

À des fins de recherche uniquement

# Anticorps Polyclonal de lapin anti- Phospho-PERK/EIF2AK3 (Ser719)

Numéro de catalogue: 29546-1-AP    5 Publications



## Informations de base

Numéro de catalogue:	BC126354	Méthode de purification:
29546-1-AP		Purification par affinité contre l'antigène
Taille:	Identification du gène (NCBI):	Dilutions recommandées:
100ul , Concentration: 250 µg/ml by Nanodrop;	9451	WB 1:500-1:2000
Hôte:	Nom complet:	
Lapin	eukaryotic translation initiation factor 2-alpha kinase 3	
Isotype:	MW calculé	
IgG	1116 aa, 125 kDa	
	MW observés:	
	140 kDa	

## Applications

Applications testées:	Contrôles positifs:
WB, ELISA	WB: cellules HeLa traitées à la λ phosphatase,
Demandes citées:	Thapsigargin treated HeLa cells
WB	
Spécificité de l'espèce:	
Humain	
Espèces citées:	
Humain, rat	

## Informations générales

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 (eif2α), to attenuate global protein translation and reduce incoming ER protein load via upregulated ER chaperone expression. (PMID: 35922637, PMID: 32029570)

## Publications notables

Autrice	Pubmed ID	Journal	Application
Jie Cui	37348685	Fish Shellfish Immunol	WB
Ziwei Cui	37329685	Tissue Cell	WB
Xiao-Xue Hou	37213406	World J Gastroenterol	WB

## Stockage

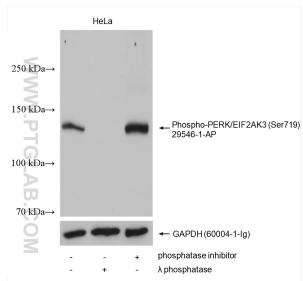
**Stockage:**  
Stocker à -20°C. Stable pendant un an après l'expédition.  
**Tampon de stockage:**  
PBS avec azoture de sodium à 0,02 % et glycérol à 50 % pH 7,3  
L'aliquotage n'est pas nécessaire pour le stockage à -20C

\*\*\* Les 20ul contiennent 0,1% de BSA.

For technical support and original validation data for this product please contact:  
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## Données de validation sélectionnées



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.

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.