

PKR (PRKR) Antibody (C-term T535) Blocking peptide
Synthetic peptide
Catalog # BP8151d**Specification**

PKR (PRKR) Antibody (C-term T535) Blocking peptide - Product Information

Primary Accession [P19525](#)
Other Accession [Q52M43](#)

PKR (PRKR) Antibody (C-term T535) Blocking peptide - Additional Information

Gene ID 5610

Other Names

Interferon-induced, double-stranded RNA-activated protein kinase, Eukaryotic translation initiation factor 2-alpha kinase 2, eIF-2A protein kinase 2, Interferon-inducible RNA-dependent protein kinase, P1/eIF-2A protein kinase, Protein kinase RNA-activated, PKR, Protein kinase R, Tyrosine-protein kinase EIF2AK2, p68 kinase, EIF2AK2, PKR, PRKR

Target/Specificity

The synthetic peptide sequence used to generate the antibody [AP8151d](/product/products/AP8151d) was selected from the C-term region of human PRKR-T535 (C-term). A 10 to 100 fold molar excess to antibody is recommended. Precise conditions should be optimized for a particular assay.

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

Precautions

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

PKR (PRKR) Antibody (C-term T535) Blocking peptide - Protein Information

Name EIF2AK2

Synonyms PKR, PRKR

Function

IFN-induced dsRNA-dependent serine/threonine-protein kinase that phosphorylates the alpha subunit of eukaryotic translation initiation factor 2 (EIF2S1/eIF-2-alpha) and plays a key role in the innate immune response to viral infection (PubMed:[18835251](http://www.uniprot.org/citations/18835251), PubMed:[19507191](http://www.uniprot.org/citations/19507191), PubMed:[19189853](http://www.uniprot.org/citations/19189853), PubMed:

[21123651](http://www.uniprot.org/citations/21123651) PubMed: [21072047](http://www.uniprot.org/citations/21072047) PubMed: [22948139](http://www.uniprot.org/citations/22948139) PubMed: [23229543](http://www.uniprot.org/citations/23229543) PubMed: [22381929](http://www.uniprot.org/citations/22381929)). Inhibits viral replication via the integrated stress response (ISR): EIF2S1/eIF-2- alpha phosphorylation in response to viral infection converts EIF2S1/eIF-2-alpha in a global protein synthesis inhibitor, resulting to a shutdown of cellular and viral protein synthesis, while concomitantly initiating the preferential translation of ISR-specific mRNAs, such as the transcriptional activator ATF4 (PubMed: [19189853](http://www.uniprot.org/citations/19189853) PubMed: [21123651](http://www.uniprot.org/citations/21123651) PubMed: [22948139](http://www.uniprot.org/citations/22948139) PubMed: [23229543](http://www.uniprot.org/citations/23229543)). Exerts its antiviral activity on a wide range of DNA and RNA viruses including hepatitis C virus (HCV), hepatitis B virus (HBV), measles virus (MV) and herpes simplex virus 1 (HHV-1) (PubMed: [11836380](http://www.uniprot.org/citations/11836380) PubMed: [19189853](http://www.uniprot.org/citations/19189853) PubMed: [20171114](http://www.uniprot.org/citations/20171114) PubMed: [19840259](http://www.uniprot.org/citations/19840259) PubMed: [21710204](http://www.uniprot.org/citations/21710204) PubMed: [23115276](http://www.uniprot.org/citations/23115276) PubMed: [23399035](http://www.uniprot.org/citations/23399035)). Also involved in the regulation of signal transduction, apoptosis, cell proliferation and differentiation: phosphorylates other substrates including p53/TP53, PPP2R5A, DHX9, ILF3, IRS1 and the HHV-1 viral protein US11 (PubMed: [11836380](http://www.uniprot.org/citations/11836380) target="_blank">22214662 target="_blank">19229320). In addition to serine/threonine- protein kinase activity, also has tyrosine-protein kinase activity and phosphorylates CDK1 at 'Tyr-4' upon DNA damage, facilitating its ubiquitination and proteasomal degradation (PubMed: [20395957](http://www.uniprot.org/citations/20395957)). Either as an adapter protein and/or via its kinase activity, can regulate various signaling pathways (p38 MAP kinase, NF-kappa-B and insulin signaling pathways) and transcription factors (JUN, STAT1, STAT3, IRF1, ATF3) involved in the expression of genes encoding pro-inflammatory cytokines and IFNs (PubMed: [22948139](http://www.uniprot.org/citations/22948139) PubMed: [23084476](http://www.uniprot.org/citations/23084476) PubMed: [23372823](http://www.uniprot.org/citations/23372823)). Activates the NF-kappa-B pathway via interaction with IKBKB and TRAF family of proteins and activates the p38 MAP kinase pathway via interaction with MAP2K6 (PubMed: [10848580](http://www.uniprot.org/citations/10848580) PubMed: [15121867](http://www.uniprot.org/citations/15121867) PubMed: [15229216](http://www.uniprot.org/citations/15229216)). Can act as both a positive and negative regulator of the insulin signaling pathway (ISP) (PubMed: [20685959](http://www.uniprot.org/citations/20685959)). Negatively regulates ISP by inducing the inhibitory phosphorylation of insulin receptor substrate 1 (IRS1) at 'Ser-312' and positively regulates ISP via phosphorylation of PPP2R5A which activates FOXO1, which in turn up-regulates the expression of insulin receptor substrate 2 (IRS2) (PubMed: [20685959](http://www.uniprot.org/citations/20685959)). Can regulate NLRP3 inflammasome assembly and the activation of NLRP3, NLRP1, AIM2 and NLRC4 inflammasomes (PubMed: [22801494](http://www.uniprot.org/citations/22801494)). Plays a role in the regulation of the cytoskeleton by binding to gelsolin (GSN), sequestering the protein in an inactive conformation away from actin (By similarity).

Cellular Location

Cytoplasm. Nucleus. Cytoplasm, perinuclear region. Note=Nuclear localization is elevated in acute leukemia, myelodysplastic syndrome (MDS), melanoma, breast, colon, prostate and lung cancer

patient samples or cell lines as well as neurocytes from advanced Creutzfeldt- Jakob disease patients.

Tissue Location

Highly expressed in thymus, spleen and bone marrow compared to non-hematopoietic tissues such as small intestine, liver, or kidney tissues. Colocalizes with GSK3B and TAU in the Alzheimer disease (AD) brain. Elevated levels seen in breast and colon carcinomas, and which correlates with tumor progression and invasiveness or risk of progression.

PKR (PRKR) Antibody (C-term T535) Blocking peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

PKR (PRKR) Antibody (C-term T535) Blocking peptide - Images

PKR (PRKR) Antibody (C-term T535) Blocking peptide - Background

Interferon-induced, double-stranded RNA-activated protein kinase (PRKR) is a serine-threonine kinase. Activation by dsRNAs leads to autophosphorylation of PRKR and allows the kinase to phosphorylate its natural substrate, the alpha subunit of eukaryotic protein synthesis initiation factor-2 (EIF2-alpha), leading to the inhibition of protein synthesis. Human gamma-interferon (IFNG) mRNA exploits localized activation of PRKR in the cell to regulate its own translation. IFNG mRNA activates PRKR through a pseudoknot in its 5-prime untranslated region. The HCV envelope protein E2 contains a sequence identical with phosphorylation sites of the interferon-inducible protein kinase PRKR and the translation initiation factor EIF2-alpha, a target of PRKR. E2 inhibits the kinase activity of PRKR and blocks its inhibitory effect on protein synthesis and cell growth, which provides one mechanism by which HCV may circumvent the antiviral effect of interferon. PRKR, which is involved in TLR signaling and mediates apoptosis in fibroblasts in response to viral infection and inflammatory cytokines, also activates IKK and NFkB, thereby suppressing apoptosis. Apoptosis induced by live pathogenic gram-positive and gram-negative bacteria requires both TLR4 and PRKR, possibly representing a major mechanism for pathogenic bacteria that use specific virulence factors to avoid detection and destruction by the innate immune system. Roles for PRKR activation in Huntington disease and Fanconi anemia have also been suggested.

PKR (PRKR) Antibody (C-term T535) Blocking peptide - References

Beisser, P.S., et al., J. Virol. 79(1):441-449 (2005). Gerotto, M., et al., Antivir. Ther. (Lond.) 9(5):763-770 (2004). Zhang, X., et al., J. Biol. Chem. 279(42):43910-43919 (2004). Gil, J., et al., Mol. Cell. Biol. 24(10):4502-4512 (2004). Hii, S.I., et al., Int. J. Cancer 109(3):329-335 (2004).