abcam

Product datasheet

Recombinant Rafl (mutated Y340E + Y341E) protein (Active) ab62292

2 Images

Description

Product name Recombinant Raf1 (mutated Y340E + Y341E) protein (Active)

Biological activity The specific activity of RAF1 was determined to be ~6,000 nmol/min/mg in a coupled assay as

per antibody assay protocol.

Purity > 80 % SDS-PAGE.

Purity >80% as determined by SDS-PAGE and Coomassie blue staining. Affinity purified.

Expression system Baculovirus infected Sf9 cells

Accession P04049-1

Protein length Protein fragment

Animal free No

Nature Recombinant

Predicted molecular weight 63 kDa including tags

Amino acids 306 to 648

Modifications mutated Y340E + Y341E

Tags GST tag N-Terminus

Specifications

Our Abpromise guarantee covers the use of ab62292 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications Western blot

Functional Studies

Form Liquid

Preparation and Storage

Stability and Storage Shipped at 4°C. Upon delivery aliquot. Store at -80°C. Avoid freeze / thaw cycle.

pH: 7.50

Constituents: 0.00385% DTT, 0.79% Tris HCl, 25% Glycerol (glycerin, glycerine), 0.87% Sodium

chloride, 0.31% Glutathione, 0.0029% EDTA, 0.0017% PMSF

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General Info

Function

Involved in the transduction of mitogenic signals from the cell membrane to the nucleus. Part of the Ras-dependent signaling pathway from receptors to the nucleus. Protects cells from apoptosis mediated by STK3.

Tissue specificity

Involvement in disease

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

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 heterogeneous and relatively common syndrome, with an estimated incidence of 1 in 1000-2500 live births.

Defects in RAF1 are the cause of LEOPARD syndrome type 2 (LEOPARD2) [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.

Sequence similarities

Belongs to the protein kinase superfamily. TKL Ser/Thr protein kinase family. RAF subfamily.

Contains 1 phorbol-ester/DAG-type zinc finger.

Contains 1 protein kinase domain.

Contains 1 RBD (Ras-binding) domain.

Post-translational modifications

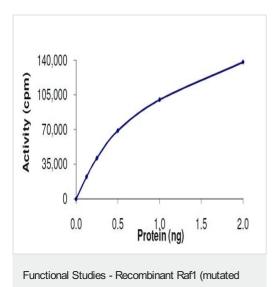
Phosphorylated upon DNA damage, probably by ATM or ATR. Phosphorylation at Thr-269 increases its kinase activity. Phosphorylation at Ser-259 induces the interaction with YWHAZ and inactivates kinase activity. Dephosphorylation of Ser-259 by the complex containing protein phosphatase 1, SHOC2 and M-Ras/MRAS relieves inactivation, leading to stimulate RAF1 activity.

Cellular localization

 $\label{eq:cytoplasm.coll} \textbf{Cytoplasm}. \textbf{ Cell membrane}. \textbf{ Colocalizes with RGS14} \textbf{ and BRAF in both the cytoplasm and}$

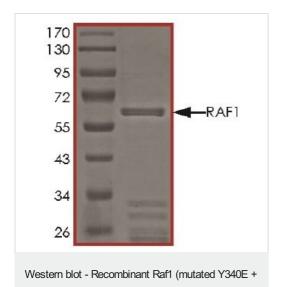
membranes.

Images



Y340E + Y341E) protein (Active) (ab62292)

ab62292 Kinase assay plot



Y341E) protein (Active) (ab62292)

Recombinant Raf1 (mutated Y340E + Y341E) protein (Active) (ab62292) + RAF1

Predicted band size: 63 kDa **Observed band size:** 63 kDa

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