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Datasheet for ABIN6264241 anti-PKC alpha antibody (C-Term)

3 Images



Overview

Quantity:	100 µL
Target:	PKC alpha (PKCa)
Binding Specificity:	C-Term
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This PKC alpha antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF), Immunocytochemistry (ICC)

Product Details

Immunogen:	A synthesized peptide derived from human PKC alpha, corresponding to a region within C- terminal amino acids.
Isotype:	lgG
Specificity:	PKC alpha Antibody detects endogenous levels of total PKC alpha.
Predicted Reactivity:	Bovine,Horse,Sheep,Rabbit,Dog,Chicken,Xenopus
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific).

Target Details

Target:

PKC alpha (PKCa)

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Alternative Name:	PRKCA (PKCa Products)
Background:	Description: Calcium-activated, phospholipid- and diacylglycerol (DAG)-dependent
	serine/threonine-protein kinase that is involved in positive and negative regulation of cell
	proliferation, apoptosis, differentiation, migration and adhesion, tumorigenesis, cardiac
	hypertrophy, angiogenesis, platelet function and inflammation, by directly phosphorylating
	targets such as RAF1, BCL2, CSPG4, TNNT2/CTNT, or activating signaling cascade involving
	MAPK1/3 (ERK1/2) and RAP1GAP. Involved in cell proliferation and cell growth arrest by
	positive and negative regulation of the cell cycle. Can promote cell growth by phosphorylating
	and activating RAF1, which mediates the activation of the MAPK/ERK signaling cascade,
	and/or by up-regulating CDKN1A, which facilitates active cyclin-dependent kinase (CDK)
	complex formation in glioma cells. In intestinal cells stimulated by the phorbol ester PMA, ca
	trigger a cell cycle arrest program which is associated with the accumulation of the hyper-
	phosphorylated growth-suppressive form of RB1 and induction of the CDK inhibitors CDKN1
	and CDKN1B. Exhibits anti-apoptotic function in glioma cells and protects them from apopto
	by suppressing the p53/TP53-mediated activation of IGFBP3, and in leukemia cells mediates
	anti-apoptotic action by phosphorylating BCL2. During macrophage differentiation induced b
	macrophage colony-stimulating factor (CSF1), is translocated to the nucleus and is associat
	with macrophage development. After wounding, translocates from focal contacts to
	lamellipodia and participates in the modulation of desmosomal adhesion. Plays a role in cell
	motility by phosphorylating CSPG4, which induces association of CSPG4 with extensive
	lamellipodia at the cell periphery and polarization of the cell accompanied by increases in ce
	motility. During chemokine-induced CD4+ T cell migration, phosphorylates CDC42-guanine
	exchange factor DOCK8 resulting in its dissociation from LRCH1 and the activation of GTPa
	CDC42 (PubMed:28028151). Is highly expressed in a number of cancer cells where it can ac
	a tumor promoter and is implicated in malignant phenotypes of several tumors such as
	gliomas and breast cancers. Negatively regulates myocardial contractility and positively
	regulates angiogenesis, platelet aggregation and thrombus formation in arteries. Mediates
	hypertrophic growth of neonatal cardiomyocytes, in part through a MAPK1/3 (ERK1/2)-
	dependent signaling pathway, and upon PMA treatment, is required to induce cardiomyocyte
	hypertrophy up to heart failure and death, by increasing protein synthesis, protein-DNA ratio
	cell surface area. Regulates cardiomyocyte function by phosphorylating cardiac troponin T
	(TNNT2/CTNT), which induces significant reduction in actomyosin ATPase activity,
	myofilament calcium sensitivity and myocardial contractility. In angiogenesis, is required for
	endothelial cell migration, adhesion to vitronectin (VTN), and vascular endothelial growth fac
	A (VEGFA)-dependent regulation of kinase activation and vascular tube formation. Involved i

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Preservative:

	the stabilization of VEGFA mRNA at post-transcriptional level and mediates VEGFA-induced cell proliferation. In the regulation of calcium-induced platelet aggregation, mediates signals from the CD36/GP4 receptor for granule release, and activates the integrin heterodimer ITGA2B- ITGB3 through the RAP1GAP pathway for adhesion. During response to lipopolysaccharides (LPS), may regulate selective LPS-induced macrophage functions involved in host defense and inflammation. But in some inflammatory responses, may negatively regulate NF-kappa-B- induced genes, through IL1A-dependent induction of NF-kappa-B inhibitor alpha (NFKBIA/IKBA). Upon stimulation with 12-O-tetradecanoylphorbol-13-acetate (TPA), phosphorylates EIF4G1, which modulates EIF4G1 binding to MKNK1 and may be involved in the regulation of EIF4E phosphorylation. Phosphorylates KIT, leading to inhibition of KIT activity. Phosphorylates ATF2 which promotes cooperation between ATF2 and JUN, activating transcription.
Molecular Weight:	80kDa
Gene ID:	5578
UniProt:	P17252
Pathways:	WNT Signaling, TCR Signaling, EGFR Signaling Pathway, Neurotrophin Signaling Pathway, Thyroid Hormone Synthesis, cAMP Metabolic Process, Myometrial Relaxation and Contraction, Cell-Cell Junction Organization, Regulation of G-Protein Coupled Receptor Protein Signaling, G- protein mediated Events, Signaling Events mediated by VEGFR1 and VEGFR2, Interaction of EGFR with phospholipase C-gamma, Thromboxane A2 Receptor Signaling, VEGFR1 Specific Signals, VEGF Signaling
Application Details	
Application Notes:	WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.

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Sodium azide

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Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months

Images





Immunofluorescence (fixed cells)

Image 1. ABIN6269167 staining NIH-3T3 by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary antibody.

Immunohistochemistry

Image 2. ABIN6269167 at 1/100 staining human Kidney tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the antibody for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit antibody was used as the secondary.

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Western Blotting

Image 3. Western blot analysis of PKC alpha expression in UV treated NIH-3T3 whole cell lysates,The lane on the left is treated with the antigen-specific peptide.

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