For Research Use Only

Phospho-p38 MAPK (Thr180/Tyr182) Recombinant antibody

Catalog Number:81212-2-RR



Basic Information

Catalog Number: GenBank Accession Number:

81212-2-RR BC031574 Protein A purfication

GeneID (NCBI): CloneNo.: 100ul , Concentration: 1000 $\mu g/ml$ by 1432 242308D3

Nanodrop: **UNIPROT ID:** Recommended Dilutions: Source: Q16539 WB 1:2000-1:10000

Rabbit Full Name:

Isotype: mitogen-activated protein kinase 14

IgG Calculated MW:

> 360 aa, 41 kDa Observed MW: 38-42 kDa

Applications

Tested Applications:

WB, ELISA Species Specificity:

human, mouse

Positive Controls:

WB: Anisomycin treated HeLa cells, UV treated NIH/3T3 cells, Anisomycin treated NIH/3T3 cells

Purification Method:

Background Information

A stress-activated serine/threonine protein kinase, p38 mitogen-activated protein kinase (p38 MAPK), belongs to the MAP kinase superfamily. Diverse extracellular stimuli, including ultraviolet light, irradiation, heat shock, high osmotic stress, proinflammatory cytokines and certain mitogens, trigger a stress-regulated protein kinase cascade culminating in activation of p38 MAPK through phosphorylation on a TGY motif within the kinase activation loop. The p38 MAPK undergoes dual phosphorylation at Thr182 and Tyr180 in the Thr-Gly-Tyr activation loop by MAP kinase kinase 6 (MKK6). Upon activation, p38 MAPK phosphorylates multiple substrates, including MAPK activated protein kinase 2 (MAPKAPK2) and activating transcription factor 2 (ATF-2). (PMID: 26901653, PMID: 10807318)

Storage

Storage:

Store at -20°C. Stable for one year after shipment.

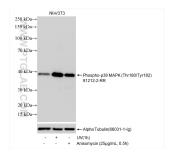
Storage Buffer:

PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

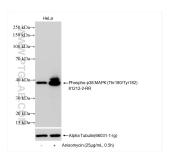
Aliquoting is unnecessary for -20°C storage

*** 20ul sizes contain 0.1% BSA

Selected Validation Data



Non-treated NIH/3T3 cells, UV treated and Anisomycin treated NIH/3T3 cells were subjected to SDS PAGE followed by western blot with 81212-2-RR (Phospho-p38 MAPK (Thr180/Tyr182) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Alpha Tubulin (66031-1-Ig) antibody as a loading control.



Non-treated HeLa cells and Anisomycin treated HeLa cells were subjected to SDS PAGE followed by western blot with 81212-2-RR (Phospho-p38 MAPK (Thr180/Tyr182) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Alpha Tubulin (66031-1-lg) antibody as a loading control.