

Park6(PINK1) Antibody(C-term)
Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP6406D

Specification

Park6(PINK1) Antibody(C-term) - Product Information

Application	WB,E
Primary Accession	Q9BXM7
Reactivity	Human, Mouse
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Antigen Region	493-526

Park6(PINK1) Antibody(C-term) - Additional Information

Gene ID 65018

Other Names

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

Target/Specificity

This Park6(PINK1) antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 493-526 amino acids from the C-terminal region of human Park6(PINK1).

Dilution

WB~~1:1000

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

Park6(PINK1) Antibody(C-term) is for research use only and not for use in diagnostic or therapeutic procedures.

Park6(PINK1) Antibody(C-term) - Protein Information

Name PINK1

Function Serine/threonine-protein kinase which protects against mitochondrial dysfunction during cellular stress by phosphorylating mitochondrial proteins such as PRKN and DNM1L, to coordinate mitochondrial quality control mechanisms that remove and replace dysfunctional mitochondrial

components (PubMed:[14607334](#), PubMed:[18957282](#), PubMed:[18443288](#), PubMed:[15087508](#), PubMed:[19229105](#), PubMed:[19966284](#), PubMed:[20404107](#), PubMed:[22396657](#), PubMed:[20798600](#), PubMed:[23620051](#), PubMed:[23754282](#), PubMed:[23933751](#), PubMed:[24660806](#), PubMed:[24898855](#), PubMed:[24751536](#), PubMed:[24784582](#), PubMed:[24896179](#), PubMed:[25527291](#), PubMed:[32484300](#), PubMed:[20547144](#)). 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:[18443288](#), PubMed:[23620051](#), PubMed:[24898855](#), PubMed:[20798600](#), PubMed:[20404107](#), PubMed:[19966284](#), PubMed:[32484300](#), PubMed:[22396657](#), PubMed:[32047033](#), PubMed:[15087508](#)). Mediates the translocation and activation of PRKN at the outer membrane (OMM) of dysfunctional/depolarized mitochondria (PubMed:[19966284](#), PubMed:[20404107](#), PubMed:[20798600](#), PubMed:[23754282](#), PubMed:[24660806](#), PubMed:[24751536](#), PubMed:[24784582](#), PubMed:[25474007](#), PubMed:[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:[19966284](#), PubMed:[20404107](#), PubMed:[20798600](#), PubMed:[23754282](#), PubMed:[24660806](#), PubMed:[24751536](#), PubMed:[24784582](#), PubMed:[25474007](#), PubMed:[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](#)). Also promotes mitochondrial fission independently of PRKN and ATG7-mediated mitophagy, via the phosphorylation and activation of DNMT1 (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 NDUF10 (By similarity). Phosphorylates LETM1, positively regulating its mitochondrial calcium transport activity (PubMed:[29123128](#)).

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

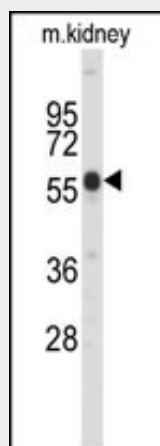
Highly expressed in heart, skeletal muscle and testis, and at lower levels in brain, placenta, liver, kidney, pancreas, prostate, ovary and small intestine. Present in the embryonic testis from an early stage of development

Park6(PINK1) Antibody(C-term) - Protocols

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

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

Park6(PINK1) Antibody(C-term) - Images



Western blot analysis of Park6 (PINK1) C-term (Cat. #AP6406d) in mouse kidney tissue lysates (35ug/lane). Park6 (arrow) was detected using the purified Pab.

Park6(PINK1) Antibody(C-term) - Background

Parkinson is the second most common neurodegenerative disease after Alzheimers. About 1 percent of people over the age of 65 and 3 percent of people over the age of 75 are affected by the disease. The mutation is the most common cause of Parkinson disease identified to date. 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.

Park6(PINK1) Antibody(C-term) - References

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., et al., Oncogene 20(33):4457-4465 (2001).

Park6(PINK1) Antibody(C-term) - Citations

- [CLEC16A regulates splenocyte and NK cell function in part through MEK signaling.](#)
- [PINK1 regulates histone H3 trimethylation and gene expression by interaction with the polycomb protein EED/WAIT1.](#)