For Research Use Only

## Phospho-MEK6 (Ser207) Polyclonal antibody

Catalog Number: 28903-1-AP



**Basic Information** 

Catalog Number: GenBank Accession Number:

28903-1-AP BC012009 GeneID (NCBI):

100ul, Concentration: 600 ug/ml by 5608 Nanodrop: **UNIPROT ID:** Source: P52564 Rabbit Full Name:

Isotype: mitogen-activated protein kinase

kinase 6 IgG

> 334 aa, 37 kDa Observed MW: 35-37 kDa

Calculated MW:

**Applications** 

**Tested Applications:** 

WB, ELISA

Species Specificity:

Human

Recommended Dilutions: WB 1:1000-1:6000

Antigen affinity purification

**Purification Method:** 

Positive Controls:

WB: UV treated HEK-293 cells, HEK-293 cells, HeLa

cells, Anisomycin treated HeLa cells

## **Background Information**

MAPKK6 (Dual specificity mitogen activated protein kinase kinase 6), also known as MAP2K6 / MEK6 / MKK6, is a member of MAPKK protein kinase family. MKK6 plays an important role in intracellular signaling pathways leading toward activation of the p38 MAP kinase. MEK6 phosphorylates and activates p38 MAP kinase in response to inflammatory cytokines or environmental stress. Phosphorylation of Ser207 and Thr211 by MAP3Ks is critical for MEK6 activation, and acetylation of these resides by VopA blocks MEK6 activity(PMID: 26648936). The antibody also recognizes the ser218 phosphorylation site of MEK3.

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

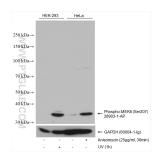
\*\*\* 20ul sizes contain 0.1% BSA

in USA), or 1(312) 455-8498 (outside USA)

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This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

## Selected Validation Data



Non-treated HEK-293 cells, UV treated HEK-293 cells, non-treated HeLa cells and Anisomycin treated HeLa cells were subjected to SDS PAGE followed by western blot with 28903-1-AP (Phospho-MEK6 (Ser207) antibody) at dilution of 1:3000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.