For Research Use Only

Phospho-AKT (Thr308) Recombinant antibody

Catalog Number:81232-10-RR



Basic Information

Catalog Number: GenBank Accession Number: 81232-10-RR BC000479

Concentration: GenelD (NCBI): 1000 µ g/ml 207

Source: UNIPROT ID:
Rabbit P31749
Isotype: Full Name:

gG v-akt murine thymoma viral oncogene homolog 1

> Calculated MW: 56 kDa Observed MW: 60 kDa

Purification Method:

242063B2

Protein A purification CloneNo.:

Recommended Dilutions: WB 1:2000-1:10000

Applications

Tested Applications: WB, FC (Intra), ELISA Species Specificity: human, mouse, rat

Positive Controls:

WB: Calyculin A treated HSC-T6 cells, Calyculin A treated HEK-293 cells, Calyculin A treated NIH/3T3

cells

Background Information

AKT is a serine/threonine kinase and it participates in the key role of the PI3K signaling pathway. Phosphatidylinositol-3 kinase (PI3K) is the key regulator of AKT activation. The recruitment of inactive AKT protein to PIP3-rich areas of the plasma membrane results in a conformational change that exposes the activation loop of AKT. AKT's activating kinase, phosphoinositide-dependent protein kinase (PDK1), is also recruited to PIP3 microdomains. PDK1 phosphorylates AKT on threonine 308 (Thr308) of the exposed activation loop, activating AKT and leading to a second phosphorylation of AKT at serine 473 (Ser473) by a kinase presumed to be mTORC2 that further potentiates kinase activity. Active AKT will phosphorylate various downstream protein targets that control cell growth and translational control and act to suppress apoptosis. (PMID: 31594388, PMID: 30808672)

Storage

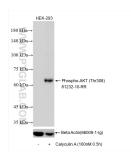
Storage:

Store at -20°C. Stable for one year after shipment. Storage Buffer:

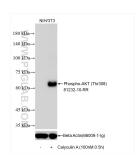
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

Aliquoting is unnecessary for -20°C storage

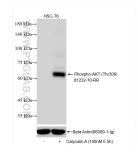
Selected Validation Data



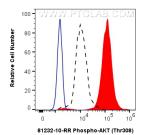
Non-treated HEK-293 cells and Calyculin A treated HEK-293 cells were subjected to SDS PAGE followed by western blot with 81232-10-RR (Phospho-AKT (Thr308) antibody) at dilution of 1.5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Actin (66009-1-1g) antibody as a loading control.



Non-treated NIH/3T3 cells and Calyculin A treated NIH/3T3 cells were subjected to SDS PAGE followed by western blot with 81232-10-RR (Phospho-AKT (Thr308) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Actin (66009-1-lg) antibody as a loading control



Non-treated HSC-T6 cells and Calyculin A treated HSC-T6 cells were subjected to SDS PAGE followed by western blot with 81232-10-RR (Phospho-AKT (Thr308) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Actin (66009-1-lg) antibody as a loading control.



1X10^6 HeLa cells untreated (dashed lines) or treated with Calyculin A which intracellularly stained with 0.13 ug Phospho-AKT (Thr308) Recombinant antibody (81232-10-RR, Clone:242063B2) and Coralite®488-Conjugated Goat Anti-Rabbit IgG(H+L) (SA00013-2)(red), or 0.13 ug Rabbit IgG Isotype Control RecAb (98136-1-RR, Clone: 240953C9) (blue). Cells were fixed with 4% PFA and permeabilized with 90% MeOH.