# abcam

### Product datasheet

# Anti-Dystrophia myotonica protein kinase / DMPK antibody ab102804

## 2 Images

#### Overview

Product name Anti-Dystrophia myotonica protein kinase / DMPK antibody

**Description**Rabbit polyclonal to Dystrophia myotonica protein kinase / DMPK

Host species Rabbit

Tested applications Suitable for: WB

Species reactivity Reacts with: Human

Immunogen Recombinant full length protein corresponding to Human Dystrophia myotonica protein kinase/

DMPK aa 1-629.

Database link: NP 004400.4

Positive control WB: HeLa and Dystrophia myotonica protein kinase / DMPK transfected 293T cell lysates.

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

#### **Properties**

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

Storage buffer pH: 7.40

Constituent: 100% PBS

Purity Protein A purified

**Clonality** Polyclonal

**Isotype** IgG

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab102804 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		1/500 - 1/1000. Predicted molecular weight: 70 kDa.

#### **Target**

**Function** 

Critical to the modulation of cardiac contractility and to the maintenance of proper cardiac conduction activity. Phosphorylates phospholamban.

**Tissue specificity** 

Most isoforms are expressed in many tissues including heart, skeletal muscle, liver and brain, except for isoform 2 which is only found in the heart and skeletal muscle, and isoform 14 which is only found in the brain, with high levels in the striatum, cerebellar cortex and pons.

Involvement in disease

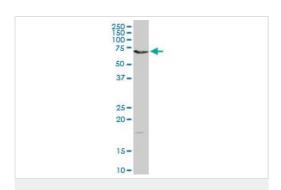
Defects in DMPK are the cause of dystrophia myotonica type 1 (DM1) [MIM:160900]; also known as Steinert disease. A muscular disorder characterized by myotonia, muscle wasting in the distal extremities, cataract, hypogonadism, defective endocrine functions, male baldness and cardiac arrhythmias. Note=The causative mutation is a CTG expansion in the 3'-UTR of the DMPK gene. A length exceeding 50 CTG repeats is pathogenic, while normal individuals have 5 to 37 repeats. Intermediate alleles with 35-49 triplets are not disease-causing but show instability in intergenerational transmissions. Disease severity varies with the number of repeats: mildly affected persons have 50 to 150 repeats, patients with classic DM have 100 to 1,000 repeats, and those with congenital onset can have more than 2,000 repeats.

Sequence similarities

Belongs to the protein kinase superfamily. AGC Ser/Thr protein kinase family. DMPK subfamily. Contains 1 AGC-kinase C-terminal domain.

Contains 1 protein kinase domain.

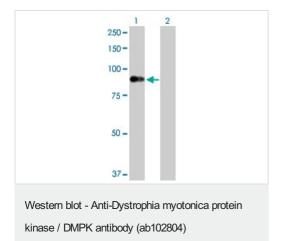
#### **Images**



Western blot - Anti-Dystrophia myotonica protein kinase / DMPK antibody (ab102804)

Anti-Dystrophia myotonica protein kinase / DMPK antibody (ab102804) at 1/500 dilution + HeLa cell lysate at 50 µg

Predicted band size: 70 kDa



**All lanes :** Anti-Dystrophia myotonica protein kinase / DMPK antibody (ab102804) at 1/500 dilution

**Lane 1 :** Dystrophia myotonica protein kinase / DMPK transfected 293T cell lysate

Lane 2: Non-transfected 293T cell line

Lysates/proteins at 25 µg per lane.

Predicted band size: 70 kDa

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

#### Our Abpromise to you: Quality guaranteed and expert technical support

- · Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- · Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit <a href="https://www.abcam.com/abpromise">https://www.abcam.com/abpromise</a> or contact our technical team.

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