abcam

Product datasheet

Anti-PKLR antibody ab125697

3 图像

概述

产品名称 Anti-PKLR抗体

描述 兔多克隆抗体to PKLR

宿主 Rabbit

经测试应用 适用于: WB, IHC-P, ICC/IF

种属反应性 与反应: Human

免疫原 Recombinant fragment containing a sequence corresponding to a region within amino acids 178-

510 of Human PKLR.(Uniprot:P30613)

阳性对照 A549 cell lysate, HeLa whole cell lysate (ab150035), HepG2 whole cell lysate (ab7900), HTC116

cell lysate and Human breast cancer tissue.

常规说明

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

性能

形式 Liquid

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

cycles.

存储溶液 pH: 7.00

Preservative: 0.01% Thimerosal (merthiolate)

Constituents: 1.21% Tris, 0.75% Glycine, 20% Glycerol (glycerin, glycerine)

纯**度** Immunogen affinity purified

克隆 多克隆

同种型 IgG

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab125697于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应 用	Ab评论	说明
WB		1/500 - 1/3000. Predicted molecular weight: 62 kDa.
IHC-P		1/100 - 1/500. Perfrom antigen retrieval using heat mediated 10mM Citrate buffer(pH6.0) or Tris-EDTA buffer(pH8.0).
ICC/IF		1/100 - 1/500.

靶标

疾病相关

功能 Plays a key role in glycolysis.

通路 Carbohydrate degradation; glycolysis; pyruvate from D-glyceraldehyde 3-phosphate: step 5/5.

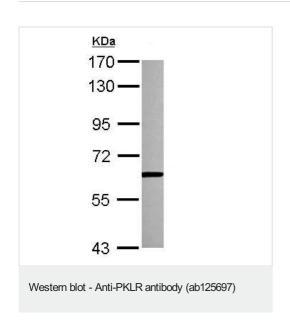
Defects in PKLR are the cause of pyruvate kinase hyperactivity (PKHYP) [MIM:102900]; also known as high red cell ATP syndrome. This autosomal dominant phenotype is characterized by increase of red blood cell ATP.

Defects in PKLR are the cause of pyruvate kinase deficiency of red cells (PKRD) [MIM:266200]. A frequent cause of hereditary non-spherocytic hemolytic anemia. Clinically, pyruvate kinase-deficient patients suffer from a highly variable degree of chronic hemolysis, ranging from severe neonatal jaundice and fatal anemia at birth, severe transfusion-dependent chronic hemolysis, moderate hemolysis with exacerbation during infection, to a fully compensated hemolysis without

apparent anemia.

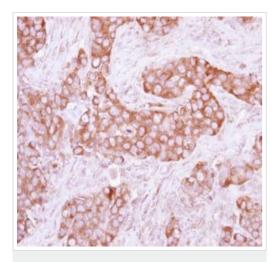
序列相似性 Belongs to the pyruvate kinase family.

图片



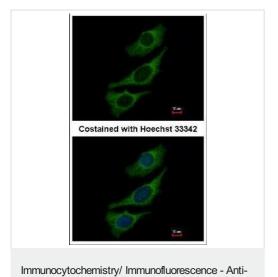
Anti-PKLR antibody (ab125697) at 1/1000 dilution + A549 cell lysate at 30 μg

Predicted band size: 62 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PKLR antibody (ab125697)

ab125697, at 1/250 dilution, staining PKLR in Formalin-fixed, Paraffin-embedded Human breast cancer tissue by Immunohistochemistry.



PKLR antibody (ab125697)

ab125697, at 1/500 dilution, staining PKLR in Methanol-fixed HeLa cells by Immunocytochemistry/Immunofluorescence.

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

Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- · We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit https://www.abcam.cn/abpromise or contact our technical team.

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