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## Phospho-PERK (Thr980) Rabbit pAb

Catalog Number: bs-3330R

Target Protein: Phospho-PERK (Thr980)

Concentration: 1mg/ml

Form: Liquid

Host: Rabbit

Clonality: Polyclonal

Isotype: IgG

Applications: WB (1:500-2000), IHC-P (1:100-500), IHC-F (1:100-500), IF (1:100-500), Flow-Cyt (2ug/Test)

Reactivity: Human, Mouse, Rat (predicted:Rabbit, Pig, Cow, Dog)

Predicted MW: 119 kDa Entrez Gene: 13666 Swiss Prot: Q9Z2B5

Source: KLH conjugated synthesised phosphopeptide derived from mouse PERK around the

phosphorylation site of Thr980: H(p-T)GQ.

Purification: affinity purified by Protein A

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

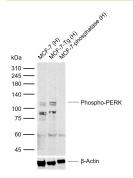
Background: The protein encoded by this gene phosphorylates the alpha subunit of eukaryotic

translation-initiation factor 2 (EIF2), leading to its inactivation, and thus to a rapid reduction of translational initiation and repression of global protein synthesis. It is a type I membrane protein located in the endoplasmic reticulum (ER), where it is induced by ER stress caused

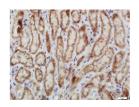
by malfolded proteins. Mutations in this gene are associated with Wolcott-Rallison

syndrome. [provided by RefSeq, Jan 2010].

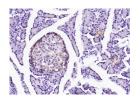
## **VALIDATION IMAGES**



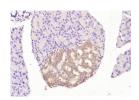
Sample: Lane 1: Human MCF-7 cell lysates Lane 2: Human MCF-7-Tg (1  $\mu$ M thapsigargin, 20 min) cell lysates Lane 3: Human MCF-7-phosphatase cell lysates Primary: Anti-Phospho-PERK (Thr980) (bs-3330R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 119 kDa Observed band size: 120 kDa



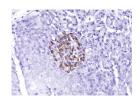
Paraformaldehyde-fixed, paraffin embedded (rat kidney tissue); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-PERK(Thr980)) Polyclonal Antibody, Unconjugated (bs-3330R) at 1:400 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.



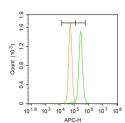
Paraformaldehyde-fixed, paraffin embedded (mouse pancreas); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-PERK (Thr980)) Polyclonal Antibody, Unconjugated (bs-3330R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (rat pancreas); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-PERK (Thr980)) Polyclonal Antibody, Unconjugated (bs-3330R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (human pancreas); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-PERK (Thr980)) Polyclonal Antibody, Unconjugated (bs-3330R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Blank control: A431. Primary Antibody (green line): Rabbit Anti-PERK(Thr980) antibody (bs-3330R) Dilution:  $3\mu g/10^6$  cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody: Goat anti-rabbit IgG-AF647 Dilution:  $3\mu g/test$ . Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 0.1% PBST for 20 min at room temperature. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature. Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

## PRODUCT SPECIFIC PUBLICATIONS

[IF=41.845] Ashley R. Helseth. et al. Cholinergic neurons constitutively engage the ISR for dopamine modulation and skill learning in mice. Science. 2021 Apr;372(6540): IHC; Mouse . 33888613

[IF=25.606] Raines, Lydia N.. et al. PERK is a critical metabolic hub for immunosuppressive function in macrophages. Nat Immunol. 2022 Feb;23(3):431-445 FCM,WB; Mouse . 10.1038/s41590-022-01145-x

[IF=14.7] Anna Flury. et al. A neurodegenerative cellular stress response linked to dark microglia and toxic lipid secretion. NEURON. 2024 Dec 23 IF; Mouse . 39719704

[IF=12.8] Xinli Wang. et al. Sustained therapeutic effects of self-assembled hyaluronic acid nanoparticles loaded with  $\alpha$ -Ketoglutarate in various osteoarthritis stages. BIOMATERIALS. 2024 Sep;:122845 WB; MOUSE . 39326362

[IF=9.988] Yue Zhang. et al. Endoplasmic reticulum stress-controlled autophagic pathway promotes polystyrene microplastics-induced myocardial dysplasia in birds. ENVIRON POLLUT. 2022 Oct;311:119963 WB; Chicken. 35973452