translational modifications

Phosphorylated in the C-terminal upon stimulation by nerve growth factor (NGF) and epidermal growth factor (EGF). Phosphorylated, in vitro, by MAPK and RSK1. Phosphorylation on both Ser-362 and Ser-374 by MAPK1/2 and RSK1/2 leads to protein stabilization with phosphorylation on Ser-374 being the major site for protein stabilization on NGF stimulation. Phosphorylation on Ser-362 and Ser-374 primes further phosphorylations on Thr-325 and Thr-331 through promoting docking of MAPK to the DEF domain. Phosphorylation on Thr-232, induced by HA-RAS, activates the transcriptional activity and antagonizes sumoylation. Phosphorylation on Ser-362 by

RSK2 in osteoblasts contributes to osteoblast transformation.

Constitutively sumoylated by SUMO1, SUMO2 and SUMO3. Desumoylated by SENP2. Sumoylation requires heterodimerization with JUN and is enhanced by mitogen stimulation. Sumoylation inhibits the AP-1 transcriptional activity and is, itself, inhibited by Ras-activated phosphorylation on Thr-232.

#### **Images**



Immunocytochemistry/ Immunofluorescence - Antic-Fos antibody [EPR20769] (ab214672)

Immunoprecipitation - Anti-c-Fos antibody [EPR20769] (ab214672)

15 kDa 🗕

10 kDa

Immunofluorescent analysis of 4% paraformaldehyde-fixed. 0.1% Triton X-100 permeabilized serum treated and non-treated HeLa (human cervix adenocarcinoma epithelial cell) cells labeling c-Fos with ab214672 at 1/500 dilution followed by Goat Anti-Rabbit IgG H&L (Alexa Fluor<sup>®</sup> 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing weakly nuclear staining on HeLa cells grown in serum free medium for 36 hours. Expression of c-Fos increased in HeLa cells grown in serum free medium for 36 hours followed by addition of 20% FBS for 2 hours.

The nuclear counter stain is DAPI (blue). Tubulin is detected with Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor<sup>®</sup> 594) (**ab195889**) (red) at 1/200 dilution.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (Alexa Fluor<sup>®</sup> 488) (ab150077) secondary antibody at 1/1000 dilution.

c-Fos was immunoprecipitated from 0.35 mg of HeLa (human epithelial cell line from cervix adenocarcinoma) whole cell lysate with ab214672 at 1/40 dilution. Western blot was performed from the immunoprecipitate using ab214672 at 1/500 dilution. VeriBlot for IP Detection Reagent (HRP) (ab131366), was used for detection at 1/10,000 dilution

Lane 1: HeLa (human epithelial cell line from cervix adenocarcinoma) grown in serum free medium for 36 hours, followed by addition of 20%FBS for 2 hours, whole cell lysate, 10  $\mu$ g (lnput).

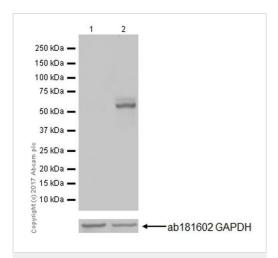
Lane 2: ab214672 IP in HeLa grown in serum free medium for 36 hours, followed by addition of 20% FBS for 2 hours, whole cell lysate.

Lane 3: Rabbit monoclonal IgG (<u>ab172730</u>) instead of ab214672 in HeLa grown in serum free medium for 36 hours, followed by addition of 20% FBS for 2 hours, whole cell lysate.

Blocking and dilution buffer and concentration: 5% NFDM/TBST.

Exposure time: 1 second.

The observed lower band is a proteasomal degradation fragment (PMID: 9737957).



Western blot - Anti-c-Fos antibody [EPR20769] (ab214672)

**All lanes :** Anti-c-Fos antibody [EPR20769] (ab214672) at 1/1000 dilution

Lane 1: Untreated HeLa (human epithelial cell line from cervix adenocarcinoma) grown in serum free medium for 36 hours, whole cell lysate

Lane 2: HeLa (human epithelial cell line from cervix adenocarcinoma) grown in serum free medium for 36 hours, followed by addition of 20% FBS for 2 hours, whole cell lysate

Lysates/proteins at 10 µg per lane.

## **Secondary**

**All lanes :** Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/50000 dilution

Developed using the ECL technique.

**Predicted band size:** 40 kDa **Observed band size:** 55-60 kDa

Exposure time: 15 seconds.

Blocking/Dilution buffer: 5% NFDM/TBST.

The expression of c-Fos is induced by the addition of 20%FBS (PMID: 24386331, PMID: 23300800, PMID: 25695333).

250 kDa — 150 kDa — 100 kDa — 75 kDa — 50 kDa — 37 kDa — 25 kDa — 20 kDa — 20 kDa — 4 LLCC (a) 10 kDa —

-ab181602 GAPDH

Western blot - Anti-c-Fos antibody [EPR20769] (ab214672)

**All lanes :** Anti-c-Fos antibody [EPR20769] (ab214672) at 1/5000 dilution

Lane 1: Untreated Jurkat (human T cell leukemia cell line from peripheral blood) grown in serum free medium overnight, whole cell lysate

**Lane 2**: Jurkat (human T cell leukemia cell line from peripheral blood) grown in serum free medium overnight, followed by treatment with 200 nM PMA for 4 hours, whole cell lysate

Lysates/proteins at 10 µg per lane.

#### Secondary

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at

1/100000 dilution

Developed using the ECL technique.

**Predicted band size:** 40 kDa **Observed band size:** 55-60 kDa

Exposure time: 3 minutes.

Blocking/Dilution buffer: 5% NFDM/TBST.

PMA treatment induces expression of c-Fos, as documented in the literature (PMID: 24386331, PMID: 23300800, PMID: 25695333).

**All lanes :** Anti-c-Fos antibody [EPR20769] (ab214672) at 1/5000 dilution

**Lane 1 :** Untreated RAW264.7 (mouse macrophage cell line transformed with Abelson murine leukemia virus grown in serum free medium overnight, whole cell lysate

Lane 2: RAW264.7 (mouse macrophage cell line transformed with Abelson murine leukemia virus) grown in serum free medium overnight, followed by treatment with 200 nM PMA for 4 hours, whole cell lysate

Lysates/proteins at 10 µg per lane.

1 2

250 kDa —
150 kDa —
150 kDa —
75 kDa —
37 kDa —
37 kDa —
37 kDa —
31 kDa —
4 20 kDa

Western blot - Anti-c-Fos antibody [EPR20769] (ab214672)

## Secondary

**All lanes :** Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/100000 dilution

Developed using the ECL technique.

Predicted band size: 40 kDa Observed band size: 55-60 kDa

Exposure time: 1 minute.

Blocking/Dilution buffer: 5% NFDM/TBST.

PMA treatment induces expression of c-Fos, as documented in the literature (PMID: 24386331, PMID: 23300800, PMID: 25695333).



Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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