

**PLK1 Antibody (C-term) Blocking Peptide**  
**Synthetic peptide**  
**Catalog # BP7937a****Specification****PLK1 Antibody (C-term) Blocking Peptide -  
Product Information**Primary Accession [P53350](#)**PLK1 Antibody (C-term) Blocking Peptide -  
Additional Information****Gene ID** 5347**Other Names**Serine/threonine-protein kinase PLK1,  
Polo-like kinase 1, PLK-1,  
Serine/threonine-protein kinase 13, STPK13,  
PLK1, PLK**Target/Specificity**

The synthetic peptide sequence used to generate the antibody [AP7937a](#) was selected from the C-term region of human PLK1 . 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.

**PLK1 Antibody (C-term) Blocking Peptide -  
Protein Information****Name** PLK1**PLK1 Antibody (C-term) Blocking Peptide -  
Background**

Protein kinases are enzymes that transfer a phosphate group from a phosphate donor, generally the  $\gamma$  phosphate of ATP, onto an acceptor amino acid in a substrate protein. By this basic mechanism, protein kinases mediate most of the signal transduction in eukaryotic cells, regulating cellular metabolism, transcription, cell cycle progression, cytoskeletal rearrangement and cell movement, apoptosis, and differentiation. With more than 500 gene products, the protein kinase family is one of the largest families of proteins in eukaryotes. The family has been classified in 8 major groups based on sequence comparison of their tyrosine (PTK) or serine/threonine (STK) kinase catalytic domains.

**PLK1 Antibody (C-term) Blocking Peptide -  
References**

Ree, A.H., et al., *Oncogene* 22(55):8952-8955 (2003). Cheng, K.Y., et al., *EMBO J.* 22(21):5757-5768 (2003). Elia, A.E., et al., *Cell* 115(1):83-95 (2003). Lin, H.R., et al., *J. Biol. Chem.* 278(38):35979-35987 (2003). Neef, R., et al., *J. Cell Biol.* 162(5):863-875 (2003).

## Synonyms PLK

### Function

Serine/threonine-protein kinase that performs several important functions throughout M phase of the cell cycle, including the regulation of centrosome maturation and spindle assembly, the removal of cohesins from chromosome arms, the inactivation of anaphase-promoting complex/cyclosome (APC/C) inhibitors, and the regulation of mitotic exit and cytokinesis. Polo-like kinase proteins acts by binding and phosphorylating proteins are that already phosphorylated on a specific motif recognized by the POLO box domains. Phosphorylates BORA, BUB1B/BUBR1, CCNB1, CDC25C, CEP55, ECT2, ERCC6L, FBXO5/EMI1, FOXM1, KIF20A/MKLP2, CENPU, NEDD1, NINL, NPM1, NUDC, PKMYT1/MYT1, KIZ, PPP1R12A/MYPT1, PRC1, RACGAP1/CYK4, SGO1, STAG2/SA2, TEX14, TOPORS, p73/TP73, TPT1, WEE1 and HNRNPU. Plays a key role in centrosome functions and the assembly of bipolar spindles by phosphorylating KIZ, NEDD1 and NINL. NEDD1 phosphorylation promotes subsequent targeting of the gamma-tubulin ring complex (gTuRC) to the centrosome, an important step for spindle formation. Phosphorylation of NINL component of the centrosome leads to NINL dissociation from other centrosomal proteins. Involved in mitosis exit and cytokinesis by phosphorylating CEP55, ECT2, KIF20A/MKLP2, CENPU, PRC1 and RACGAP1. Recruited at the central spindle by phosphorylating and docking PRC1 and KIF20A/MKLP2; creates its own docking sites on PRC1 and KIF20A/MKLP2 by mediating phosphorylation of sites subsequently recognized by the POLO box domains. Phosphorylates RACGAP1, thereby creating a docking site for the Rho GTP exchange factor ECT2 that is essential for the cleavage furrow formation. Promotes the central spindle recruitment of ECT2. Plays a central role in G2/M transition of mitotic cell cycle by phosphorylating CCNB1, CDC25C, FOXM1, CENPU, PKMYT1/MYT1, PPP1R12A/MYPT1 and WEE1. Part of a regulatory circuit that promotes the activation of CDK1 by phosphorylating the positive regulator CDC25C and inhibiting the negative regulators WEE1 and PKMYT1/MYT1. Also acts by mediating

phosphorylation of cyclin-B1 (CCNB1) on centrosomes in prophase. Phosphorylates FOXM1, a key mitotic transcription regulator, leading to enhance FOXM1 transcriptional activity. Involved in kinetochore functions and sister chromatid cohesion by phosphorylating BUB1B/BUBR1, FBXO5/EMI1 and STAG2/SA2. PLK1 is high on non-attached kinetochores suggesting a role of PLK1 in kinetochore attachment or in spindle assembly checkpoint (SAC) regulation. Required for kinetochore localization of BUB1B. Regulates the dissociation of cohesin from chromosomes by phosphorylating cohesin subunits such as STAG2/SA2. Phosphorylates SGO1: required for spindle pole localization of isoform 3 of SGO1 and plays a role in regulating its centriole cohesion function. Mediates phosphorylation of FBXO5/EMI1, a negative regulator of the APC/C complex during prophase, leading to FBXO5/EMI1 ubiquitination and degradation by the proteasome. Acts as a negative regulator of p53 family members: phosphorylates TOPORS, leading to inhibit the sumoylation of p53/TP53 and simultaneously enhance the ubiquitination and subsequent degradation of p53/TP53. Phosphorylates the transactivation domain of the transcription factor p73/TP73, leading to inhibit p73/TP73-mediated transcriptional activation and pro-apoptotic functions. Phosphorylates BORA, and thereby promotes the degradation of BORA. Contributes to the regulation of AURKA function. Also required for recovery after DNA damage checkpoint and entry into mitosis. Phosphorylates MISP, leading to stabilization of cortical and astral microtubule attachments required for proper spindle positioning (PubMed:<a href="http://www.uniprot.org/citations/8991084" target="\_blank">8991084</a>, PubMed:<a href="http://www.uniprot.org/citations/11202906" target="\_blank">11202906</a>, PubMed:<a href="http://www.uniprot.org/citations/12207013" target="\_blank">12207013</a>, PubMed:<a href="http://www.uniprot.org/citations/12447691" target="\_blank">12447691</a>, PubMed:<a href="http://www.uniprot.org/citations/12524548" target="\_blank">12524548</a>, PubMed:<a href="http://www.uniprot.org/ci

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target="\_blank">23509069</a>). Together  
with MEIKIN, acts as a regulator of  
kinetochore function during meiosis I:  
required both for mono-orientation of  
kinetochores on sister chromosomes and  
protection of centromeric cohesin from  
separase- mediated cleavage (By  
similarity). Phosphorylates CEP68 and is  
required for its degradation (PubMed:<a href="http://www.uniprot.org/citations/25503564" target="\_blank">25503564</a>).  
Regulates nuclear envelope breakdown  
during prophase by phosphorylating DCTN1  
resulting in its localization in the nuclear  
envelope (PubMed:<a href="http://www.uniprot.org/citations/20679239" target="\_blank">20679239</a>).  
Phosphorylates the heat shock transcription  
factor HSF1, promoting HSF1 nuclear  
translocation upon heat shock (PubMed:<a href="http://www.uniprot.org/citations/15661742" target="\_blank">15661742</a>).  
Phosphorylates HSF1 also in the early  
mitotic period; this phosphorylation  
regulates HSF1 localization to the spindle  
pole, the recruitment of the SCF(BTRC)

ubiquitin ligase complex inducing HSF1 degradation, and hence mitotic progression (PubMed:<a href="http://www.uniprot.org/citations/18794143" target="\_blank">18794143</a>). Regulates mitotic progression by phosphorylating ROK2 (PubMed:<a href="http://www.uniprot.org/citations/21880710" target="\_blank">21880710</a>).

#### **Cellular Location**

Nucleus. Chromosome, centromere, kinetochore. Cytoplasm, cytoskeleton, microtubule organizing center, centrosome. Cytoplasm, cytoskeleton, spindle. Midbody  
Note=localization at the centrosome starts at the G1/S transition (PubMed:24018379). During early stages of mitosis, the phosphorylated form is detected on centrosomes and kinetochores. Localizes to the outer kinetochore. Presence of SGO1 and interaction with the phosphorylated form of BUB1 is required for the kinetochore localization. Localizes onto the central spindle by phosphorylating and docking at midzone proteins KIF20A/MKLP2 and PRC1. Colocalizes with FRY to separating centrosomes and spindle poles from prophase to metaphase in mitosis, but not in other stages of the cell cycle. Localization to the centrosome is required for S phase progression (PubMed:24018379) Colocalizes with HSF1 at the spindle poles during prometaphase (PubMed:18794143).

#### **Tissue Location**

Placenta and colon.

### **PLK1 Antibody (C-term) Blocking Peptide - Protocols**

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

- [Blocking Peptides](#)