

## DMPK rabbit pAb

Cat No.: ES2180

For research use only

## Overview

Product Name DMPK rabbit pAb

Host species Rabbit
Applications WB;ELISA

**Species Cross-Reactivity** Human;Rat;Mouse;

Recommended dilutions Western Blot: 1/500 - 1/2000. ELISA: 1/40000. Not

yet tested in other applications.

Immunogen The antiserum was produced against synthesized

peptide derived from human DMPK. AA range:11-60

**Specificity** DMPK Polyclonal Antibody detects endogenous

levels of DMPK protein.

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and

0.02% sodium azide.

**Storage** Store at -20°C. Avoid repeated freeze-thaw cycles.

**Protein Name** Myotonin-protein kinase

Gene Name DMPK

**Cellular localization** Endoplasmic reticulum membrane; Single-pass type

IV membrane protein; Cytoplasmic side. Nucleus outer membrane; Single-pass type IV membrane protein; Cytoplasmic side. Mitochondrion outer membrane; Single-pass type IV membrane protein.

Sarcoplasmic reticulum membrane . Cell membrane . Cytoplasm, cytosol . Localizes to

sarcoplasmic reticulum membranes of

cardiomyocytes. .; [Isoform 1]: Mitochondrion membrane.; [Isoform 3]: Mitochondrion membrane.

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**Purification** The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

Clonality Polyclonal
Concentration 1 mg/ml
Observed band 70kD
Human Gene ID 1760
Human Swiss-Prot Number Q09013



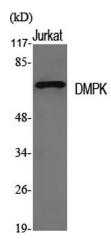
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**Alternative Names** 

**Background** 

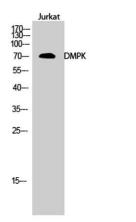
DMPK; DM1PK; MDPK; Myotonin-protein kinase; MT-PK; DM-kinase; DMK; DM1 protein kinase; DMPK; Myotonic dystrophy protein kinase The protein encoded by this gene is a serine-threonine kinase that is closely related to other kinases that interact with members of the Rho family of small GTPases. Substrates for this enzyme include myogenin, the beta-subunit of the L-type calcium channels, and phospholemman. The 3' untranslated region of this gene contains 5-38 copies of a CTG trinucleotide repeat. Expansion of this unstable motif to 50-5,000 copies causes myotonic dystrophy type I, which increases in severity with increasing repeat element copy number. Repeat expansion is associated with condensation of local chromatin structure that disrupts the expression of genes in this region. Several alternatively spliced transcript variants of this gene have been described, but the full-length nature of some of these variants has not been determined. [provided by RefSeq, Jul 2016],



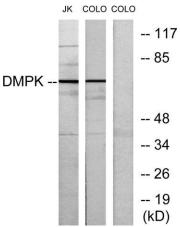
Western Blot analysis of various cells using DMPK Polyclonal Antibody



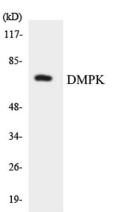




Western Blot analysis of Jurkat cells using DMPK Polyclonal Antibody



Western blot analysis of lysates from Jurkat and COLO205 cells, using DMPK Antibody. The lane on the right is blocked with the synthesized peptide.



Western blot analysis of the lysates from HT-29 cells using DMPK antibody.

