

# Phospho-MEK1/MEK2-S217/S221 Rabbit mAb

Catalog No.: AP1349

Recombinant

2 Publications

## Basic Information

**Observed MW**

45kDa

**Calculated MW**

40kDa/43kDa/44kDa

**Category**

Primary antibody

**Applications**

WB,IP,ELISA

**Cross-Reactivity**

Human, Mouse, Rat

**CloneNo number**

ARC55711

## Background

The protein encoded by this gene is a member of the dual specificity protein kinase family, which acts as a mitogen-activated protein (MAP) kinase kinase. MAP kinases, also known as extracellular signal-regulated kinases (ERKs), act as an integration point for multiple biochemical signals. This protein kinase lies upstream of MAP kinases and stimulates the enzymatic activity of MAP kinases upon wide variety of extra- and intracellular signals. As an essential component of MAP kinase signal transduction pathway, this kinase is involved in many cellular processes such as proliferation, differentiation, transcription regulation and development.

## Recommended Dilutions

**WB** 1:2000 - 1:10000**IP** 1µg-4µg antibody for  
800µg-1000µg extracts  
of whole cells**ELISA** Recommended starting  
concentration is 1 µg/mL.  
Please optimize the  
concentration based on  
your specific assay  
requirements.

## Immunogen Information

**Gene ID**

5604/5605

**Swiss Prot**

Q02750/P36507

**Immunogen**

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

**Synonyms**

MEK1/MEK2; Phospho-MEK1/MEK2-S217/S221

## Contact

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## Product Information

**Source**

Rabbit

**Isotype**

IgG

**Purification**

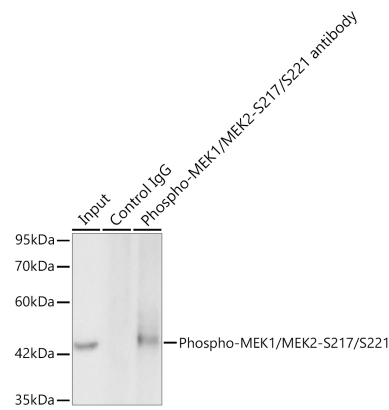
Affinity purification

**Storage**

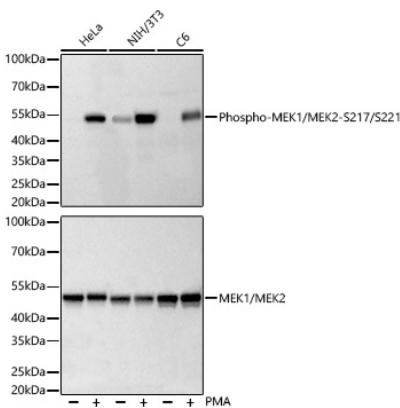
Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% Sodium azide, 0.05% BSA, 50% glycerol, pH7.3.

Validation Data



Immunoprecipitation of Phospho-MEK1/MEK2-S217/S221 from 1000 µg extracts of HeLa cells treated with PMA (200nM, 1h) was performed using 2 µg of Phospho-MEK1/MEK2-S217/S221 Rabbit mAb (AP1349). Rabbit IgG isotype control (AC042) was used to precipitate the Control IgG sample. IP samples were eluted with 1X Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using Phospho-MEK1/MEK2-S217/S221 Rabbit mAb (AP1349) at a dilution of 1:5000.



Western blot analysis of various lysates, using Phospho-MEK1/MEK2-S217/S221 Rabbit mAb (AP1349) at 1:3000 dilution (upper) or MEK1/MEK2 Rabbit mAb (A4868) at 1:1000 dilution (lower). incubated overnight at 4°C. HeLa cells and NIH/3T3 cells were treated with PMA (100 nM) at 37°C for 30 minutes after serum-starvation overnight. C6 cells were treated with PMA (200 nM) 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: 30 µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 45s.