

# Anti-JNK1 antibody [EPR17557] - BSA and Azide free ab251271


KO VALIDATED

Recombinant

RabMAb

10 Images

### Overview

Product name	Anti-JNK1 antibody [EPR17557] - BSA and Azide free
Description	Rabbit monoclonal [EPR17557] to JNK1 - BSA and Azide free
Host species	Rabbit
Tested applications	<b>Suitable for:</b> WB
Species reactivity	<p><b>Reacts with:</b> Mouse, Rat, Chicken, Hamster, Dog, Human, African green monkey, <i>Xenopus tropicalis</i></p> <p><b>Predicted to work with:</b> Monkey </p>
Immunogen	Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.
General notes	<p>ab251271 is the carrier-free version of <a href="#">ab199380</a>.</p> <p>Our <b>carrier-free</b> antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.</p> <p>This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.</p> <p>Use our <b>conjugation kits</b> for antibody conjugates that are ready-to-use in as little as 20 minutes with &lt;1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.</p> <p>This product is compatible with the Maxpar<sup>®</sup> Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.</p> <p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> <li>- High batch-to-batch consistency and reproducibility</li> <li>- Improved sensitivity and specificity</li> <li>- Long-term security of supply</li> <li>- Animal-free production</li> </ul> <p>For more information <a href="#">see here</a>.</p> <p>Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAb<sup>®</sup> patents</a>.</p>

## Properties

Form	Liquid
Storage instructions	Shipped at 4°C. Store at +4°C. Do Not Freeze.
Storage buffer	pH: 7.2 Constituent: PBS
Carrier free	Yes
Purity	Protein A purified
Clonality	Monoclonal
Clone number	EPR17557
Isotype	IgG

## Applications

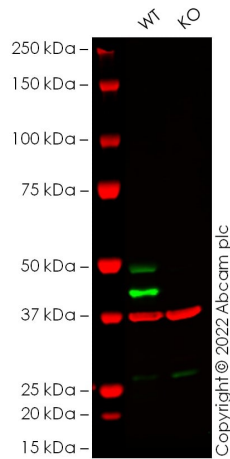
**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab251271 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		Use at an assay dependent concentration. Detects a band of approximately 54, 46 kDa (predicted molecular weight: 44 kDa).

## Target

Function	<p>Responds to activation by environmental stress and pro-inflammatory cytokines by phosphorylating a number of transcription factors, primarily components of AP-1 such as JUN, JDP2 and ATF2 and thus regulates AP-1 transcriptional activity. In T-cells, JNK1 and JNK2 are required for polarized differentiation of T-helper cells into Th1 cells (By similarity). Phosphorylates heat shock factor protein 4 (HSF4).</p> <p>JNK1 isoforms display different binding patterns: beta-1 preferentially binds to c-Jun, whereas alpha-1, alpha-2, and beta-2 have a similar low level of binding to both c-Jun or ATF2. However, there is no correlation between binding and phosphorylation, which is achieved at about the same efficiency by all isoforms.</p>
Sequence similarities	<p>Belongs to the protein kinase superfamily. CMGC Ser/Thr protein kinase family. MAP kinase subfamily.</p> <p>Contains 1 protein kinase domain.</p>
Domain	The TXY motif contains the threonine and tyrosine residues whose phosphorylation activates the MAP kinases.
Post-translational modifications	Dually phosphorylated on Thr-183 and Tyr-185, which activates the enzyme.

## Images



Western blot - Anti-JNK1 antibody [EPR17557] -  
BSA and Azide free (ab251271)

**All lanes :** Anti-JNK1 antibody [EPR17557] ([ab199380](#)) at 1/2500 dilution

**Lane 1 :** Wild-type U-2 OS cell lysate

**Lane 2 :** MAPK8 knockout U-2 OS cell lysate

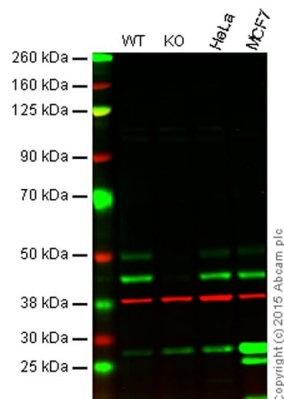
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

**Predicted band size:** 44 kDa

**Observed band size:** 42-48 kDa

False colour image of Western blot: Anti-JNK1 antibody [EPR17557] staining at 1/2500 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] ([ab8245](#)) loading control staining at 1/20000 dilution, shown in red. In Western blot, [ab199380](#) was shown to bind specifically to JNK1. A band was observed at 42/48 kDa in wild-type U-2 OS cell lysates with no signal observed at this size in mapk8 knockout cell line [ab277181](#) (knockout cell lysate [ab277223](#)). To generate this image, wild-type and mapk8 knockout U-2 OS cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 5 % milk in TBS-0.1 % Tween® 20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L 800CW and Goat anti-Mouse IgG H&L 680RD at 1/20000 dilution.



Western blot - Anti-JNK1 antibody [EPR17557] - BSA and Azide free (ab251271)

This data was developed using [ab199380](#), the same antibody clone in a different buffer formulation.

**Lane 1:** Wild-type HAP1 cell lysate (20 µg)

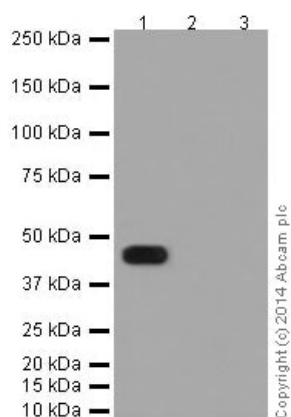
**Lane 2:** JNK1 knockout HAP1 cell lysate (20 µg)

**Lane 3:** HeLa cell lysate (20 µg)

**Lane 4:** MCF7 cell lysate (20 µg)

**Lanes 1 - 4:** Merged signal (red and green). Green - [ab199380](#) observed at 46 and 54 kDa. Red - loading control, [ab8226](#), observed at 42 kDa.

[ab199380](#) was shown to recognize JNK1 when JNK1 knockout samples were used, along with additional cross-reactive bands. Wild-type and JNK1 knockout samples were subjected to SDS-PAGE. [ab199380](#) and [ab8226](#) (loading control to beta Actin) were diluted to 1/2500 and 1/1000 respectively and incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed ([ab216773](#)) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed ([ab216776](#)) secondary antibodies at 1/10 000 dilution for 1 h at room temperature before imaging.



Western blot - Anti-JNK1 antibody [EPR17557] - BSA and Azide free (ab251271)

**All lanes :** Anti-JNK1 antibody [EPR17557] ([ab199380](#)) at 1/5000 dilution

**Lane 1 :** Full length Human JNK1 recombinant protein

**Lane 2 :** Full length Human JNK2 recombinant protein

**Lane 3 :** Full length Human JNK3 recombinant protein

Lysates/proteins at 0.01 µg per lane.

### Secondary

**All lanes :** Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 44 kDa

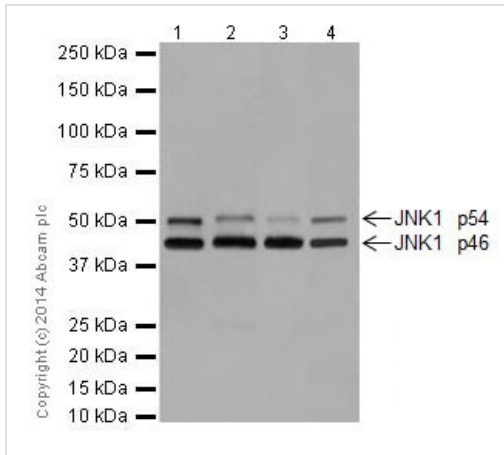
**Observed band size:** 48 kDa

This data was developed using [ab199380](#), the same antibody clone in a different buffer formulation.

**Blocking and dilution buffer:** 5% NFDm/TBST.

Recombinant full length JNK1 protein contains aa1-427 with His-Tag®. Recombinant full length JNK2 protein contains aa1-424 with

GST-tag and JNK3 protein contains aa1-464 with GST-tag.



Western blot - Anti-JNK1 antibody [EPR17557] - BSA and Azide free (ab251271)

**All lanes :** Anti-JNK1 antibody [EPR17557] ([ab199380](#)) at 1/10000 dilution

**Lane 1 :** HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate

**Lane 2 :** U937 (Human histiocytic lymphoma cells) whole cell lysate

**Lane 3 :** MCF7 (Human breast adenocarcinoma cell line) whole cell lysate

**Lane 4 :** Jurkat (Human T cell leukemia cells from peripheral blood) whole cell lysate

Lysates/proteins at 20 µg per lane.

### Secondary

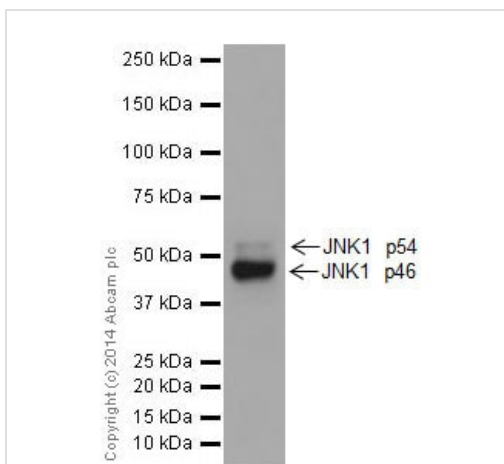
**All lanes :** Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 44 kDa

**Observed band size:** 46,54 kDa

This data was developed using [ab199380](#), the same antibody clone in a different buffer formulation.

**Blocking and dilution buffer:** 5% NFDm/TBST.



Western blot - Anti-JNK1 antibody [EPR17557] - BSA and Azide free (ab251271)

Anti-JNK1 antibody [EPR17557] ([ab199380](#)) at 1/2500 dilution + Human fetal brain lysate at 10 µg

### Secondary

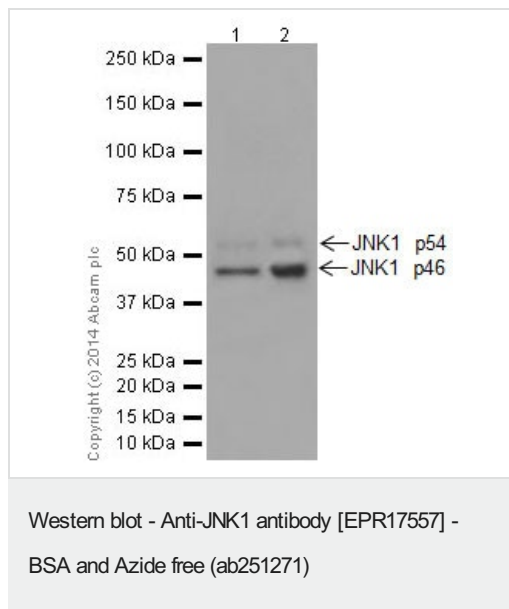
Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/1000 dilution

**Predicted band size:** 44 kDa

**Observed band size:** 46,54 kDa

This data was developed using [ab199380](#), the same antibody clone in a different buffer formulation.

**Blocking and dilution buffer:** 5% NFDm/TBST.



**All lanes :** Anti-JNK1 antibody [EPR17557] (**ab199380**) at 1/2500 dilution

**Lane 1 :** Human fetal heart lysate

**Lane 2 :** Human fetal kidney lysate

Lysates/proteins at 10 µg per lane.

#### Secondary

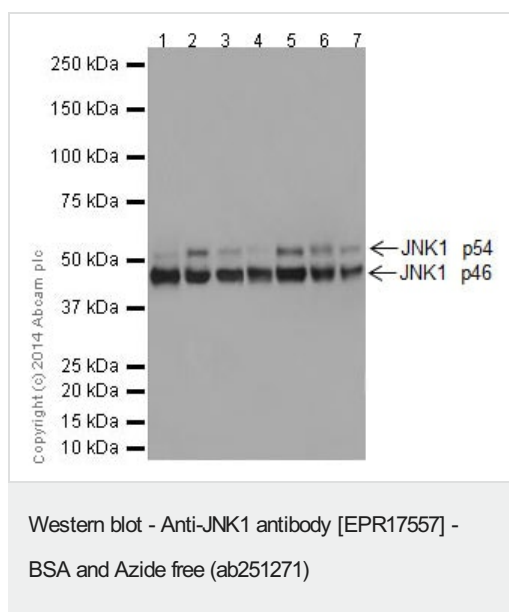
**All lanes :** Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/1000 dilution

**Predicted band size:** 44 kDa

**Observed band size:** 46,54 kDa

This data was developed using **ab199380**, the same antibody clone in a different buffer formulation.

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



**All lanes :** Anti-JNK1 antibody [EPR17557] (**ab199380**) at 1/5000 dilution

**Lane 1 :** Rat brain lysate

**Lane 2 :** Rat heart lysate

**Lane 3 :** Rat spleen lysate

**Lane 4 :** C6 (Rat glial tumor cells) whole cell lysate

**Lane 5 :** RAW 264.7 (Mouse macrophage cells transformed with Abelson murine leukemia virus) whole cell lysate

**Lane 6 :** PC-12 (Rat adrenal gland pheochromocytoma) whole cell lysate

**Lane 7 :** NIH/3T3 (Mouse embryo fibroblast cells) whole cell lysate

Lysates/proteins at 10 µg per lane.

#### Secondary

**All lanes :** Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 44 kDa

**Observed band size:** 46,54 kDa

This data was developed using **ab199380**, the same antibody clone in a different buffer formulation.

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

**All lanes :** Anti-JNK1 antibody [EPR17557] (**ab199380**) at 1/2500 dilution

**Lane 1 :** MDCK (Canine kidney cell line) whole cell lysates

**Lane 2 :** COS-1 (African green monkey kidney fibroblast-like cell line) whole cell lysates

**Lane 3 :** UMNSAH/DF-1 (Transformed chicken embryonic fibroblast cells) whole cell lysates

Lysates/proteins at 10 µg per lane.

#### Secondary

**All lanes :** Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 44 kDa

**Observed band size:** 46,54 kDa

This data was developed using **ab199380**, the same antibody clone in a different buffer formulation.

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

**All lanes :** Anti-JNK1 antibody [EPR17557] (**ab199380**) at 1/2500 dilution

**Lane 1 :** Xenopus (X. tropicalis) muscle lysates

**Lane 2 :** BHK (Hamster kidney fibroblast cells) whole cell lysates

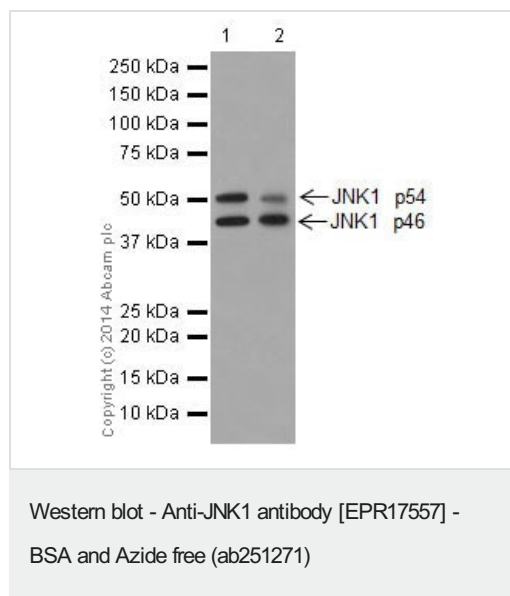
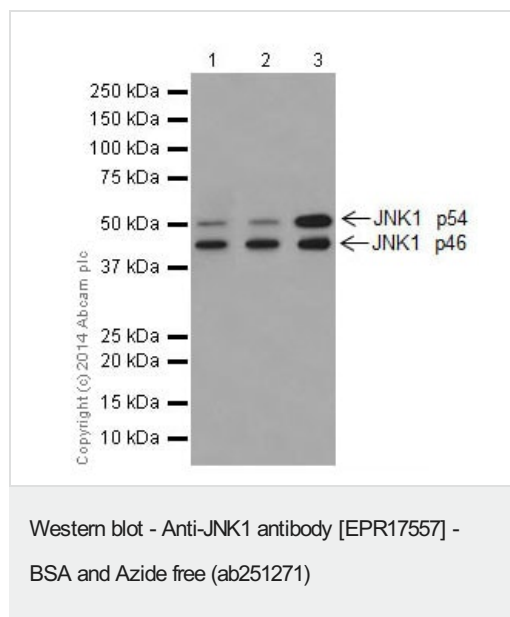
Lysates/proteins at 10 µg per lane.

#### Secondary

**All lanes :** Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 44 kDa

**Observed band size:** 46,54 kDa



This data was developed using **ab199380**, the same antibody clone in a different buffer formulation.

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

Why choose a recombinant antibody?

 <p><b>Research with confidence</b> Consistent and reproducible results</p>	 <p><b>Long-term and scalable supply</b> Recombinant technology</p>
 <p><b>Success from the first experiment</b> Confirmed specificity</p>	 <p><b>Ethical standards compliant</b> Animal-free production</p>

Anti-JNK1 antibody [EPR17557] - BSA and Azide free (ab251271)

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

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