abcam

Product datasheet

Anti-PERK antibody ab65142

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Overview

Product name Anti-PERK antibody

Description Rabbit polyclonal to PERK

Host species Rabbit

Tested applications
Suitable for: ICC/IF, WB
Species reactivity
Reacts with: Human

Immunogen A synthesized peptide derived from an internal region of human PERK

General notes

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

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: PBS, 50% Glycerol, 0.87% Sodium chloride

Without Mg2+ and Ca2+

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab65142 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
ICC/IF	★ ☆ ☆ ☆ ☆ (1)	1/500 - 1/1000.
WB	*****(1)	1/500 - 1/1000. Detects a band of approximately 125 kDa (predicted molecular weight: 125 kDa).

Target

Function Phosphorylates the alpha subunit of eukaryotic translation-initiation factor 2 (EIF2), leading to its inactivation and thus to a rapid reduction of translational initiation and repression of global protein synthesis. Serves as a critical effector of unfolded protein response (UPR)-induced G1 growth arrest due to the loss of cyclin D1.

Tissue specificity Ubiquitous. A high level expression is seen in secretory tissues.

Involvement in disease

Defects in EIF2AK3 are the cause of Wolcott-Rallison syndrome (WRS) [MIM:226980]; also known as multiple epiphyseal dysplasia with early-onset diabetes mellitus. WRS is a rare autosomal recessive disorder, characterized by permanent neonatal or early infancy insulindependent diabetes and, at a later age, epiphyseal dysplasia, osteoporosis, growth retardation and other multisystem manifestations, such as hepatic and renal dysfunctions, mental retardation

and cardiovascular abnormalities.

Sequence similaritiesBelongs to the protein kinase superfamily. Ser/Thr protein kinase family. GCN2 subfamily.

Contains 1 protein kinase domain.

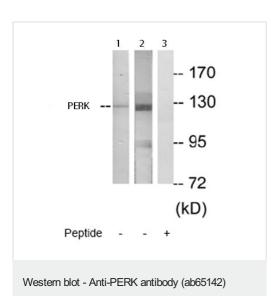
DomainThe lumenal domain senses perturbations in protein folding in the ER, probably through reversible

interaction with HSPA5/BIP.

Post-translationalAutophosphorylated.modificationsN-glycosylated.

Cellular localization Endoplasmic reticulum membrane.

Images



All lanes: Anti-PERK antibody (ab65142) at 1/500 dilution

Lane 1 : Extracts from Hela cells

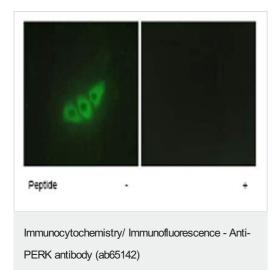
Lane 2: Extracts from HepG2 cells

Lane 3: Extracts from HepG2 cells, plus immunising peptide

Predicted band size: 125 kDa **Observed band size:** 125 kDa

The amount of positive control loading for the WB is 5-30 ug of total protein.

The amount of the peptide for the WB is 5-10 ug.



Immunofluorescence analysis of HepG2 cells using ab65142 at a 1/500-1/1000 dilution. Left image untreated. Right image treated with immunising peptide.

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