

Phospho-PKD1/2/3/PKC μ (Ser738+Ser742) Ab

Cat.#: AF3444 Concn.: 1mg/ml Mol.Wt.: 101kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000 IHC 1:50-1:200, 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-PKD1/2/3/PKC μ (Ser738+Ser742) Ab detects

endogenous levels of PKD1/2/3/PKC μ only when phosphorylated at Serine 738+Serine 742.

Immunogen: A synthesized peptide derived from human PKD1/2/3/PKC µ

around the phosphorylation site of Serine 738+Serine 742.

Uniprot: Q15139/Q9BZL6/O94806

Description: PKD3 a CAMK kinase of the PKD family. An important

component of signaling pathways downstream from novel

PKC enzymes after B-cell receptor engagement.

Subcellular Location: Cytoplasm. Membrane. Translocation to the cell membrane

is required for kinase activation.

Tissue Specificity: Up-regulated by the intestine-specific transcription factor

CDX1 in an activated KRAS-dependent manner in colorectal

cancer (CRC) cells (PubMed:24623306).

Similarity: Belongs to the protein kinase superfamily. CAMK Ser/Thr

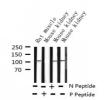
protein kinase family. PKD subfamily.

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.

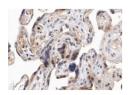


Western blot analysis of Phospho-PKD1/2/3/PKC μ (Ser738+Ser742) expression in various lysates



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AF3444 at 1/100 staining human Placenta 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 Ab for 1.5 hours at 22°C. An HRP conjugated goat antirabbit Ab was used as the secondary.



AF3444 staining A549 cells 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(Red), diluted at 1/600, was used as 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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