

Recombinant Human PCK1/PEPC protein ab119469

1 图像

描述

产品名称 重组人PCK1/PEPC蛋白

纯度 > 90 % SDS-PAGE.
ab119469 was purified using conventional chromatography.

表达系统 Escherichia coli

Accession [P35558](#)

蛋白长度 Full length protein

无动物成分 No

性质 Recombinant

种属 Human

序列

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MGSSHHHHH SSGLVPRGSH MGSHPMPQLQ
NGLNLSAKVV QGSLDSLPA VREFLENAE
LCQPDHIHIC DGSEENGRL LGQMEEEGIL
RRLKKYDNCW LALDPRDVA RIESKTVIVT
QEQRDTVPIP KTGLSQLGRW MSEEDFEKAF
NARFPGCMKG RTMYVIPFSM GPLGSPLSKI
GIELTDSYV VASMRIMTRM GTPVLEALGD
GEFVKCLHSV GCPLPLQKPL VNNWPCNPEL
TLIAHLPDRR EIISFGSGYG GNSLLGKKCF
ALRMASRLAK EEGWLAEHML VLGITNPEGE
KKYLAAAFPS ACGKTNLMM NPSLPGWKVE
CVGDDIAWMK FDAQGHLRAI NPENFFGVA
PGTSVKTNPN AIKTIQKNTI FTNVAETSDG
GVYWEGIDEP LASGVTITSW KNKEWSSDGE
EPCAHNSRF CTPASQCPII DAAWESPEGV
PIEGIIFGGR RPAGVPLVYE ALSWQHGVFV
GAAMRSEATA AAHKGKIIM HDPFAMRPF
GYNFGKYLAH WLSMAQPAA KLPKIFHVNW
FRKDKGKFL WPGFGNSRV LEWMFNRIDG
KASTKLTPIG YIPKEDALNL KGLGHINMME
LFSISKEFWE KEVEDIEKYL EDQVNADLPC
EIEREILALK QRISQM
    
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预测分子量 72 kDa including tags

氨基酸 1 to 622

标签 His tag N-Terminus

技术指标

Our **Abpromise guarantee** covers the use of **ab119469** in the following tested applications.

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

应用	SDS-PAGE
形式	Liquid
补充说明	This product was previously labelled as PCK1

制备和贮存

稳定性和存储 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long term. Avoid freeze / thaw cycle.

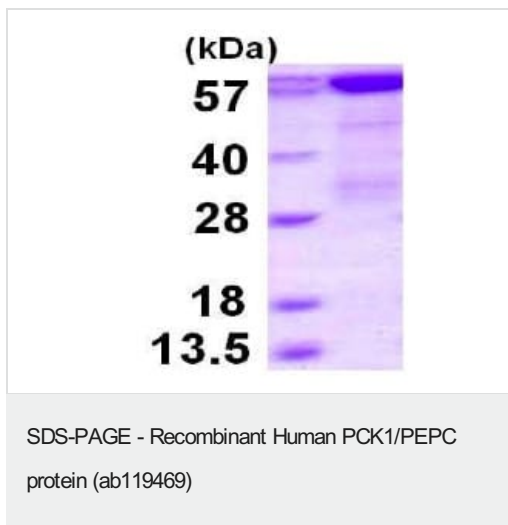
pH: 8.00

Constituents: 0.02% DTT, 0.32% Tris HCl, 10% Glycerol (glycerin, glycerine), 0.58% Sodium chloride

常规信息

功能	Catalyzes the conversion of oxaloacetate (OAA) to phosphoenolpyruvate (PEP), the rate-limiting step in the metabolic pathway that produces glucose from lactate and other precursors derived from the citric acid cycle.
组织特异性	Major sites of expression are liver, kidney and adipocytes.
通路	Carbohydrate biosynthesis; gluconeogenesis.
疾病相关	Defects in PCK1 are the cause of cytosolic phosphoenolpyruvate carboxykinase deficiency (cytosolic PEPCK deficiency) [MIM:261680]. PEPCK deficiency is a metabolic disorder resulting from impaired gluconeogenesis. It is a rare disease with less than 10 cases reported in the literature. Clinical characteristics include hypotonia, hepatomegaly, failure to thrive, lactic acidosis and hypoglycemia. Autopsy reveals fatty infiltration of both the liver and kidneys. The disorder is transmitted as an autosomal recessive trait.
序列相似性	Belongs to the phosphoenolpyruvate carboxykinase [GTP] family.
翻译后修饰	Acetylation is increased on addition of glucose and appears to regulate the protein stability.
细胞定位	Cytoplasm.

图片



15% SDS-PAGE analysis of ab119469 (3µg)

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

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