

DNA PK (PRKDC) Antibody (N-term) Blocking peptide
Synthetic peptide
Catalog # BP7053a**Specification**

DNA PK (PRKDC) Antibody (N-term) Blocking peptide - Product InformationPrimary Accession [P78527](#)**DNA PK (PRKDC) Antibody (N-term) Blocking peptide - Additional Information****Gene ID** 5591**Other Names**

DNA-dependent protein kinase catalytic subunit, DNA-PK catalytic subunit, DNA-PKcs, DNPK1, p460, PRKDC, HYRC, HYRC1

Target/Specificity

The synthetic peptide sequence used to generate the antibody [AP7053a](/product/products/AP7053a) was selected from the N-term region of human PRKDC. 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.

DNA PK (PRKDC) Antibody (N-term) Blocking peptide - Protein Information**Name** PRKDC**Synonyms** HYRC, HYRC1**Function**

Serine/threonine-protein kinase that acts as a molecular sensor for DNA damage (PubMed: [11955432](http://www.uniprot.org/citations/11955432), PubMed: [12649176](http://www.uniprot.org/citations/12649176), PubMed: [14734805](http://www.uniprot.org/citations/14734805), PubMed: [33854234](http://www.uniprot.org/citations/33854234)). Involved in DNA non-homologous end joining (NHEJ) required for double-strand break (DSB) repair and V(D)J recombination (PubMed: [11955432](http://www.uniprot.org/citations/11955432), PubMed: [12649176](http://www.uniprot.org/citations/12649176), PubMed: [14734805](http://www.uniprot.org/citations/14734805)).

target="_blank">14734805, PubMed:33854234, PubMed:34352203). Must be bound to DNA to express its catalytic properties (PubMed:11955432). Promotes processing of hairpin DNA structures in V(D)J recombination by activation of the hairpin endonuclease artemis (DCLRE1C) (PubMed:11955432). Recruited by XRCC5 and XRCC6 to DNA ends and is required to (1) protect and align broken ends of DNA, thereby preventing their degradation, (2) and sequester the DSB for repair by NHEJ (PubMed:15574326, PubMed:11955432, PubMed:12649176, PubMed:14734805, PubMed:33854234). Act as a scaffold protein to aid the localization of DNA repair proteins to the site of damage (PubMed:15574326, PubMed:11955432, PubMed:12649176, PubMed:14734805). The assembly of the DNA-PK complex at DNA ends is also required for the NHEJ ligation step (PubMed:15574326, PubMed:11955432, PubMed:12649176, PubMed:14734805). Found at the ends of chromosomes, suggesting a further role in the maintenance of telomeric stability and the prevention of chromosomal end fusion (By similarity). Also involved in modulation of transcription (PubMed:15574326, PubMed:11955432, PubMed:12649176, PubMed:14734805). As part of the DNA-PK complex, involved in the early steps of ribosome assembly by promoting the processing of precursor rRNA into mature 18S rRNA in the small-subunit processome (PubMed:32103174). Binding to U3 small nucleolar RNA, recruits PRKDC and XRCC5/Ku86 to the small-subunit processome (PubMed:32103174). Recognizes the substrate consensus sequence [ST]-Q (PubMed:15574326, PubMed:11955432, PubMed:12649176, PubMed:14734805). Phosphorylates 'Ser-139' of histone variant H2AX, thereby regulating DNA damage response mechanism (PubMed:14627815, PubMed:16046194). Phosphorylates ASF1A, DCLRE1C, c-Abl/ABL1, histone H1, HSPCA, c-jun/JUN, p53/TP53, PARP1, POU2F1, DHX9, FH, SRF, NHEJ1/XLF, XRCC1, XRCC4, XRCC5, XRCC6, WRN, MYC and RFA2 (PubMed:2507541, PubMed:2247066, PubMed:1597196, PubMed:8407951, PubMed:8464713, PubMed:9362500, PubMed:9139719, PubMed:10026262, PubMed:10467406, PubMed:12509254, PubMed:11889123, PubMed:14612514).

target="_blank">14612514, PubMed:14599745, PubMed:15177042, PubMed:18644470, PubMed:26666690, PubMed:30247612, PubMed:14704337, PubMed:16397295, PubMed:26237645, PubMed:28712728, PubMed:29478807). Can phosphorylate C1D not only in the presence of linear DNA but also in the presence of supercoiled DNA (PubMed:9679063). Ability to phosphorylate p53/TP53 in the presence of supercoiled DNA is dependent on C1D (PubMed:9363941). Contributes to the determination of the circadian period length by antagonizing phosphorylation of CRY1 'Ser-588' and increasing CRY1 protein stability, most likely through an indirect mechanism (By similarity). Plays a role in the regulation of DNA virus-mediated innate immune response by assembling into the HDP-RNP complex, a complex that serves as a platform for IRF3 phosphorylation and subsequent innate immune response activation through the cGAS-STING pathway (PubMed:28712728). Also regulates the cGAS-STING pathway by catalyzing phosphorylation of CGAS, thereby impairing CGAS oligomerization and activation (PubMed:33273464). Also regulates the cGAS-STING pathway by mediating phosphorylation of PARP1 (PubMed:35460603).

Cellular Location

Nucleus. Nucleus, nucleolus

DNA PK (PRKDC) Antibody (N-term) Blocking peptide - Protocols

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

- [Blocking Peptides](#)

DNA PK (PRKDC) Antibody (N-term) Blocking peptide - Images

DNA PK (PRKDC) Antibody (N-term) Blocking peptide - Background

The PRKDC gene encodes the catalytic subunit of a nuclear DNA-dependent serine/threonine protein kinase (DNA-PK). The second component is the autoimmune antigen Ku (MIM 152690), which is encoded by the G22P1 gene on chromosome 22q. On its own, the catalytic subunit of DNA-PK is inactive and relies on the G22P1 component to direct it to the DNA and trigger its kinase activity; PRKDC must be bound to DNA to express its catalytic properties.[supplied by OMIM]

DNA PK (PRKDC) Antibody (N-term) Blocking peptide - References

Goudelock, D.M., et al., J. Biol. Chem. 278(32):29940-29947 (2003). Ding, Q., et al., Mol. Cell. Biol. 23(16):5836-5848 (2003). Lucero, H., et al., J. Biol. Chem. 278(24):22136-22143 (2003). Calsou, P., et al., J. Mol. Biol. 326(1):93-103 (2003). Karpova, A.Y., et al., Proc. Natl. Acad. Sci. U.S.A. 99(5):2818-2823 (2002).