

## Phospho-PKCD (Tyr64) Ab

Cat.#: AF3410 Concn.: 1mg/ml Mol.Wt.: 78kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000, IF/ICC 1:100-1:500

Reactivity: Human, Mouse, Rat

Purification: The Ab is from purified rabbit serum by affinity purification

via sequential chromatography on phospho- and non-

phospho-peptide affinity columns.

Specificity: Phospho-PKCD (Tyr64) Ab detects endogenous levels of

PKCD only when phosphorylated at Tyrosine 64.

Immunogen: A synthesized peptide derived from human PKCD around the

phosphorylation site of Tyrosine 64.

Uniprot: Q05655

Description: Protein kinase C (PKC) is a family of serine- and threonine-

specific protein kinases that can be activated by calcium and the second messenger diacylglycerol. PKC family members phosphorylate a wide variety of protein targets and are known to be involved in diverse cellular signaling pathways.

Subcellular Location: Cytoplasm. Membrane.

Similarity: The C1 domain, containing the phorbol ester/DAG-type

region 1 (C1A) and 2 (C1B), is the diacylglycerol sensor. The C2 domain is a non-calcium binding domain. It binds proteins

containing phosphotyrosine in a sequence-specific manner.Belongs to the protein kinase superfamily. AGC

Ser/Thr protein kinase family. PKC subfamily.

Storage Condition and

Buffer:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

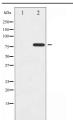
°C.Stable for 12 months from date of receipt.



Western blot analysis of Phospho-PKCD (Tyr64) expression in various lysates



## Affinity Biosciences website:www.affbiotech.com order:order@affbiotech.com



Western blot analysis of PKCD phosphorylation expression in PMA treated HepG2 whole cell lysates, The lane on the left is treated with the antigen-specific peptide.



AF3410 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 Ab 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 Ab.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1% TBS, 0.1% Tween@20 at 4°C with gentle shaking, overnight.

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