

Raf-1 (phospho Tyr341) Polyclonal Antibody

Catalog No: YP0670

Reactivity: Human; Mouse; Rat

Applications: WB;IHC;IF;ELISA

Target: Raf-1

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;>>Chemokine 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;>>Axon guidance;>>VEGF signaling pathway;>>Apelin signaling pathway;>>Focal adhesion;>>Gap junction;>>Signaling pathways

regulating pluripotency of stem cells;>>Neutrophil extracellular trap formation;>>C-type lectin receptor signaling pathway;>>JAK-STAT signaling pathway;>>Natural killer cell mediated cytotoxicity;>>T cell receptor signaling pathway;>>B cell receptor signaling pathway;>>Fc epsilon RI signaling pathway;>>Fc gamma R-mediated phagocytosis:>>Long-term potentiation;>>Neurotrophin signaling pathway;>>Fc epsilon RI signaling pathway;>>Fc epsilon RI signaling pathway;>>Fc gamma R-mediated phagocytosis:>>Long-term potentiation;>>Neurotrophin signaling pathway;>>Fc epsilon RI signaling pathway;>>Fc epsilon RI signaling pathway;>>Fc gamma R-mediated phagocytosis:>>Long-term potentiation;>>Neurotrophin signaling pathway;>>Fc epsilon RI signaling pathway;

Gene Name: RAF1

Protein Name: RAF proto-oncogene serine/threonine-protein kinase

Human Gene Id: 5894

Human Swiss Prot P04049

No:

Mouse Gene Id: 110157

Mouse Swiss Prot

No:

Q99N57

Rat Gene ld: 24703

Rat Swiss Prot No: P11345

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

C-RAF around the phosphorylation site of Tyr341. AA range:311-360

Specificity: Phospho-Raf-1 (Y341) Polyclonal Antibody detects endogenous levels of Raf-1

protein only when phosphorylated at Y341.

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. 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)

Observed Band: 73kD

Cell Pathway: MAPK_ERK_Growth;MAPK_G_Protein;ErbB_HER;Chemokine;Vascular

smooth muscle contraction; VEGF; Focal adhesion; Gap junction; Natural killer cell

mediated cytotoxicity; T_Cell_Receptor; B_Cell_Antigen; Fc epsilo

Background: This gene is the cellular homolog of viral raf gene (v-raf). The encoded protein is

a MAP kinase kinase kinase (MAP3K), which functions downstream of the Ras family of membrane associated GTPases to which it binds directly. Once activated, the cellular RAF1 protein can phosphorylate to activate the dual specificity protein kinases MEK1 and MEK2, which in turn phosphorylate to activate the serine/threonine specific protein kinases, ERK1 and ERK2. Activated ERKs are pleiotropic effectors of cell physiology and play an important role in the control of gene expression involved in the cell division cycle, apoptosis, cell differentiation and cell migration. Mutations in this gene are associated with

Noonan syndrome 5 and LEOPARD syndrome 2. [provided by RefSeq, Jul 2008],

Function : catalytic activity:ATP + a protein = ADP + a phosphoprotein.,cofactor:Binds 2

zinc ions per subunit., disease: Defects in RAF1 are the cause of LEOPARD syndrome type 2 (LEOPARD syndrome-2) [MIM:611554]. LEOPARD syndrome is an autosomal dominant disorder allelic with Noonan syndrome. The acronym LEOPARD stands for lentigines, electrocardiographic conduction abnormalities, ocular hypertelorism, pulmonic stenosis, abnormalities of genitalia, retardation of growth, and deafness., disease: Defects in RAF1 are the cause of Noonan syndrome type 5 (NS5) [MIM:611553]. Noonan syndrome (NS) is a disorder

characterized by dysmorphic facial features, short stature, hypertelorism, cardiac anomalies, deafness, motor delay, and a bleeding diathesis. It is a genetically

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heterogeneous and relatively common syndrome, with an estimated incidence of 1 in 1000-2500 live births.,function:Involved in the transducti

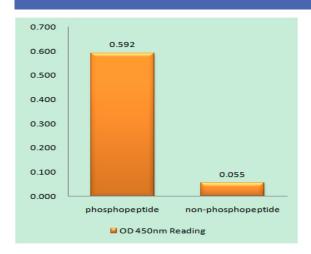
Subcellular Location:

Cytoplasm. Cell membrane. Mitochondrion. Nucleus. Colocalizes with RGS14 and BRAF in both the cytoplasm and membranes. Phosphorylation at Ser-259 impairs its membrane accumulation. Recruited to the cell membrane by the active Ras protein. Phosphorylation at Ser-338 and Ser-339 by PAK1 is required for its mitochondrial localization. Retinoic acid-induced Ser-621 phosphorylated form of RAF1 is predominantly localized at the nucleus.

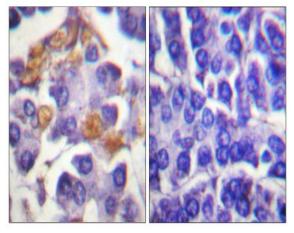
Expression:

In skeletal muscle, isoform 1 is more abundant than isoform 2.

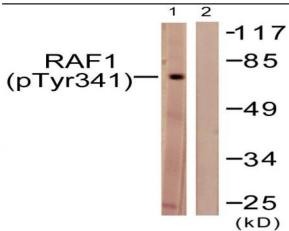
Products Images



Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-Phosphopeptide (Phospho-right), using C-RAF (Phospho-Tyr341) Antibody



Immunohistochemistry analysis of paraffin-embedded human pancreas, using C-RAF (Phospho-Tyr341) Antibody. The picture on the right is blocked with the phospho peptide.



Western blot analysis of lysates from Jurkat cells treated with Paclitaxel 1uM 24h, using C-RAF (Phospho-Tyr341) Antibody. The lane on the right is blocked with the phospho peptide.