

Product datasheet

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ARG42606 anti-PFKM antibody

Package: 50 μl Store at: -20°C

Summary

Product Description Recombinant Rabbit Monoclonal antibody recognizes PFKM

Tested Reactivity Hu, Ms, Rat

Tested Application FACS, ICC/IF, IHC-P, WB

Host Rabbit

Clonality Monoclonal

Isotype IgG

Target Name PFKM

Species Human

Immunogen Recombinant protein of Human PFKM.

Conjugation Un-conjugated

Alternate Names PFK-A; 6-phosphofructokinase type A; PPP1R122; PFKX; ATP-dependent 6-phosphofructokinase, muscle

type; EC 2.7.1.11; Phosphofructo-1-kinase isozyme A; PFK1; ATP-PFK; GSD7; PFK-1; Phosphohexokinase;

PFK-M; PFKA

Application Instructions

Application table	Application	Dilution
	FACS	1:50 - 1:100
	ICC/IF	1:50 - 1:200
	IHC-P	1:50 - 1:200
	WB	1:500 - 1:1000
Application Note	* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	
Positive Control	PC-3M	
Observed Size	~ 82 kDa	

Properties

Form Liquid

Purification Purification with Protein A.

Buffer TBS (pH 7.4), 0.05% Sodium azide, 40% Glycerol and 0.05% BSA.

Preservative 0.05% Sodium azide

Stabilizer 40% Glycerol and 0.05% BSA

Concentration 1 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. 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 before use.

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

Bioinformation

Gene Symbol PFKM

Gene Full Name phosphofructokinase, muscle

Background Three phosphofructokinase isozymes exist in humans: muscle, liver and platelet. These isozymes

function as subunits of the mammalian tetramer phosphofructokinase, which catalyzes the phosphorylation of fructose-6-phosphate to fructose-1,6-bisphosphate. Tetramer composition varies depending on tissue type. This gene encodes the muscle-type isozyme. Mutations in this gene have been associated with glycogen storage disease type VII, also known as Tarui disease. Alternatively

spliced transcript variants have been described. [provided by RefSeq, Nov 2009]

Function Catalyzes the phosphorylation of D-fructose 6-phosphate to fructose 1,6-bisphosphate by ATP, the first

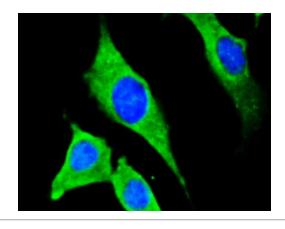
committing step of glycolysis. [UniProt]

Calculated Mw 85 kDa

PTM GlcNAcylation decreases enzyme activity. [UniProt]

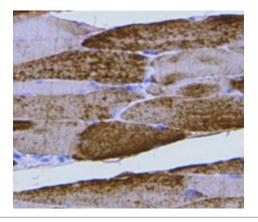
Cellular Localization Cytoplasm. [UniProt]

Images



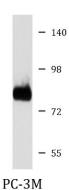
ARG42606 anti-PFKM antibody ICC/IF image

Immunofluorescence: SH-SY5Y cells were fixed in paraformaldehyde and permeabilized with 0.25% Triton X100/PBS. Cells were stained with ARG42606 anti-PFKM antibody (green). DAPI (blue) for nuclear staining.



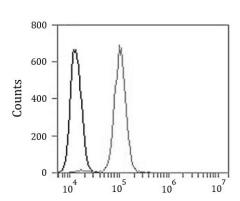
ARG42606 anti-PFKM antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Mouse skeletal muscle tissue stained with ARG42606 anti-PFKM antibody and counter stained with hematoxylin.



ARG42606 anti-PFKM antibody WB image

Western blot: PC-3M cell lysate stained with ARG42606 anti-PFKM antibody at 1:500 dilution.



ARG42606 anti-PFKM antibody FACS image

Flow Cytometry: SH-SY5Y cells stained with ARG42606 anti-PFKM antibody (right histogram) at 1:100 dilution or without primary antibody as control (left histogram), followed by incubation with Alexa Fluor® 488 labelled secondary antibody.