# abcam

## Product datasheet

# Recombinant Human PFKM protein ab95304

# 1 Image

#### **Description**

Product name Recombinant Human PFKM protein

Purity > 80 % SDS-PAGE.

Purified using conventional chromatography techniques.

Expression system Escherichia coli

Protein length Full length protein

Animal free No

Nature Recombinant

**Species** Human

Sequence

MGSSHHHHHH SSGLVPRGSH MTHEEHHAAK

TLGIGKAIAV LTSGGDAQGM NAAVRAVVRV GIFTGARVFF VHEGYQGLVD GGDHIKEATW ESVSMMLQLG GTVIGSARCK DFREREGRLR AAYNLVKRGI TNLCVIGGDG SLTGADTFRS

EWSDLLSDLQ KAGKITDEEA TKSSYLNIVG LVGSIDNDFC GTDMTIGTDS ALHRIMEIVD AITTTAQSHQ RTFVLEVMGR

HCGYLALVTS LSCGADWVFLPECPPDDDWE

EHLCRRLSET RTRGSRLNII IVAEGAIDKN GKPITSEDIK

NLVVKRLGYD TRVTVLGHVQ RGGTPSAFDR
ILGSRMGVEA VMALLEGTPD TPACVVSLSG
NQAVRLPLME CVQVTKDVTK AMDEKKFDEA
LKLRGRSFMN NWEVYKLLAH VRPPVSKSGS
HTVAVMNVGA PAAGMNAAVR STVRIGLIQG
NRVLVVHDGF EGLAKGQIEE AGWSYVGGWT

GQGGSKLGTK RTLPKKSFEQ ISANITKFNI QGLVIIGGFE

AYTGGLELME GRKQFDELCIPFVVIPATVS

NNVPGSDFSV GADTALNTIC TTCDRIKQSA AGTKRRVFII

ETMGGYCGYL ATMAGLAAGA DAAYIFEEPF

TIRDLQANVE HLVQKMKTTV KRGLVLRNEK CNENYTTDFI

FNLYSEEGKG IFDSRKNVLG HMQQGGSPTP FDRNFATKMG AKAMNWMSGK IKESYRNGRI FANTPDSGCV LGMRKRALVF QPVAELKDQT

DFEHRIPKEQ WWLKLRPILK ILAKYEIDLD TSDHAHLEHI

**TRKRSGEAAV** 

#### **Specifications**

Our Abpromise guarantee covers the use of ab95304 in the following tested applications.

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

Applications SDS-PAGE

Form Liquid

Additional notes This product was previously labelled as Fructose 6 Phosphate Kinase

#### **Preparation and Storage**

Stability and Storage Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

pH: 8.00

Constituents: 0.077% DTT, 0.316% Tris HCI, 20% Glycerol (glycerin, glycerine), 1.16% Sodium

chloride

#### **General Info**

Pathway Carbohydrate degradation; glycolysis; D-glyceraldehyde 3-phosphate and glycerone phosphate

from D-glucose: step 3/4.

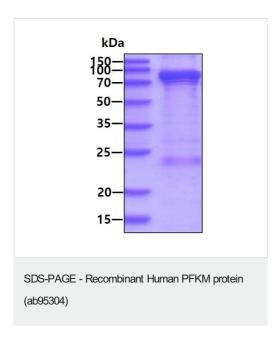
**Involvement in disease** Defects in PFKM are the cause of glycogen storage disease type 7 (GSD7) [MIM:232800]; also

known as Tarui disease. GSD7 is an autosomal recessive disorder characterized by exercise intolerance with associated nausea and vomiting. Short bursts of intense activity are particularly difficult. Severe muscle cramps and myoglobinuria develop after vigorous exercise. Most patients obtain a "second wind" when the onset of exercise is followed by a brief rest period. In time

patients adjust their activity level and are well compensated.

Sequence similarities Belongs to the phosphofructokinase family. Two domains subfamily.

# **Images**



SDS-PAGE analysis of ab95304 (3 µg) under reducing conditions and visualized by coomassie blue stain.

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

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