

## ARG66184 anti-SAPK / JNK phospho (Thr183) antibody

Package: 100 μg Store at: -20°C

# Summary

| Product Description | Rabbit Polyclonal antibody recognizes SAPK / JNK phospho (Thr183)  |
|---------------------|--|
| Tested Reactivity   | Hu, Ms, Rat  |
| Tested Application  | IHC-P, WB  |
| Specificity         | The antibody detects endogenous SAPK / JNK protein only when phosphorylated at Thr183.   |
| Host                | Rabbit   |
| Clonality           | Polyclonal   |
| Isotype             | IgG  |
| Target Name         | SAPK / JNK   |
| Species             | Human  |
| Immunogen           | Synthetic phospho-peptide around Thr183 of Human SAPK / JNK.   |
| Conjugation         | Un-conjugated  |
| Alternate Names     | MAP kinase 8; PRKM8; JNK1; c-Jun N-terminal kinase 1; Stress-activated protein kinase JNK1; MAPK 8;<br>SAPK1c; JNK21B1/2; JNK-46; Mitogen-activated protein kinase 8; EC 2.7.11.24; JNK1A2; JNK; Stress-<br>activated protein kinase 1c; SAPK1 |

## **Application Instructions**

| Application table | Application  | Dilution       |
|-------------------|--|----------------|
|                   | IHC-P  | 1:100 - 1:300  |
|                   | WB   | 1:500 - 1:2000 |
| Application Note  | IHC-P: Antigen Retrieval: Boil tissue section in Sodium citrate buffer (pH 6.0) for 20 min.<br>* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations<br>should be determined by the scientist. |                |
| Observed Size     | ~ 48 kDa   |                |

## Properties

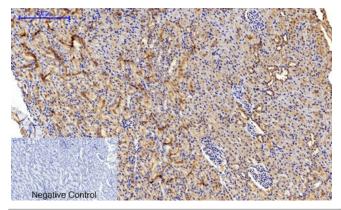
| Form                | Liquid   |
|---------------------|--|
| Purification        | Affinity purification with immunogen.  |
| Buffer              | PBS, 0.02% Sodium azide, 50% Glycerol and 0.5% BSA.  |
| Preservative        | 0.02% Sodium azide   |
| Stabilizer          | 50% Glycerol and 0.5% BSA  |
| Concentration       | 1 mg/ml  |
| Storage instruction | For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw |

Note

For laboratory research only, not for drug, diagnostic or other use.

## Bioinformation

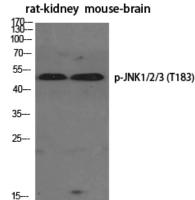
| Gene Symbol    | MAPK8  |
|----------------|--|
| Gene Full Name | mitogen-activated protein kinase 8   |
| Background     | The protein encoded by this gene is a member of the MAP kinase family. MAP kinases act as an integration point for multiple biochemical signals, and are involved in a wide variety of cellular processes such as proliferation, differentiation, transcription regulation and development. This kinase is activated by various cell stimuli, and targets specific transcription factors, and thus mediates immediate-early gene expression in response to cell stimuli. The activation of this kinase by tumor-necrosis factor alpha (TNF-alpha) is found to be required for TNF-alpha induced apoptosis. This kinase is also involved in UV radiation induced apoptosis, which is thought to be related to cytochrom c-mediated cell death pathway. Studies of the mouse counterpart of this gene suggested that this kinase play a key role in T cell proliferation, apoptosis and differentiation. Five alternatively spliced transcript variants encoding distinct isoforms have been reported. [provided by RefSeq, Jun 2013]  |
| Function       | Serine/threonine-protein kinase involved in various processes such as cell proliferation, differentiation, migration, transformation and programmed cell death. Extracellular stimuli such as proinflammatory cytokines or physical stress stimulate the stress-activated protein kinase/c-Jun N-terminal kinase (SAP/JNK) signaling pathway. In this cascade, two dual specificity kinases MAP2K4/MKK4 and MAP2K7/MKK7 phosphorylate and activate MAPK8/JNK1. In turn, MAPK8/JNK1 phosphorylates a number of transcription factors, primarily components of AP-1 such as JUN, JDP2 and ATF2 and thus regulates AP-1 transcriptional activity. Phosphorylates the replication licensing factor CDT1, inhibiting the interaction between CDT1 and the histone H4 acetylase HBO1 to replication origins. Loss of this interaction abrogates the acetylation required for polarized differentiation of T-helper cells into Th1 cells. Contributes to the survival of erythroid cells by phosphorylating the antagonist of cell death BAD upon EPO stimulation. Mediates starvation-induced BCL2 phosphorylating the antagonist of cell death BAD upon EPO stimulation. Mediates starvation-induced BCL2 phosphorylating the antagonist of radial migration from the ventricular zone. Phosphorylates several other substrates including heat shock factor protein 4 (HSF4), the deacetylase SIRT1, ELK1, or the E3 ligase ITCH. Phosphorylates the CLOCK-ARNTL/BMAL1 heterodimer and plays a role in the regulation of the circadian clock. |
| Calculated Mw  | 48 kDa   |
| РТМ            | Dually phosphorylated on Thr-183 and Tyr-185 by MAP2K7 and MAP2K4, which activates the enzyme.<br>Phosphorylated by TAOK2.   |



### ARG66184 anti-SAPK / JNK phospho (Thr183) antibody IHC-P image

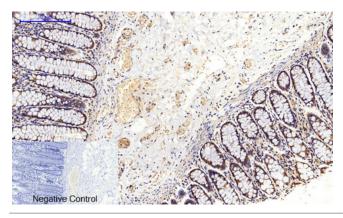
Immunohistochemistry: Paraffin-embedded Rat kidney tissue stained with ARG66184 anti-SAPK / JNK phospho (Thr183) antibody at 1:200 dilution (4°C, overnight). Antigen Retrieval: Boil tissue section in Sodium citrate buffer (pH 6.0) for 20 min.

Negative control was used by secondary antibody only.



### ARG66184 anti-SAPK / JNK phospho (Thr183) antibody WB image

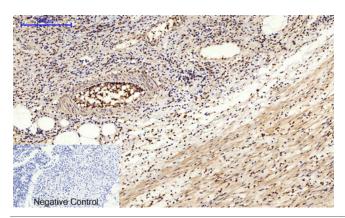
Western blot: Rat kidney and Mouse brain lysates stained with ARG66184 anti-SAPK / JNK phospho (Thr183) antibody at 1:1000 dilution.



### ARG66184 anti-SAPK / JNK phospho (Thr183) antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human colon tissue stained with ARG66184 anti-SAPK / JNK phospho (Thr183) antibody at 1:200 dilution (4°C, overnight). Antigen Retrieval: Boil tissue section in Sodium citrate buffer (pH 6.0) for 20 min.

Negative control was used by secondary antibody only.

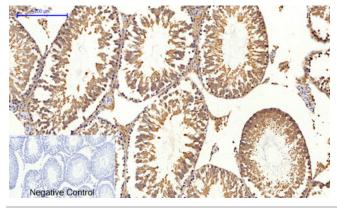


#### ARG66184 anti-SAPK / JNK phospho (Thr183) antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human Appendix tissue stained with ARG66184 anti-SAPK / JNK phospho (Thr183) antibody at 1:200 dilution (4°C, overnight). Antigen Retrieval: Boil tissue section in Sodium citrate buffer (pH 6.0) for 20 min.

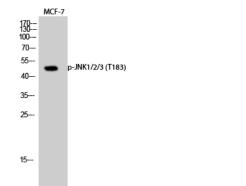
Negative control was used by secondary antibody only.

### ARG66184 anti-SAPK / JNK phospho (Thr183) antibody IHC-P image



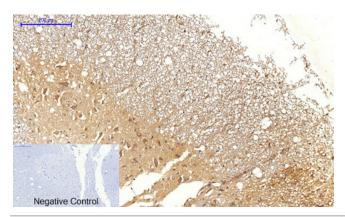
Immunohistochemistry: Paraffin-embedded Rat testis tissue stained with ARG66184 anti-SAPK / JNK phospho (Thr183) antibody at 1:200 dilution (4°C, overnight). Antigen Retrieval: Boil tissue section in Sodium citrate buffer (pH 6.0) for 20 min.

Negative control was used by secondary antibody only.



#### ARG66184 anti-SAPK / JNK phospho (Thr183) antibody WB image

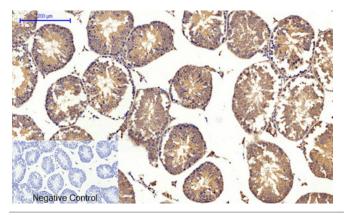
Western blot: MCF-7 cell lysate stained with ARG66184 anti-SAPK / JNK phospho (Thr183) antibody at 1:1000 dilution.



#### ARG66184 anti-SAPK / JNK phospho (Thr183) antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Rat spinal-cord tissue stained with ARG66184 anti-SAPK / JNK phospho (Thr183) antibody at 1:200 dilution (4°C, overnight). Antigen Retrieval: Boil tissue section in Sodium citrate buffer (pH 6.0) for 20 min.

Negative control was used by secondary antibody only.



#### ARG66184 anti-SAPK / JNK phospho (Thr183) antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Mouse testis tissue stained with ARG66184 anti-SAPK / JNK phospho (Thr183) antibody at 1:200 dilution (4°C, overnight). Antigen Retrieval: Boil tissue section in Sodium citrate buffer (pH 6.0) for 20 min.

Negative control was used by secondary antibody only.