

PINK1 (PARK6) Antibody (N-term T133) Blocking peptide
Synthetic peptide
Catalog # BP6406a**Specification****PINK1 (PARK6) Antibody (N-term T133) Blocking peptide - Product Information**

Primary Accession [Q9BXM7](#)
Other Accession [NP_115785](#)

PINK1 (PARK6) Antibody (N-term T133) Blocking peptide - Additional Information

Gene ID 65018

Other Names

Serine/threonine-protein kinase PINK1,
mitochondrial, BRPK, PTEN-induced putative
kinase protein 1, PINK1

Target/Specificity

The synthetic peptide sequence used to generate the antibody AP6406a was selected from the PINK1 region of human PARK6 (PINK1). 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.

PINK1 (PARK6) Antibody (N-term T133) Blocking peptide - Protein Information

Name PINK1

PINK1 (PARK6) Antibody (N-term T133) Blocking peptide - Background

Defects in PINK1 are the cause of autosomal recessive early-onset Parkinson's disease 6 (PARK6). Six novel pathogenic PINK1 mutations suggest that PINK1 may be the second most common causative gene next to parkin in parkinsonism with the recessive mode of inheritance. Strong evidence indicates that, although important in mendelian forms of Parkinson's disease (PD), PINK1 does not influence the cause of sporadic nonmendelian forms of PD.

PINK1 (PARK6) Antibody (N-term T133) Blocking peptide - References

Rogaeva,E.,et al. Arch. Neurol. 61 (12), 1898-1904 (2004)Hatano,Y., et al. Ann. Neurol. 56 (3), 424-427 (2004)Healy,D.G., et al. Ann. Neurol. 56 (3), 329-335 (2004)Valente,E.M., et al. Science 304 (5674), 1158-1160 (2004)Nakajima,A., et al. Cancer Lett. 201 (2), 195-201 (2003)Unoki,M. and Nakamura,Y. Oncogene 20 (33), 4457-4465 (2001)

Serine/threonine-protein kinase which protects against mitochondrial dysfunction during cellular stress by phosphorylating mitochondrial proteins such as PRKN and DNMT1L, to coordinate mitochondrial quality control mechanisms that remove and replace dysfunctional mitochondrial components ([PubMed](#):<http://www.uniprot.org/citations/14607334>[target="_blank">14607334](#), PubMed:<http://www.uniprot.org/citations/18957282>[target="_blank">18957282](#), PubMed:<http://www.uniprot.org/citations/18443288>[target="_blank">18443288](#), PubMed:<http://www.uniprot.org/citations/15087508>[target="_blank">15087508](#), PubMed:<http://www.uniprot.org/citations/19229105>[target="_blank">19229105](#), PubMed:<http://www.uniprot.org/citations/19966284>[target="_blank">19966284](#), PubMed:<http://www.uniprot.org/citations/20404107>[target="_blank">20404107](#), PubMed:<http://www.uniprot.org/citations/22396657>[target="_blank">22396657](#), PubMed:<http://www.uniprot.org/citations/20798600>[target="_blank">20798600](#), PubMed:<http://www.uniprot.org/citations/23620051>[target="_blank">23620051](#), PubMed:<http://www.uniprot.org/citations/23754282>[target="_blank">23754282](#), PubMed:<http://www.uniprot.org/citations/23933751>[target="_blank">23933751](#), PubMed:<http://www.uniprot.org/citations/24660806>[target="_blank">24660806](#), PubMed:<http://www.uniprot.org/citations/24898855>[target="_blank">24898855](#), PubMed:<http://www.uniprot.org/citations/24751536>[target="_blank">24751536](#), PubMed:<http://www.uniprot.org/citations/24784582>[target="_blank">24784582](#)

tations/24896179"
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tations/32484300"
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PubMed:<a href="http://www.uniprot.org/ci
tations/20547144"
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Depending on the severity of mitochondrial
damage and/or dysfunction, activity ranges
from preventing apoptosis and stimulating
mitochondrial biogenesis to regulating
mitochondrial dynamics and eliminating
severely damaged mitochondria via
mitophagy (PubMed:<a href="http://www.u
niprot.org/citations/18443288"
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PubMed:<a href="http://www.uniprot.org/ci
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tations/32047033"
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tations/15087508"
target="_blank">15087508). Mediates
the translocation and activation of PRKN at
the outer membrane (OMM) of
dysfunctional/depolarized mitochondria
(PubMed:<a href="http://www.uniprot.org/c
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target="_blank">19966284,
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tations/25527291"
target="_blank">25527291). At the
OMM of damaged mitochondria,
phosphorylates pre-existing polyubiquitin
chains at 'Ser-65', the
PINK1-phosphorylated polyubiquitin then
recruits PRKN from the cytosol to the OMM
where PRKN is fully activated by
phosphorylation at 'Ser-65' by PINK1
(PubMed:<a href="http://www.uniprot.org/c
itations/19966284"
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PubMed:<a href="http://www.uniprot.org/ci
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tations/25474007"
target="_blank">25474007,
PubMed:<a href="http://www.uniprot.org/ci
tations/25527291"
target="_blank">25527291). In
damaged mitochondria, mediates the
decision between mitophagy or preventing
apoptosis by promoting PRKN-dependent

poly- or monoubiquitination of VDAC1; polyubiquitination of VDAC1 by PRKN promotes mitophagy, while monoubiquitination of VDAC1 by PRKN decreases mitochondrial calcium influx which ultimately inhibits apoptosis (PubMed:32047033). When cellular stress results in irreversible mitochondrial damage, functions with PRKN to promote clearance of damaged mitochondria via selective autophagy (mitophagy) (PubMed:14607334, PubMed:20798600, PubMed:20404107, PubMed:19966284, PubMed:23933751, PubMed:15087508). The PINK1-PRKN pathway also promotes fission of damaged mitochondria by phosphorylating and thus promoting the PRKN-dependent degradation of mitochondrial proteins involved in fission such as MFN2 (PubMed:18443288, PubMed:23620051, PubMed:24898855). This prevents the refusion of unhealthy mitochondria with the mitochondrial network or initiates mitochondrial fragmentation facilitating their later engulfment by autophagosomes (PubMed:18443288, PubMed:23620051, PubMed:23620051). Also promotes mitochondrial fission

independently of PRKN and ATG7-mediated mitophagy, via the phosphorylation and activation of DNM1L (PubMed:18443288, PubMed:32484300). Regulates motility of damaged mitochondria by promoting the ubiquitination and subsequent degradation of MIRO1 and MIRO2; in motor neurons, this likely inhibits mitochondrial intracellular anterograde transport along the axons which probably increases the chance of the mitochondria undergoing mitophagy in the soma (PubMed:22396657). Required for ubiquinone reduction by mitochondrial complex I by mediating phosphorylation of complex I subunit NDUFA10 (By similarity).

Cellular Location

Mitochondrion outer membrane; Single-pass membrane protein. Mitochondrion inner membrane {ECO:0000250|UniProtKB:Q99MQ3}; Single-pass membrane protein. Cytoplasm, cytosol. Note=Localizes mostly in mitochondrion and the two smaller proteolytic processed fragments localize mainly in cytosol (PubMed:19229105). When mitochondria lose mitochondrial membrane potential following damage, PINK1 import is arrested, which induces its accumulation in the outer mitochondrial membrane, where it acquires kinase activity (PubMed:18957282)

Tissue Location

Highly expressed in heart, skeletal muscle and testis, and at lower levels in brain, placenta, liver, kidney, pancreas, prostate, ovary and small intestine. Present in the embryonic testis from an early stage of development

PINK1 (PARK6) Antibody (N-term T133) Blocking peptide - Protocols

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

- [Blocking Peptides](#)