# abcam

# Product datasheet

# Anti-ACK1 antibody ab65108

★★★★★ 1 Abreviews 3 References 2 Images

#### Overview

Product name Anti-ACK1 antibody

**Description** Rabbit polyclonal to ACK1

Host species Rabbit

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

Predicted to work with: Mouse, Cow, Dog, Rhesus monkey

Immunogen Synthetic peptide conjugated to KLH derived from within residues 1000 to the C-terminus of

Human ACK1.Read Abcam's proprietary immunogen policy(Peptide available as ab74487.)

**Positive control** This antibody gave a positive signal in the following Whole Cell Lysates: HeLa, HepG2, PC12,

SHSY-5Y, T24/83, HT 1080

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 +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

 $80^{\circ}\text{C}\,.$  Avoid freeze / thaw cycle.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising

agent. If you would like information about the formulation of a specific lot, please contact our

scientific support team who will be happy to help.

Purity Immunogen affinity purified

1

**Clonality** Polyclonal

**Isotype** IgG

#### **Applications**

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab65108 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	<b>★</b> ★ ★ ★ ★ <b>(1)</b>	Use a concentration of 1 µg/ml.
WB		Use a concentration of 1 µg/ml. Detects a band of approximately 120 kDa (predicted molecular weight: 114 kDa).

## **Target**

**Function** 

Non-receptor tyrosine kinase that regulates the activity of a number of proteins by tyrosine phosphorylation especially proteins critical for cell survival, cell growth, and proliferation. Activates AKT1 by phosphorylating it on 'Tyr-176' resulting in its activation. Phosphorylates AR on 'Tyr-267' and 'Tyr-363' and promotes its recruitment to the androgen-responsive enhancers (AREs). Phosphorylates WWOX on 'Tyr-287'. Downstream effector of CDC42 which mediates CDC42-dependent cell migration via phosphorylation of BCAR1. Binds to both poly- and mono-ubiquitin and regulates ligand-induced degradation of EGFR. Participates in clathrin-mediated endocytosis. May be involved both in adult synaptic function and plasticity and in brain development.

**Tissue specificity** 

The Tyr-284 phosphorylated form shows a significant increase in expression in breast cancers during the progressive stages i.e. normal to hyperplasia (ADH), ductal carcinoma in situ (DCIS), invasive ductal carcinoma (IDC) and lymph node metastatic (LNMM) stages. It also shows a significant increase in expression in prostate cancers during the progressive stages.

Sequence similarities

Belongs to the protein kinase superfamily. Tyr protein kinase family.

Contains 1 CRIB domain.

Contains 1 protein kinase domain.

Contains 1 SH3 domain. Contains 1 UBA domain.

Domain

The EBD (EGFR-binding domain) domain is necessary for interaction with EGFR.

The SAM-like domain is necessary for NEDD4-mediated ubiquitination. Promotes membrane

 $\label{localization} \mbox{localization and dimerization to allow for autophosphorylation}.$ 

The UBA domain binds both poly- and mono-ubiquitin.

Post-translational modifications

interaction with SRC.

Ubiquitinated by NEDD4. Its EGF-induced degradation is EGFR activation-dependent and is

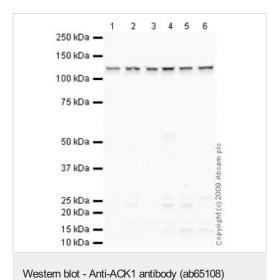
processed by lysosomes, not proteasomes.

**Cellular localization** 

Cell membrane. Nucleus. Endosome. Cell junction > adherens junction. Cytoplasmic vesicle membrane. The Tyr-284 phosphorylated form is expressed both in the membrane and nucleus.

Co-localizes with EGFR on the endosomes.

## **Images**



All lanes: Anti-ACK1 antibody (ab65108) at 1 µg/ml

Lane 1 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

**Lane 2 :** HepG2 (Human hepatocellular liver carcinoma cell line) Whole Cell Lysate

Lane 3: PC12 (Rat adrenal pheochromocytoma cell line) Whole Cell Lysate

Lane 4 : SHSY-5Y (Human neuroblastoma cell line) Whole Cell Lysate

Lane 5 : T24/83 (Human bladder carcinoma) Whole Cell LysateLane 6 : HT 1080 (Human fibrosarcoma) Whole Cell Lysate

Lysates/proteins at 10 µg per lane.

# **Secondary**

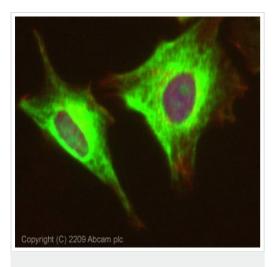
 $\begin{tabular}{ll} \textbf{All lanes:} Goat polyclonal to Rabbit lgG - H\&L - Pre-Adsorbed (HRP) at 1/3000 dilution \end{tabular}$ 

Developed using the ECL technique.

Performed under reducing conditions.

**Predicted band size:** 114 kDa **Observed band size:** 120 kDa

Exposure time: 4 minutes



Immunocytochemistry/ Immunofluorescence - Anti-ACK1 antibody (ab65108)

ICC/IF image of ab65108 stained HeLa cells. The cells were 4% PFA fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab65108, 1µg/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat anti-rabbit lgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.

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

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