

## Product datasheet

# Anti-PKLR antibody ab137787

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### Overview

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<b>Product name</b>	Anti-PKLR antibody
<b>Description</b>	Rabbit polyclonal to PKLR
<b>Host species</b>	Rabbit
<b>Tested applications</b>	<b>Suitable for:</b> WB
<b>Species reactivity</b>	<b>Reacts with:</b> Mouse, Rat, Human
<b>Immunogen</b>	Recombinant fragment, corresponding to a region within amino acids 254-574 of Human PKLR
<b>Positive control</b>	Jurkat, Raji, K562, THP-1 and NCI-H929 cell extracts; Mouse and Rat liver extracts
<b>General notes</b>	<p>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.</p> <p>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&amp;As</p>

### Properties

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<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.
<b>Storage buffer</b>	pH: 7.00 Preservative: 0.01% Thimerosal (merthiolate) Constituents: 1.21% Tris, 0.75% Glycine, 10% Glycerol (glycerin, glycerine)
<b>Purity</b>	Immunogen affinity purified
<b>Clonality</b>	Polyclonal
<b>Isotype</b>	IgG

### Applications

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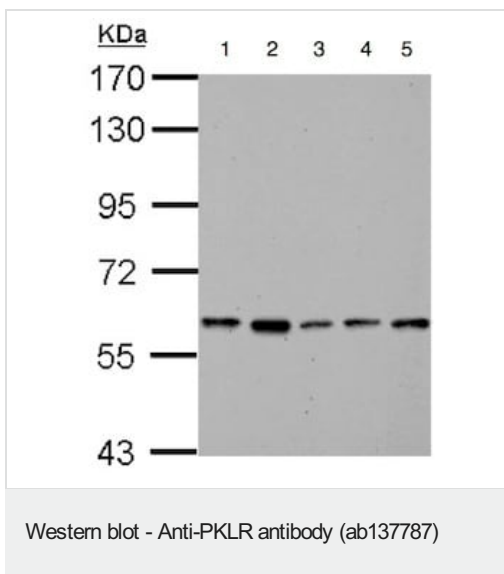
**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab137787 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		1/500 - 1/3000. Predicted molecular weight: 62 kDa.

## Target

<b>Function</b>	Plays a key role in glycolysis.
<b>Pathway</b>	Carbohydrate degradation; glycolysis; pyruvate from D-glyceraldehyde 3-phosphate: step 5/5.
<b>Involvement in disease</b>	<p>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.</p> <p>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.</p>
<b>Sequence similarities</b>	Belongs to the pyruvate kinase family.

## Images



**All lanes :** Anti-PKLR antibody (ab137787) at 1/1000 dilution

**Lane 1 :** Jurkat cell extract

**Lane 2 :** Raji cell extract

**Lane 3 :** K562 cell extract

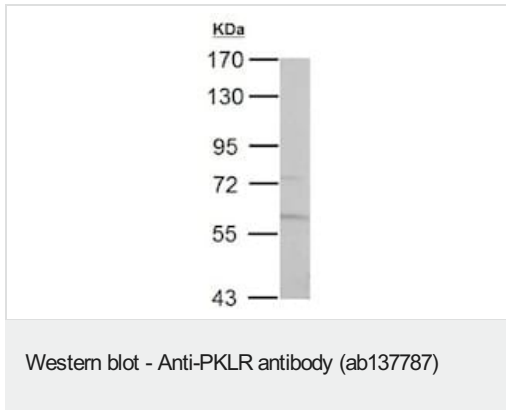
**Lane 4 :** THP-1 cell extract

**Lane 5 :** NCI-H929 cell extract

Lysates/proteins at 30 µg per lane.

**Predicted band size:** 62 kDa

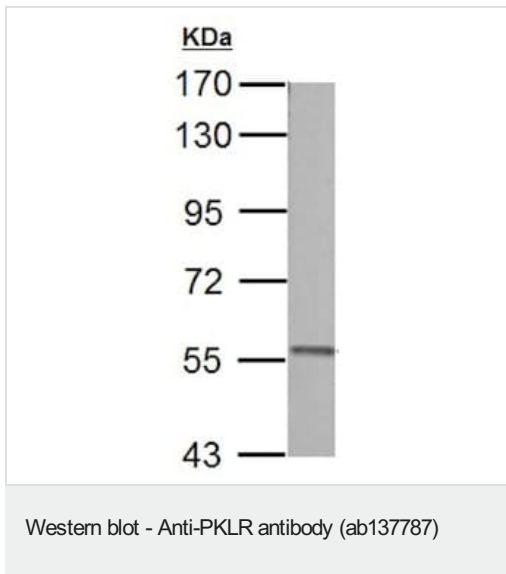
7.5% SDS PAGE



Anti-PKLR antibody (ab137787) at 1/500 dilution + Mouse liver extract at 50 µg

**Predicted band size:** 62 kDa

7.5% SDS PAGE



Anti-PKLR antibody (ab137787) at 1/500 dilution + Rat liver extract at 50 µg

**Predicted band size:** 62 kDa

7.5% SDS PAGE

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

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