

MEK-2 (phospho Thr394) Polyclonal Antibody

Catalog No: YP0169

Reactivity: Human; Mouse; Rat

Applications: WB;IHC;IF;IP;ELISA

Target: MEK2

Fields: >>EGFR tyrosine kinase inhibitor resistance;>>Endocrine resistance;>>MAPK

signaling pathway;>>ErbB signaling pathway;>>Ras signaling pathway;>>Rap1

signaling pathway;>>cGMP-PKG signaling pathway;>>cAMP signaling

pathway;>>HIF-1 signaling pathway;>>FoxO signaling pathway;>>Sphingolipid

signaling pathway;>>Phospholipase D signaling pathway;>>Autophagy -

animal;>>mTOR signaling pathway;>>PI3K-Akt signaling

pathway;>>Apoptosis;>>Cellular senescence;>>Vascular smooth muscle contraction;>>VEGF signaling pathway;>>Apelin signaling pathway;>>Gap junction;>>Signaling pathways regulating pluripotency of stem cells;>>Neutrophil extracellular trap formation;>>Toll-like receptor signaling pathway;>>Natural killer cell mediated cytotoxicity;>>T cell receptor signaling pathway;>>B cell receptor

signaling pathway;>>Fc epsilon RI signaling pathway;>>Long-term potentiation;>>Neurotrophin signaling pathway;>>Long-term depression;>>Regulation of actin cytoskeleton;>>Insulin signaling

pathway;>>GnRH signal

Gene Name: MAP2K2

Protein Name: Dual specificity mitogen-activated protein kinase kinase 2

Human Gene Id: 5606

Human Swiss Prot P36507

No:

Mouse Gene Id: 26396

Q63932

Mouse Swiss Prot

No:

Rat Gene ld: 58960

Rat Swiss Prot No: P36506

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Immunogen: The antiserum was produced against synthesized peptide derived from human

MEK2 around the phosphorylation site of Thr394. AA range:261-310

Specificity: Phospho-MEK-2 (T394) Polyclonal Antibody detects endogenous levels of

MEK-2 protein only when phosphorylated at T394.

Formulation : Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

Dilution: WB 1:500 - 1:2000. IHC 1:100 - 1:300. Immunoprecipitation: 2-5 ug:mg lysate.

ELISA: 1:10000.. IF 1:50-200

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Molecularweight: 44kD

Cell Pathway: Regulates Angiogenesis; Regulation of Actin Dynamics; Stem cell pathway;

T_Cell_Receptor; Insulin Receptor; Cell Growth; Toll_Like;

MAPK_ERK_Growth;MAPK_G_Protein; B_Cell_Antigen; PI3K/Akt

Background: The protein encoded by this gene is a dual specificity protein kinase that belongs

to the MAP kinase kinase family. This kinase is known to play a critical role in mitogen growth factor signal transduction. It phosphorylates and thus activates MAPK1/ERK2 and MAPK2/ERK3. The activation of this kinase itself is dependent on the Ser/Thr phosphorylation by MAP kinase kinase kinases. Mutations in this gene cause cardiofaciocutaneous syndrome (CFC syndrome), a disease characterized by heart defects, mental retardation, and distinctive facial features

similar to those found in Noonan syndrome. The inhibition or degradation of this kinase is also found to be involved in the pathogenesis of Yersinia and anthrax. A pseudogene, which is located on chromosome 7, has been identified for this

gene. [provided by RefSeq, Jul 2008],

Function: catalytic activity:ATP + a protein = ADP + a phosphoprotein.,disease:Defects in

MAP2K2 are a cause of cardiofaciocutaneous syndrome (CFC syndrome) [MIM:115150]; also known as cardio-facio-cutaneous syndrome. CFC syndrome is characterized by a distinctive facial appearance, heart defects and mental retardation. Heart defects include pulmonic stenosis, atrial septal defects and hypertrophic cardiomyopathy. Some affected individuals present with ectodermal abnormalities such as sparse, friable hair, hyperkeratotic skin lesions and a

generalized ichthyosis-like condition. Typical facial features are similar to Noonan

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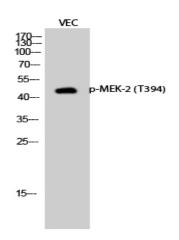
syndrome. They include high forehead with bitemporal constriction, hypoplastic supraorbital ridges, downslanting palpebral fissures, a depressed nasal bridge, and posteriorly angulated ears with prominent helices. The inheritance of CFC syndrome is autosomal dominant.,function:C

Subcellular Location:

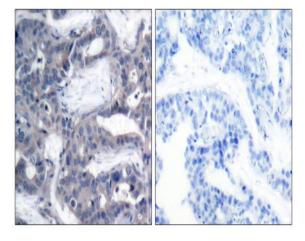
Cytoplasm . Membrane ; Peripheral membrane protein . Membrane localization is probably regulated by its interaction with KSR1. .

Expression: Colon carcinoma, Epithelium, Human cerebellum, Muscle, Platelet

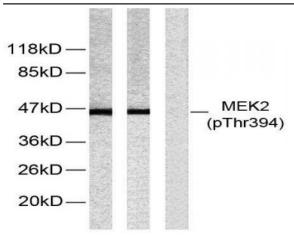
Products Images



Western Blot analysis of VEC cells using Phospho-MEK-2 (T394) Polyclonal Antibody diluted at 1:2000



Immunohistochemistry analysis of paraffin-embedded human breast carcinoma, using MEK2 (Phospho-Thr394) Antibody. The picture on the right is blocked with the phospho peptide.



Western blot analysis of lysates from ovary cancer, using MEK2 (Phospho-Thr394) Antibody. The lane on the right is blocked with the phospho peptide.