For Research Use Only

Phospho-PERK/EIF2AK3 (Thr982) Recombinant antibody

Catalog Number:82534-1-RR 2 Publications



Basic Information

Catalog Number: 82534-1-RR Concentration: 1000 ug/ml

Source: Rabbit Isotype:

GenBank Accession Number: BC126354

GeneID (NCBI): **UNIPROT ID:** Q9NZJ5

Full Name: eukaryotic translation initiation factor 2-alpha kinase 3

Calculated MW: 1116 aa, 125 kDa Observed MW: 180 kDa

Purification Method:

Protein A purification

CloneNo.: 4E16

Recommended Dilutions: WB 1:2000-1:11200

Applications

Tested Applications: WB, FC (Intra), ELISA Cited Applications:

Species Specificity: human, mouse Cited Species: human

Positive Controls:

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

Background Information

EIF2AK3 encodes the protein kinase RNA-like ER kinase (PERK), a key regulator of the unfolded protein response (UPR) in response to ER stress. Under ER stress conditions, activation of PERK is triggered by the dissociation of glucose-regulated protein (GRP) 78 (also known as BiP) from its luminal domain, followed by oligomerization and autophosphorylation. Phosphorylated PERK subsequently phosphorylates eukaryotic translation initiation factor 2 alpha (eif2 a), to attenuate global protein translation and reduce incoming ER protein load via upregulated ER chaperone expression. (PMID: 35922637, PMID: 32029570)

Notable Publications

Author	Pubmed ID	Journal	Application
Yu Feng	39138149	Cell Death Dis	WB
Yu Han	38513524	Vet Microbiol	WB

Storage

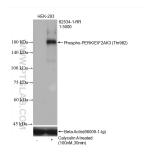
Store at -20°C. Stable for one year after shipment.

Storage Buffer:

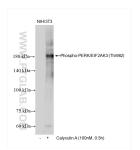
PBS with 0.02% sodium azide and 50% glycerol, pH7.3

Aliquoting is unnecessary for -20°C storage

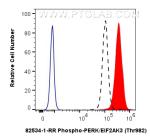
Selected Validation Data



Non-treated HEK-293 cells and Calyculin A treated HEK-293 cells were subjected to SDS PAGE followed by western blot with 82534-1-RR (Phospho-PERK/EIF2AK3 (Thr982) antibody) at dilution of 1:5000 incubated at room temperature for 1.5 hours. The membrane was stripped and reblotted with beta actin antibody (66009-1-lg) as loading control.



Non-treated NIH/3T3 cells and Calyculin A treated NIH/3T3 cells were subjected to SDS PAGE followed by western blot with 82534-1-RR (Phospho-PERK/EIF2AK3 (Thr982) antibody) at dilution of 1:2500 incubated at room temperature for 1.5 hours.



1X10^6 HEK-293 cells untreated (dashed lines) or treated with Calyculin A (red) were intracellularly stained with 0.13 ug Phospho-PERK/EIF2AK3 (Thr982) Recombinant antibody (82534-1-RR, Clone:4E16) and Coralite®488-Conjugated Goat Anti-Rabbit 1gG(H+L) (5A00013-2)(red), or 0.13 ug Rabbit 1gG Isotype Control Recombinant Antibody (98136-1-RR, Clone: 240953C9) (blue). Cells were fixed with 49% PFA and permeabilized with 90% MeOH.