

**PNKD Antibody (N-term) Blocking peptide**  
**Synthetic peptide**  
**Catalog # BP12567a****Specification**

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**PNKD Antibody (N-term) Blocking peptide - Product Information**Primary Accession [Q8N490](#)**PNKD Antibody (N-term) Blocking peptide - Additional Information****Gene ID** 25953**Other Names**

Probable hydrolase PNKD, 3---, Myofibrillogenesis regulator 1, MR-1, Paroxysmal nonkinesio-genic dyskinesia protein, Trans-activated by hepatitis C virus core protein 2, PNKD, KIAA1184, MR1, TAHCCP2

**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.

**PNKD Antibody (N-term) Blocking peptide - Protein Information****Name** PNKD**Synonyms** KIAA1184, MR1, TAHCCP2**Function**

Probable thioesterase that may play a role in cellular detoxification processes; it likely acts on a yet-unknown alpha- hydroxythioester substrate (Probable). In vitro, it is able to catalyze the hydrolysis of S-D-lactoyl-glutathione to form glutathione and D- lactic acid at very low rate, though this reaction is not physiologically relevant in vivo (PubMed:<a href="http://www.uniprot.org/citations/21487022" target="\_blank">21487022</a>).

**Cellular Location**

[Isoform 1]: Cell membrane; Peripheral membrane protein. Mitochondrion [Isoform 3]: Mitochondrion. Golgi apparatus. Endoplasmic reticulum

**Tissue Location**

Isoform 1 is only expressed in the brain. Isoform 2 is ubiquitously detected with highest expression in skeletal muscle and detected in myocardial myofibrils.

**PNKD Antibody (N-term) Blocking peptide - Protocols**

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

- [Blocking Peptides](#)

**PNKD Antibody (N-term) Blocking peptide - Images****PNKD Antibody (N-term) Blocking peptide - Background**

This gene is thought to play a role in the regulation of myofibrillogenesis. Mutations in this gene have been associated with the movement disorder paroxysmal non-kinesigenic dyskinesia. Alternative splicing results in multiple transcript variants.

**PNKD Antibody (N-term) Blocking peptide - References**

Yang, J., et al. Neuron 66(6):871-883(2010) Davila, S., et al. Genes Immun. 11(3):232-238(2010) Ghezzi, D., et al. Hum. Mol. Genet. 18(6):1058-1064(2009) Friedman, A., et al. Eur. Neurol. 61(1):39-41(2009) Ren, K., et al. J. Biol. Chem. 283(51):35598-35605(2008)