# Phospho-MEK1-S298 Rabbit pAb

Catalog No.: AP0063 3 Publications



# **Basic Information**

#### **Observed MW**

43kDa

#### **Calculated MW**

43kDa

## Category

Primary antibody

## **Applications**

WB,IP,ELISA

## **Cross-Reactivity**

Human, Mouse, Rat

# **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** 1:500 - 1:2000

**IP** 0.5μg-4μg antibody for 200μg-400μg 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**Swiss Prot
5604
Q02750

## **Immunogen**

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

## **Synonyms**

MEL; CFC3; MEK1; MKK1; MAPKK1; PRKMK1; Phospho-MEK1-S298

# **Contact**

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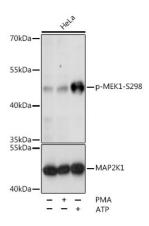
# **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

### Storage

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

Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.

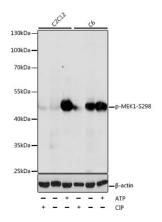


Western blot analysis of various lysates using Phospho-MEK1-S298 Rabbit pAb (AP0063) at 1:1000 dilution or MEK1 antibody (A12687). HeLa cells were treated with PMA/TPA (200 nM) at 37°C for 15 minutes after serum-starvation overnight or treated with ATP(5 mM) at 30°C for 1 hour.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane. Blocking buffer: 3% BSA. Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.



Western blot analysis of various lysates using Phospho-MEK1-S298 Rabbit pAb (AP0063) at 1:1000 dilution. C2C12 cells were treated with CIP(20uL/400ul) at  $37^{\circ}$ C for 1 hour or treated with ATP(5 mM) at  $30^{\circ}$ C for 1 hour. C6 cells were treated with CIP(20uL/400ul) at  $37^{\circ}$ C for 1 hour or treated with ATP(5 mM) at  $30^{\circ}$ C for 1 hour.

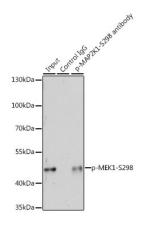
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins:  $25\mu g$  per lane.

Blocking buffer: 3% BSA.

Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.



Immunoprecipitation analysis of 200  $\mu g$  extracts of 293T cells, using 3  $\mu g$  Phospho-MEK1-S298 pAb (AP0063). Western blot was performed from the immunoprecipitate using Phospho-MEK1-S298 pAb (AP0063) at a dilution of 1:1000. 293T cells were treated with PMA/TPA (200 nM) at 37°C for 30 minutes after serum-starvation overnight.