abcam

Product datasheet

Anti-Cl4orf133 antibody ab125084

1 References 2 Images

Overview

Product name Anti-C14orf133 antibody

Description Rabbit polyclonal to C14orf133

Host species Rabbit

Tested applications Suitable for: WB, IP

Species reactivity Reacts with: Mouse, Human

Predicted to work with: Chimpanzee, Rhesus monkey, Gorilla, Orangutan

Immunogen Synthetic peptide, corresponding to a region between amino acids 443-493 of Human C14orf133

(NP_071350.2).

Positive control HeLa, 293T, Jurkat and NIH3T3 whole cell lysates.

General notesThe 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. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Storage buffer pH: 7

Preservative: 0.09% Sodium azide Constituent: 99% Tris citrate/phosphate

pH 7 to 8

Purity Immunogen affinity purified

Purification notes ab125084 was affinity purified using an epitope specific to C14orf133 immobilized on solid

support.

Clonality Polyclonal

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Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab125084 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		1/2000 - 1/10000. Predicted molecular weight: 57 kDa.
IP		Use at 2-10 µg/mg of lysate.

Target

Function Plays a role in lysosomal trafficking. May play a role in epithelial polarization through stabilization

of apical membrane protein content, possibly via the RAB11A-dependent apical recycling

pathway. Also involved in direct or indirect transcriptional regulation of E-cadherin.

Involvement in diseaseDefects in VIPAR are the cause of arthrogryposis-renal dysfunction-cholestasis syndrome type 2

(ARCS2) [MIM:613404]. ARCS2 is an autosomal recessive multisystem disorder, characterized by neurogenic arthrogryposis multiplex congenita, renal tubular dysfunction and neonatal cholestasis with bile duct hypoplasia and low gamma glutamyl transpeptidase activity. Platelet

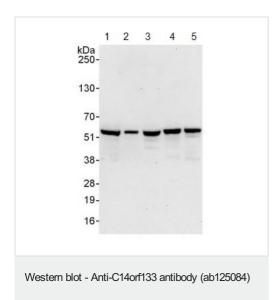
dysfunction is common. Note=In liver, CEACAM5 and ABCB11 are mislocalized and E-cadherin

expression is decreased.

Sequence similaritiesBelongs to the VIPAR family.

Cellular localization Early endosome. Recycling endosome. Late endosome.

Images



All lanes: Anti-C14orf133 antibody (ab125084) at 0.1 µg/ml

Lane 1: HeLa whole cell lysate at 50 μg **Lane 2**: HeLa whole cell lysate at 15 μg

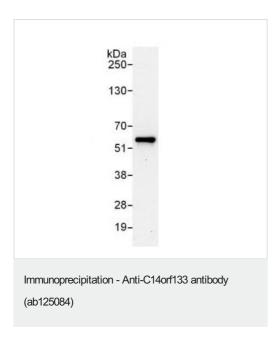
Lane 3: 293T whole cell lysate at 50 μg **Lane 4**: Jurkat whole cell lysate at 50 μg

Lane 5: NIH3T3 whole cell lysate at 50 µg

Developed using the ECL technique.

Predicted band size: 57 kDa

Exposure time: 30 seconds



Detection of C14orf133 by Western Blot of Immunprecipitate. ab125084 at 1μ g/ml staining C14orf133 in HeLa whole cell lysate immunoprecipitated using ab125084 at 6μ g/mg lysate (1 mg/IP; 20% of IP loaded/lane).

Detection: Chemiluminescence with exposure time of 30 seconds.

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

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