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# Phospho-JNK1/2/3-T183/T183/T221 Rabbit mAb

Catalog No.: AP0631 Recombinant 76 Publications

## **Basic Information**

# **Observed MW**

46kDa/54kDa

#### **Calculated MW**

35kDa/44kDa/48kDa/27kDa/52kDa

#### Category

SMab Recombinant Monoclonal Antibody

#### **Applications**

WB,IHC-P,IF/ICC,ELISA

#### **Cross-Reactivity**

Human, Mouse, Rat

#### CloneNo number

ARC0193

# **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. Several alternatively spliced transcript variants encoding distinct isoforms have been reported. [provided by RefSeq, Apr 2016]

# **Recommended Dilutions**

**WB** 1:1000 - 1:5000

**IHC-P** 1:50 - 1:200

**IF/ICC** 1:50 - 1:200

**ELISA** Recommended starting

concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

#### **Contact**

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# **Immunogen Information**

Gene ID Swiss Prot

5599/5601/5602 P45983/P45984/P53779

**Immunogen** 

Synthetic peptide. This information is considered to be commercially sensitive.

**Synonyms** 

JNK1/JNK2/JNK3; Phospho-JNK1/2/3-T183/T183/T221

### **Product Information**

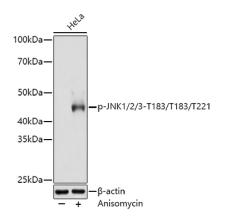
SourceIsotypePurificationRabbitIgGAffinity purification

#### Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS containing 50% glycerol and 0.05% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.

# **Validation Data**



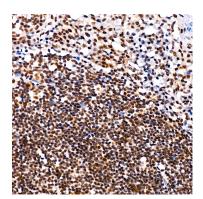
Western blot analysis of lysates from HeLa cells, using Phospho-JNK1/2/3-T183/T183/T221 Rabbit mAb (AP0631) at 1:3000 dilution. HeLa cells were treated with Anisomycin (25  $\mu$ g/mL) at 37°C for 30 minutes after serum-starvation overnight. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

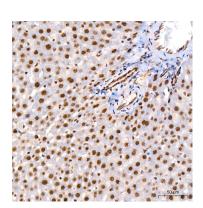
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

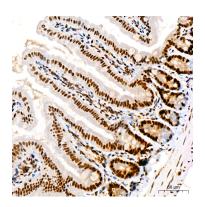
Exposure time: 180s.



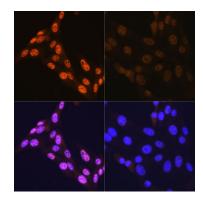
Immunohistochemistry analysis of paraffin-embedded Human tonsil using Phospho-JNK1/2/3-T183/T183/T221 Rabbit mAb (AP0631) at dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



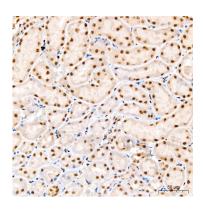
Immunohistochemistry analysis of paraffin-embedded Rat liver using Phospho-JNK1/2/3-T183/T183/T221 Rabbit mAb (AP0631) at dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffin-embedded Mouse colon using Phospho-JNK1/2/3-T183/T183/T221 Rabbit mAb (AP0631) at dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Immunofluorescence analysis of NIH-3T3 cells using Phospho-JNK1/2/3-T183/T183/T221 Rabbit mAb (AP0631).NIH-3T3 cells were treated with Anisomycin (25 μg/mL) at 37°C for 30 minutes after serumstarvation overnight. Secondary antibody: Cy3-conjugated Goat anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear



Immunohistochemistry analysis of paraffin-embedded Rat kidney using Phospho-JNK1/2/3-T183/T183/T221 Rabbit mAb (AP0631) at dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.