Rho GDIα (G-2): sc-373724



The Power to Question

BACKGROUND

Members of the Ras superfamily of small GTP-binding proteins are critical mediators of diverse cell signaling pathways, including those leading to cell proliferation, cytoskeletal organization and secretion. The counter-conversion of the active GTP-bound form of these proteins to their inactive GDP-bound form is influenced by two types of regulatory proteins: those that alter the intrinsic GTPase activity of the GTP-binding proteins and those that alter the rate of GDP/GTP exchange. Guanine nucleotide-releasing factors (GRFs) increase the GDP dissociation rate, while GDP-dissociation inhibitors (GDIs) decrease the dissociation rate. Rho GDI α , also known as ARHGDIA or GDIA1, is a 204 amino acid member of the Rho GDI family of proteins. Localized to the cytoplasm, Rho GDI α inhibits the dissociation of GDP from Rho proteins, thereby preventing GTP from binding to and subsequently activating Rho proteins. In humans, Rho GDI α can be phosphorylated at Ser 101 by p21-activated kinase (α PAK), an event that inhibits Rho GDI α activity and may result in positive feedback regulation of certain Rho GDI α target proteins.

CHROMOSOMAL LOCATION

Genetic locus: ARHGDIA (human) mapping to 17q25.3; Arhgdia (mouse) mapping to 11 E2.

SOURCE

Rho $GDl\alpha$ (G-2) is a mouse monoclonal antibody specific for an epitope mapping between amino acids 2-29 at the N-terminus of Rho $GDl\alpha$ of human origin.

PRODUCT

Each vial contains 200 $\mu g \ lgG_{2h}$ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Rho GDI α (G-2) (G-2) is available conjugated to agarose (sc-373724 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-373724 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-373724 PE), fluorescein (sc-373724 FITC), Alexa Fluor® 488 (sc-373724 AF488), Alexa Fluor® 546 (sc-373724 AF546), Alexa Fluor® 594 (sc-373724 AF594) or Alexa Fluor® 647 (sc-373724 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor® 680 (sc-373724 AF680) or Alexa Fluor® 790 (sc-373724 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

Blocking peptide available for competition studies, sc-373724 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% stabilizer protein)

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STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

PROTOCOLS

See our web site at www.scbt.com for detailed protocols and support products.

APPLICATIONS

Rho GDI α (G-2) is recommended for detection of Rho GDI α of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

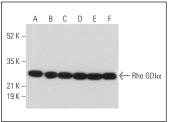
Rho GDI α (G-2) is also recommended for detection of Rho GDI α in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for Rho GDI α siRNA (h): sc-36417, Rho GDI α siRNA (m): sc-36416, Rho GDI α siRNA (r): sc-61880, Rho GDI α shRNA Plasmid (h): sc-36417-SH, Rho GDI α shRNA Plasmid (m): sc-36416-SH, Rho GDI α shRNA Plasmid (r): sc-61880-SH, Rho GDI α shRNA (h) Lentiviral Particles: sc-36417-V, Rho GDI α shRNA (m) Lentiviral Particles: sc-36416-V and Rho GDI α shRNA (r) Lentiviral Particles: sc-61880-V.

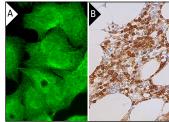
Molecular Weight of Rho GDI α : 30 kDa.

Positive Controls: MCF7 whole cell lysate: sc-2206, K-562 whole cell lysate: sc-2203 or Jurkat whole cell lysate: sc-2204.

DATA







Rho GDI α (G-2): sc-373724. Immunofluorescence staining of formalin-fixed Hep G2 cells showing cytoskeleton localization (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human bone marrow tissue showing cytoolasmic staining of hematopoietic cells (B)

SELECT PRODUCT CITATIONS

- Hernandez, G., et al. 2013. MitoTimer: a novel tool for monitoring mitochondrial turnover. Autophagy 9: 1852-1861.
- Sorrentino, S., et al. 2018. Hindlimb ischemia impairs endothelial recovery and increases neointimal proliferation in the carotid artery. Sci. Rep. 8: 761.
- 3. Mohammad, G., et al. 2019. Functional regulation of an oxidative stress mediator, Rac 1, in diabetic retinopathy. Mol. Neurobiol. 56: 8643-8655.
- 4. Kholmanskikh, S., et al. 2022. Activation of RhoC by regulatory ubiquitination is mediated by LNX1 and suppressed by LIS1. Sci. Rep. 12: 16493.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

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