

Rabbit Anti-PERK antibody

SL2469R

Product Name:	PERK
Chinese Name:	蛋白激 酶样内质网激酶抗体
Alias:	DKFZp781H1925; E2AK3_HUMAN; EC 2.7.11.1; EIF2AK3; Eukaryotic translation initiation factor 2 alpha kinase 3; Eukaryotic translation initiation factor 2-alpha kinase 3; Heme regulated EIF2 alpha kinase; HRI; HsPEK; Pancreatic eIF2 alpha kinase; Pancreatic eIF2-alpha kinase; PEK; PRKR like endoplasmic reticulum kinase; PRKR- like endoplasmic reticulum kinase; WRS.
	Specific References(4) SL2469R has been referenced in 4 publications.
	[IF=2.33]He, Yihuai, et al. "Sustained endoplasmic reticulum stress inhibits hepatocyte
	proliferation via downregulation of c-Met expression." Molecular and Cellular
	Biochemistry (2014): 1-8.WB;Human.
	PubMed:24390087
	[IF=0.00]Wang, Yu, et al. "Tanshinone II A Relieves Adriamycin-induced Myocardial
文献引用	Injury in Rat Model." International Journal of Chemistry 8.1 (2016): 40.WB;Rat.
Pub	PubMed:not posted yet
	[IF=4.65]Yu, H., et al. "Gypenoside Protects against Myocardial Ischemia-Reperfusion
	Injury by Inhibiting Cardiomyocytes Apoptosis via Inhibition of CHOP Pathway and
	Activation of PI3K/Akt Pathway In Vivo and In Vitro."Cellular Physiology and
	Biochemistry 39.1 (2016): 123-136.WB;Rat.
	<u>PubMed:27322831</u>
	[IF=3.29]Yan, Jiting, et al. "Catalpol prevents alteration of cholesterol homeostasis in
	non-alcoholic fatty liver disease via attenuating endoplasmic reticulum stress and NOX4
	over-expression." RSC Advances 7.2 (2017): 1161-1176.WB;Human.

	PubMed:0
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human, Mouse, Rat,
	ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800Flow-Cyt=1µg/TestIF=1:100- 500 (Paraffin sections need antigen repair)
Applications:	not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	122kDa
Cellular localization:	cytoplasmic
Form:	Lyophilized or Liquid
Concentration:	lmg/ml
immunogen:	KLH conjugated synthetic peptide derived from human PERK:1001-1116/1116
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20 °C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	PubMed
	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]
Product Detail:	Function: 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. Serves as a critical effector of unfolded protein response (UPR)-induced G1 growth arrest due to the loss of cyclin-D1 (CCND1). Subunit: Forms dimers with HSPA5/BIP in resting cells. Oligomerizes in ER-stressed cells.
	Subcellular Location: Endoplasmic reticulum membrane; Single-pass type I membrane protein. Tissue Specificity: Ubiquitous. A high level expression is seen in secretory tissues. Post-translational modifications:

Oligomerization of the N-terminal ER luminal domain by ER stress promotes PERK trans-autophosphorylation of the C-terminal cytoplasmic kinase domain at multiple residues including Thr-982 on the kinase activation loop. Autophosphorylated. Phosphorylated at Tyr-619 following endoplasmic reticulum stress, leading to activate its tyrosine-protein kinase activity. Dephosphorylated by PTPN1/TP1B, leading to inactivate its enzyme activity.

N-glycosylated.

ADP-ribosylated by PARP16 upon ER stress, which increases kinase activity.

DISEASE:

Wolcott-Rallison syndrome (WRS) [MIM:226980]: A rare autosomal recessive disorder, characterized by permanent neonatal or early infancy insulin-dependent diabetes and, at a later age, epiphyseal dysplasia, osteoporosis, growth retardation and other multisystem manifestations, such as hepatic and renal dysfunctions, mental retardation and cardiovascular abnormalities. Note=The disease is caused by mutations affecting the gene represented in this entry.

Similarity:

Belongs to the protein kinase superfamily. Ser/Thr protein kinase family. GCN2 subfamily.

Contains 1 protein kinase domain.

SWISS: Q9NZJ5

Gene ID: 9451

Database links:

Entrez Gene: 9451Human

Entrez Gene: 13666Mouse

Entrez Gene: 29702Rat

Omim: 604032Human

SwissProt: Q9NZJ5Human

SwissProt: Q9Z2B5Mouse

SwissProt: Q9Z1Z1Rat

Unigene: 591589Human





incubated for 30 min on the ice, followed by 1 X PBS containing 0.5% BSA + 1 0%
goat serum (15 min) to block non-specific protein-protein interactions. Then the
Goat Anti-rabbit IgG/PE antibody was added into the blocking buffer mentioned
above to react with the primary antibody at 1/200 dilution for 30 min on ice.
Acquisition of 20,000 events was performed.

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