

À des fins de recherche uniquement

Anticorps Recombinant de lapin anti- Phospho-PERK/EIF2AK3 (Thr982)



Numéro de catalogue: 82534-1-RR

Informations de base

Numéro de catalogue:	BC126354	Méthode de purification:
82534-1-RR		Purification par protéine A
Taille:	Identification du gène (NCBI):	CloneNo.:
100ul , Concentration: 500 µg/ml by Nanodrop;	9451	4E16
Hôte:	Nom complet:	Dilutions recommandées:
Lapin	eukaryotic translation initiation factor 2-alpha kinase 3	WB 1:2000-1:11200
Isotype:	MW calculé	
IgG	1116 aa, 125 kDa	
	MW observés:	
	180 kDa	

Applications

Applications testées:	Contrôles positifs:
WB, ELISA	WB : cellules HEK-293 traitées à la calyculine A,
Spécificité de l'espèce:	
Humain	

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 autoprophosphorylation. 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)

Stockage

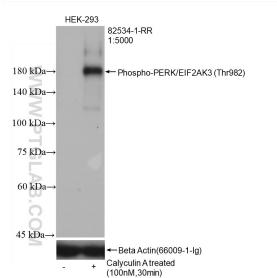
Stockage:
Stocker à -20°C. Stable pendant un an après l'expédition.
Tampon de stockage:
PBS avec azoture de sodium à 0,02 % et glycérol à 50 % pH 7,3
L'aliquotage n'est pas nécessaire pour le stockage à -20C

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

For technical support and original validation data for this product please contact:
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in USA), or 1(312) 455-8498 (outside USA) E: proteintech@ptglab.com
W: ptglab.com

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Données de validation sélectionnées



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 re-blotted with beta actin antibody (66009-1-Ig) as loading control.