abcam

Product datasheet

Anti-MTPAP antibody ab156684

2 Images

Overview

Product name Anti-MTPAP antibody

Description Rabbit polyclonal to MTPAP

Host species Rabbit

Tested applications

Suitable for: WB, IP

Species reactivity

Reacts with: Human

Predicted to work with: Chimpanzee, Gorilla, Orangutan

Immunogen Synthetic peptide corresponding to Human MTPAP aa 532-582.

Database link: **NP_060579.3**

Positive control HeLa, 293T and Jurkat whole 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.

Storage buffer pH: 7

Preservative: 0.09% Sodium azide Constituent: 99% Tris citrate/phosphate

pH 7 to 8

Purity Immunogen affinity purified

Purification notes ab156684 was affinity purified using an epitope specific to MTPAP, mitochondrial immobilized on

solid support.

Clonality Polyclonal

1

Isotype IgG

Applications

The Abpromise guarantee

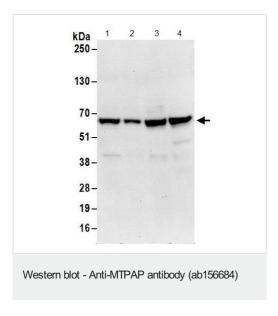
Our <u>Abpromise guarantee</u> covers the use of ab156684 in the following tested applications.

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

Application	Abreviews	Notes
WB		1/2000 - 1/10000. Predicted molecular weight: 66 kDa.
IP		Use at 2-10 μg/mg of lysate.

Target		
Function	Polymerase that creates the 3' poly(A) tail of mitochondrial transcripts. Can use all four nucleotides, but has higher activity with ATP and UTP (in vitro). Plays a role in replication-dependent histone mRNA degradation. May be involved in the terminal uridylation of mature histone mRNAs before their degradation is initiated. Might be responsible for the creation of some UAA stop codons which are not encoded in mtDNA.	
Tissue specificity	Ubiquitous, with stronger expression in tissues with high energy requirements: heart, brain, and skeletal muscle.	
Involvement in disease	Defects in MTPAP are the cause of spastic ataxia autosomal recessive type 4 (SPAX4) [MIM:613672]. A slowly progressive neurodegenerative disease characterized by cerebellar ataxia, spastic paraparesis, dysarthria, and optic atrophy. Note=Affected individuals exhibit a drastic decrease in poly(A) tail length of representative mitochondrial mRNA transcripts, including COX1 and RNA14 (PubMed:20970105).	
Sequence similarities	Belongs to the DNA polymerase type-B-like family. Contains 1 PAP-associated domain.	
Cellular localization	Cytoplasm. Mitochondrion.	

Images





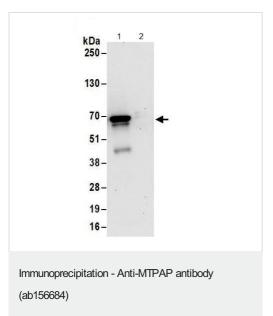
Lane 1 : HeLa whole cell lysate at 50 μg Lane 2 : HeLa whole cell lysate at 15 μg Lane 3 : 293T whole cell lysate at 50 μg

Lane 4: Jurkat whole cell lysate at 50 µg

Developed using the ECL technique.

Predicted band size: 66 kDa

Exposure time: 30 seconds



Detection of MTPAP, mitochondrial by Western blot of Immunoprecipitate:

Immunoprecipitation of HeLa whole cell lysate using ab156684 at 6 μ g/mg lysate (1 mg/IP; 20% of IP loaded/lane) followed by detection of MTPAP, mitochondrial with ab156684 at 1 μ g/ml. The lysate in lane 2 is precipitated with an IgG control antibody.

Detection: Chemiluminescence with exposure time of 30 seconds.

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

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