

Affinity Biosciences website:www.affbiotech.com order:order@affbiotech.com

MAPK1/3 Ab

Cat.#: AF0833 Concn.: 1mg/ml Mol.Wt.: 41kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000, IF/ICC 1:100-1:500

Reactivity: Human, Mouse, Rat

Purification: The antiserum was purified by peptide affinity

chromatography using SulfoLink™ Coupling Resin (Thermo

Fisher Scientific).

Specificity: MAPK1/3 Ab detects endogenous levels of MAPK1/3.

Immunogen: A synthesized peptide derived from human MAPK1/3.

Uniprot: P28482/P27361

Description: The protein encoded by this gene is a member of the MAP

kinase family. MAP kinases, also known as extracellular signal-regulated kinases (ERKs), 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. The activation of this kinase requires its phosphorylation by upstream kinases. Upon activation, this kinase translocates

to the nucleus of the stimulated cells, where it

phosphorylates nuclear targets. Two alternatively spliced transcript variants encoding the same protein, but differing in the UTRs, have been reported for this gene. [provided by

RefSeq]

Subcellular Location: Nucleus.

Similarity: The TXY motif contains the threonine and tyrosine residues

whose phosphorylation activates the MAP kinases.Belongs to the protein kinase superfamily. CMGC Ser/Thr protein kinase

family. MAP kinase subfamily.

Storage Condition and

Buffer:

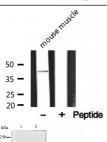
Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

°C.Stable for 12 months from date of receipt.

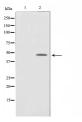


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Western blot analysis on mouse muscle lysate using MAPK1/3 Ab



Western blot analysis on JK cell lysate using MAPK1/3 Ab,The lane on the left is treated with the antigen-specific peptide.



AF0833 staining JK by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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