

Product datasheet

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ARG45604 anti-AMPD1 antibody

Package: 50 μg Store at: -20°C

Summary

Product Description Rabbit Polyclonal antibody recognizes AMPD1

Tested Reactivity Hu, Ms, Rat

Tested Application FACS, IHC-P, WB

Specificity AMPD1
Host Rabbit

Clonality Polyclonal

Isotype IgG

Target Name AMPD1
Species Human

Immunogen Recombinant protein containing to human AMPD1.

Conjugation Un-conjugated

Alternate Names AMPD1; Adenosine Monophosphate Deaminase 1; Myoadenylate Deaminase; MADA; MAD; Adenosine

Monophosphate Deaminase 1 (Isoform M); Skeletal Muscle AMPD; AMP Deaminase 1; EC 3.5.4.6; Adenosine Monophosphate Deaminase-1 (Muscle); AMP Deaminase Isoform M; AMPD Isoform M;

MMDD; AMPD

Application Instructions

Application table	Application	Dilution
	FACS	1 - 3 μg/10^6 cells
	IHC-P	2-5 μg/ml
	WB	0.25-0.5 μg/ml
Application Note	* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form Powder

Purification Affinity purified

Buffer 0.2% Na2HPO4, 0.9% NaCl and 4% Trehalose.

Stabilizer 4% Trehalose

Concentration 0.5 mg/ml

Storage instruction For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot

and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed

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before use.

Note

For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Gene Symbol AMPD1

Gene Full Name Adenosine Monophosphate Deaminase 1

Background Adenosine monophosphate deaminase 1 catalyzes the deamination of AMP to IMP in skeletal muscle

and plays an important role in the purine nucleotide cycle. Two other genes have been identified, AMPD2 and AMPD3, for the liver- and erythocyte-specific isoforms, respectively. Deficiency of the muscle-specific enzyme is apparently a common cause of exercise-induced myopathy and probably the most common cause of metabolic myopathy in the human. Alternatively spliced transcript variants encoding different isoforms have been identified in this gene.[provided by RefSeq, Feb 2010]

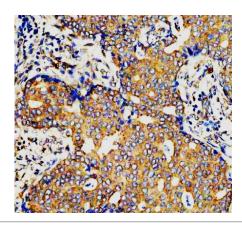
Function AMP deaminase plays a critical role in energy metabolism. [UniProt]

Calculated Mw 63 kDa

PTM Phosphoprotein. [UniProt]

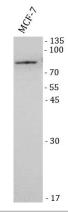
Cellular Localization Cytosol. [UniProt]

Images



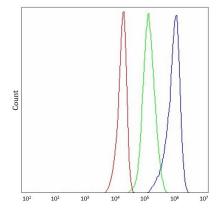
ARG45604 anti-AMPD1 antibody IHC-P image

Immunohistochemistry: Human breast cancer stained with ARG45604 anti-AMPD1 antibody at 2 $\mu g/ml$ dilution.



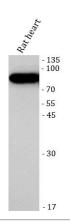
ARG45604 anti-AMPD1 antibody WB image

Western blot: MCF-7 stained with ARG45604 anti-AMPD1 antibody at 0.5 $\mu\text{g}/\text{ml}$ dilution.



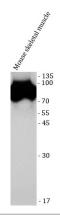
ARG45604 anti-AMPD1 antibody FACS image

Flow Cytometry: U251 stained with ARG45604 anti-AMPD1 antibody at 1 $\mu g/10^{\circ}6$ cells dilution.



ARG45604 anti-AMPD1 antibody WB image

Western blot: Rat heart stained with ARG45604 anti-AMPD1 antibody at 0.5 $\mu\text{g}/\text{ml}$ dilution.



ARG45604 anti-AMPD1 antibody WB image

Western blot: Mouse skeletal muscle stained with ARG45604 anti-AMPD1 antibody at 0.5 $\mu g/ml$ dilution.