Phospho-PRKACA-S339 Rabbit Polyclonal Antibody

CABP0558

Product Information

Size:

20uL, 50uL, 100uL, 200uL

Observed MW:

42kDa

Calculated MW:

39kDa/40kDa

Applications:

WB IP

Reactivity:

Human

Antibody Information

Recommended dilutions:
WB 1:500 - 1:2000 IP 1:50 - 1:100

Source:
Rabbit

Isotype:
IgG

Purification:
Affinity purification

Protein Background

This gene encodes one of the catalytic subunits of protein kinase A, which exists as a tetrameric holoenzyme with two regulatory subunits and two catalytic subunits, in its inactive form. cAMP causes the dissociation of the inactive holoenzyme into a dimer of regulatory subunits bound to four cAMP and two free monomeric catalytic subunits. Four different regulatory subunits and three catalytic subunits have been identified in humans. cAMP-dependent phosphorylation of proteins by protein kinase A is important to many cellular processes, including differentiation, proliferation, and apoptosis. Constitutive activation of this gene caused either by somatic mutations, or genomic duplications of regions that include this gene, have been associated with hyperplasias and adenomas of the adrenal cortex and are linked to corticotropin-independent Cushing's syndrome. Alternative splicing results in multiple transcript variants encoding different isoforms. Tissue-specific isoforms that differ at the N-terminus have been described, and these isoforms may differ in the post-translational modifications that occur at the N-terminus of some isoforms.

Immunogen information

Gene ID:
5566

Uniprot
P17612

Synonyms:
PRKACA; PKACA; PPNAD4

Immunogen:
A synthetic phosphorylated peptide around S339 of human PRKACA (NP_002721.1).

Storage:
Store at -20℃. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

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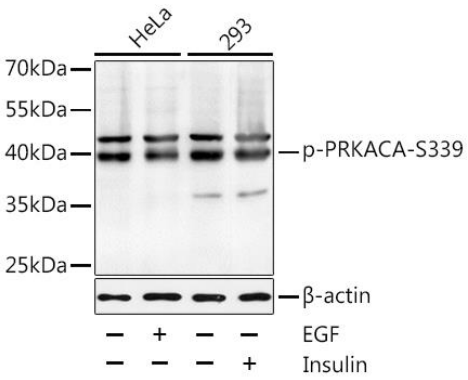
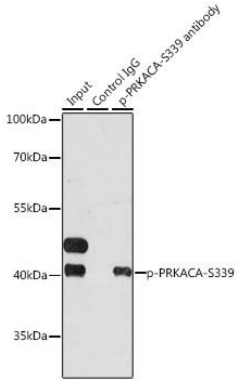
Product Images



Western blot analysis of extracts of HeLa and 293 cells, using Phospho-PRKACA-S339 antibody (CABP0558) at 1:1000 dilution. HeLa cells were treated by EGF (100ng/mL) for 30 minutes after serum-starvation overnight.293T cells were treated by Insulin (100nM) for 10 minutes after serum-starvation overnight. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (CABS014) at 1:10000 dilution. Lysates/proteins: 25ug per lane. Blocking buffer: 3% BSA.



Immunoprecipitation analysis of 200ug extracts of HeLa cells, using 3 ug Phospho-PRKACA-S339 pAb (CABP0558). Western blot was performed from the immunoprecipitate using Phospho-PRKACA-S339 pAb (CABP0558) at a dilition of 1:1000. HeLa cells were treated by EGF (100 ng/mL) at 37'C for 30 minutes after serum-starvation overnight.

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