abcam

Product datasheet

Recombinant Human RhoA protein ab91068

Description

Product name Recombinant Human RhoA protein

Purity > 90 % SDS-PAGE.

Expression system Escherichia coli

Accession P61586

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Additional sequence information AF498970

Specifications

Our Abpromise guarantee covers the use of ab91068 in the following tested applications.

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

Applications SDS-PAGE

Form Liquid

Preparation and Storage

Stability and Storage Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

pH: 7.20

Constituents: 0.00088% GDP, 0.019% Magnesium chloride, 0.077% DTT, 0.595% HEPES,

0.232% Sodium chloride

General Info

Function Regulates a signal transduction pathway linking plasma membrane receptors to the assembly of

focal adhesions and actin stress fibers. Serves as a target for the yopT cysteine peptidase from

Yersinia pestis, vector of the plague, and Yersinia pseudotuberculosis, which causes gastrointestinal disorders. May be an activator of PLCE1. Activated by ARHGEF2, which

promotes the exchange of GDP for GTP.

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Sequence similarities Belongs to the small GTPase superfamily. Rho family.

Domain The basic-rich region is essential for yopT recognition and cleavage.

Substrate for botulinum ADP-ribosyltransferase.

Post-translational

modifications Cleaved by vopT protease when the cell is infected by some Yers

Cleaved by yopT protease when the cell is infected by some Yersinia pathogens. This removes the lipid attachment, and leads to its displacement from plasma membrane and to subsequent

cytoskeleton cleavage.

AMPylation at Tyr-34 and Thr-37 are mediated by bacterial enzymes in case of infection by H.somnus and V.parahaemolyticus, respectively. AMPylation occurs in the effector region and leads to inactivation of the GTPase activity by preventing the interaction with downstream effectors, thereby inhibiting actin assembly in infected cells. It is unclear whether some human enzyme mediates AMPylation; FICD has such ability in vitro but additional experiments remain to

be done to confirm results in vivo.

Ubiquitinated by the BCR(BACURD1) and BCR(BACURD2) E3 ubiquitin ligase complexes, leading to its degradation by the proteasome, thereby regulating the actin cytoskeleton and cell

migration.

Cellular localization Cell membrane. Cytoplasm > cytoskeleton.

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