

## **Anti-PINK1** antibody



**Description** Unconjugated Rabbit polyclonal to PINK1

Model STJ192141

**Host** Rabbit

**Reactivity** Human, Mouse

**Applications** ELISA, WB

**Gene ID** <u>65018</u>

Gene Symbol PINK1

**Dilution range** WB 1:500-2000 ELISA 1:5000-20000

**Specificity** PINK1 Polyclonal Antibody detects endogenous levels of protein.

**Tissue Specificity** 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.

**Purification** PINK1 antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

**Note** For Research Use Only (RUO).

Protein Name Serine/threonine-protein kinase PINK1, mitochondrial BRPK PTEN-induced

putative kinase protein 1

Molecular Weight 63 kDa

**Clonality** Polyclonal

**Conjugation** Unconjugated

Isotype IgG

Liquid form in PBS containing 50% glycerol, and 0.02% sodium azide. **Formulation** 

Concentration 1 mg/ml

Store at -20°C, and avoid repeat freeze-thaw cycles. **Storage Instruction** 

**Database Links** HGNC:14581OMIM:168600

**Alternative Names** Serine/threonine-protein kinase PINK1, mitochondrial BRPK PTEN-induced

putative kinase protein 1

**Function** Protects against mitochondrial dysfunction during cellular stress by

> phosphorylating mitochondrial proteins. Involved in the clearance of damaged mitochondria via selective autophagy (mitophagy) by mediating activation and translocation of PRKN. Targets PRKN to dysfunctional depolarized mitochondria through the phosphorylation of MFN2. Activates PRKN in 2 steps: (1) by mediating phosphorylation at 'Ser-65' of PRKN and (2) mediating phosphorylation of ubiquitin, converting PRKN to its fully-active

form.

**Cellular Localization** Mitochondrion outer membrane. Single-pass membrane protein Cytoplasm,

cytosol. Localizes mostly in mitochondrion and the 2 proteolytic processed

fragments of 55 kDa and 48 kDa localize mainly in cytosol.

Post-translational

Autophosphorylation at Ser-228 and Ser-402 is essential for Parkin/PRKN recruitment to depolarized mitochondria. Two shorter forms of 55 kDa and 48 **Modifications** 

kDa seem to be produced by proteolytic cleavage and localize mainly in

cytosol.

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