

## Description

The ERK-MAPK Pathway Antibody Kit provides a cost-effective tool for studying key phospho-proteins involved in various steps of the ERK-MAPK pathway. Perfect for signal transduction researchers starting a new project, screening multiple prospective targets, or those who simply require less volume.

## Product Information

The ERK-MAPK Pathway Antibody Kit contains antibodies against 5 key phospho-protein targets that play critical roles in the ERK-MAPK pathway.

Antigen	Catalog No.	Host, clonality	Tested Reactivity	Applications	Volume
Phospho-MEK1 (Thr292)	<a href="#">67873-1-Ig</a>	Mouse monoclonal	H, M, R	WB, FC	20 uL
Phospho-ERK1/2 (Thr202/Tyr204)	<a href="#">80031-1-RR</a>	Rabbit Monoclonal	H	WB	20 uL
Phospho-RPS6KA1 (Ser380)	<a href="#">80108-1-RR</a>	Rabbit Monoclonal	H, M	WB, IHC, FC	20 uL
Phospho-Jun (Ser73)	<a href="#">80086-1-RR</a>	Rabbit Monoclonal	H, M	WB	20 uL
Phospho-MNK1 (Thr250/255)	<a href="#">81398-1-RR</a>	Rabbit Monoclonal	H	WB	20 uL

## Package

5×20 uL

## Storage

Store at -20°C. Stable for one year from the date of receipt.

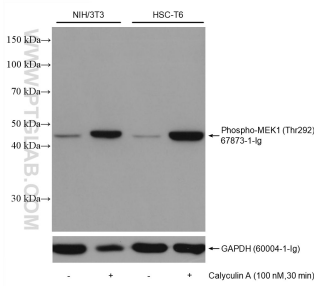
## Background Information

The ERK-MAPK signaling pathway regulates several important cellular processes including proliferation, differentiation, and survival in response to extracellular signals such as growth factor/hormone stimulation and environmental stress. The pathway is structured into several tiers in which protein kinases sequentially phosphorylate each other, resulting in the activation of gene expression downstream. Phospho-MEK1 is a MAP2K that phosphorylates and activates ERK1/2 at several sites including Thr202/Tyr204. Once activated, the phosphorylated-ERK translocates to the nucleus and then phosphorylates several transcription factors downstream including c-Jun, c-Myc, STAT, and HIF1. Ser/Thr kinases such as p90RSK/RPS6KA1 and MNK1 can also be phosphorylated by ERK, resulting in the activation of additional transcription factors and translation factors respectively.

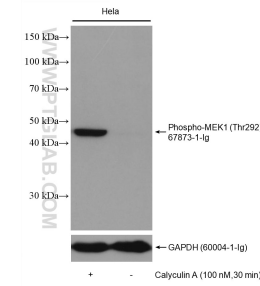
## Standard Protocols

Click [here](#) to view our standard protocols for various applications including WB, IP, IHC, IF, FC, and ELISA.

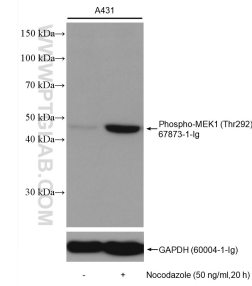
## Validation Data



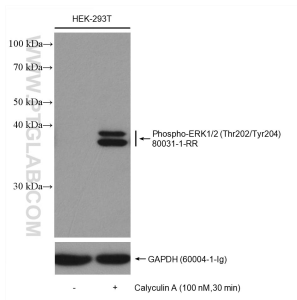
Non-treated cells and Calyculin A treated cells were subjected to SDS PAGE followed by western blot with 67873-1-Ig (Phospho-MEK1 (Thr292) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.



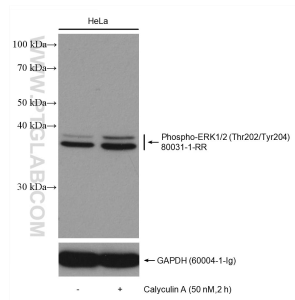
Non-treated HeLa and Calyculin A treated HeLa cells were subjected to SDS PAGE followed by western blot with 67873-1-Ig (Phospho-MEK1 (Thr292) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.



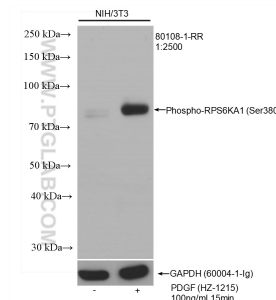
Non-treated A431 and Nocodazole treated A431 cells were subjected to SDS PAGE followed by western blot with 67873-1-Ig (Phospho-MEK1 (Thr292) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.



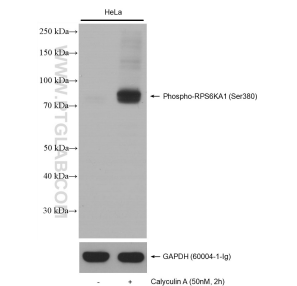
Non-treated HEK-293T and Calyculin A treated HEK-293T cells were subjected to SDS PAGE followed by western blot with 80031-1-RR (Phospho-ERK1/2 (Thr202/Tyr204) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours.



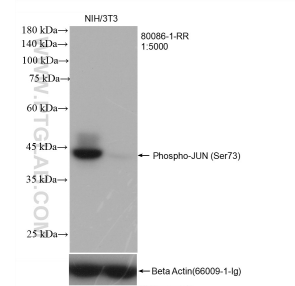
Non-treated HeLa and Calyculin A treated HeLa cells were subjected to SDS PAGE followed by western blot with 80031-1-RR (Phospho-ERK1/2 (Thr202/Tyr204) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours.



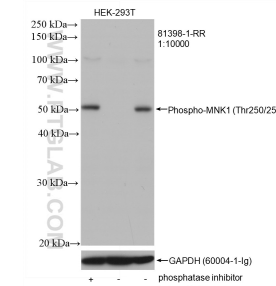
Non-treated NIH/3T3 and PDGF (HZ-1215) treated NIH/3T3 cells were subjected to SDS PAGE followed by western blot with 80108-1-RR (Phospho-RPS6KA1 (Ser380) antibody) at dilution of 1:2500 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.



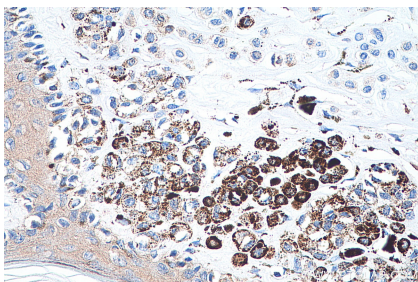
Non-treated HeLa cells and Calyculin A treated HeLa cells were subjected to SDS PAGE followed by western blot with 80108-1-RR (Phospho-RPS6KA1 (Ser380) antibody) at dilution of 1:2500 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.



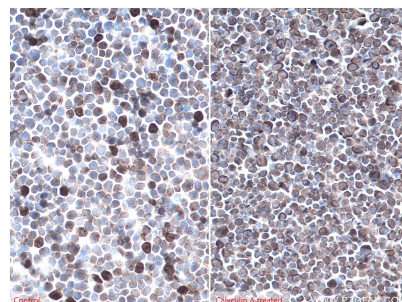
UV treated and non-treated NIH/3T3 cells were subjected to SDS PAGE followed by western blot with 80086-1-RR (Phospho-JUN (Ser73) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Actin antibody (66009-1-Ig) as loading control.



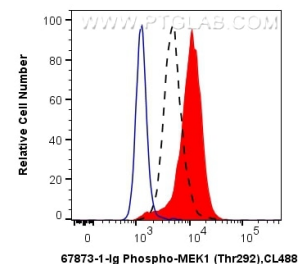
Non-treated HEK-293T cells, phosphatase inhibitor and lambda phosphatase treated HEK-293T cells were subjected to SDS PAGE followed by western blot with 81398-1-RR (Phospho-MNK1 (Thr250/255) antibody) at dilution of 1:10000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as... loading control.



Immunohistochemical analysis of paraffin-embedded human malignant melanoma tissue slide using 80108-1-RR (Phospho-RPS6KA1 (Ser380) antibody) at dilution of 1:2000 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Immunohistochemical analysis of paraffin-embedded Jurkat cells slide using 80108-1-RR (Phospho-RPS6KA1 (Ser380) antibody) at dilution of 1:2000 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



1X10<sup>6</sup> HeLa cells untreated (dashed lines) or Calyculin A (red) treated were intracellularly stained with 0.13 ug Anti-Human Phospho-MEK1 (Thr292) (67873-1-Ig, Clone:2D7A8) and CoraLite<sup>®</sup> 488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L) at dilution 1:1000, or 0.13 ug

For technical support and original validation data for this product please contact

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E: proteintech@ptglab.com  
W: ptglab.com

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