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RHOA Recombinant Monoclonal Antibody

Product Code	CSB-RA898030A0HU	
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.	
Uniprot No.	P61586	
Immunogen	A synthesized peptide derived from human Rho	
Species Reactivity	Human	
Tested Applications	ELISA, WB; Recommended dilution: WB:1:500-1:5000	
Relevance	Regulates a signal transduction pathway linking plasma membrane receptors to the assembly of focal adhesions and actin stress fibers. Involved in a microtubule-dependent signal that is required for the myosin contractile ring formation during cell cycle cytokinesis. Plays an essential role in cleavage furrow formation. Required for the apical junction formation of keratinocyte cell-cell adhesion. Stimulates PKN2 kinase activity. May be an activator of PLCE1. Activated by ARHGEF2, which promotes the exchange of GDP for GTP. Essential for the SPATA13-mediated regulation of cell migration and adhesion assembly and disassembly. The MEMO1-RHOA-DIAPH1 signaling pathway plays an important role in ERBB2-dependent stabilization of microtubules at the cell cortex. It controls the localization of APC and CLASP2 to the cell membrane, via the regulation of GSK3B activity. In turn, membrane-bound APC allows the localization of the MACF1 to the cell membrane, which is required for microtubule capture and stabilization. Regulates a signal transduction pathway linking plasma membrane receptors to the assembly of focal adhesions and actin stress fibers. Involved in a microtubule-dependent signal that is required for the myosin contractile ring formation. Required for the apical junction formation of keratinocyte cell-cell adhesion. May be an activator of PLCE1. Activated by ARHGEF2, which promotes the exchange of GDP for GTP. Essential for the SPATA13-mediated regulation of cell migration and adhesion assembly and disassembly. The MEMO1-RHOA-DIAPH1 signaling pathway plays an important role in ERBB2-dependent stabilization of microtubules at the cell cortex. It controls the localization of APC and CLASP2 to the cell membrane, via the regulation of GSK3B activity. In turn, membrane-bound APC allows the localization of the MACF1 to the cell membrane, which is required for microtubules at the cell cortex. It controls the localization of APC and CLASP2 to the cell membrane, via the regulation of GSK3B activity. In turn, membrane-boun	
Form	Liquid	
Conjugate	Non-conjugated	
Storage Buffer	Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.	
Purification Method	Affinity-chromatography	

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Isotype	Rabbit IgG		
Clonality	Monoclonal		
Product Type	Recombinant Antibody		
Immunogen Species	Homo sapiens (Human)		
Research Area	Cancer; Signal transduction		
Gene Names	RHOA		
Clone No.	1C11		
Image	$72KD \rightarrow 5KS^{51}$ $55KD \rightarrow 36KD \rightarrow 28KD \rightarrow 17KD \rightarrow 10KD \rightarrow 10K$	Western Blot Positive WB detected in: SH-SY5Y whole cell lysate All lanes: Rho antibody at 1:1500 Secondary Goat polyclonal to rabbit IgG at 1/50000 dilution Predicted band size: 22 kDa Observed band size: 22 kDa	

Description

RHOA, a member of the RHO GTPase enzyme family, is a crucial intracellular modulator of dynamics and other functions, including adhesion, proliferation, survival, and gene expression. It takes part in the entire process of cancer progression and plays an important role in tumor cell proliferation, survival and progression, regulating the formation of epithelial polarity, junction assembly, and disruption of epithelial cells. The activation of the RHOA-ROCK signaling pathway is important for both amoeboid and mesenchymal migration. Lamellipodia and uropods are absent in RHOA-depleted cells, which instead feature narrow protrusions projecting from a rounded cell body.

The main steps in the production of this RHOA recombinant antibody include immunization; harvest of positive spleen cells; obtaining the antibody sequence by screening and sequencing; expression of the target antibody in mammalian cells; purification. The RHOA antibody was produced recombinantly and has many advantages: high reproducibility, specificity and scalability. And has been validated in ELISA, WB.