For Research Use Only

## Phospho-PERK/EIF2AK3 (Ser719) Polyclonal antibody



Catalog Number: 29546-1-AP

7 Publications

**Basic Information** 

Catalog Number: 29546-1-AP Size: 250 µ g/ml Source: Rabbit

Isotype: IgG GenBank Accession Number:

BC126354

GeneID (NCBI):
9451

UNIPROT ID:
Q9NZJ5

Full Name:

eukaryotic translation initiation factor 2-alpha kinase 3

Calculated MW: 1116 aa, 125 kDa Observed MW: 140 kDa Purification Method: Antigen affinity purification Recommended Dilutions:

WB 1:500-1:2000

**Applications** 

Tested Applications: WB, ELISA
Cited Applications: WB

Species Specificity: Human Cited Species: human, rat, mouse Positive Controls:

WB:  $\lambda\,$  phosphatase treated HeLa cells, Thapsigargin treated HeLa 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
Guanjun Li	38452464	Environ Int	WB
Yun Huang	38132921	Mar Drugs	WB
Jie Cui	37348685	Fish Shellfish Immunol	WB

Storage

Storage:

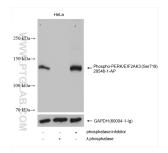
Store at -20°C. Stable for one year after shipment.

Storage Buffer:

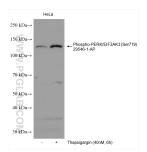
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

Aliquoting is unnecessary for -20°C storage

## **Selected Validation Data**



Non-treated HeLa cells, phosphatase inhibitor treated and  $\lambda$  phosphatase treated HeLa cells were subjected to SDS PAGE followed by western blot with 29546-1-AP (Phospho-PERK/EIF2AK3 (Ser719) antibody) at dilution of 1:1000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with GAPDH antibody as loading control.



Non-treated HeLa cells and Thapsigargin treated HeLa cells were subjected to SDS PAGE followed by western blot with 29546-1-AP (Phospho-PERK/EIF2AK3 (Ser719) antibody) at dilution of 1:1500 incubated at room temperature for 1.5 hours.