

Anti-C14orf133 antibody ab125084

1 [References](#) **2** [画像数](#)

製品の概要

製品名	Anti-C14orf133 antibody
製品の詳細	Rabbit polyclonal to C14orf133
由来種	Rabbit
アプリケーション	適用あり: WB, IP
種交差性	交差種: Mouse, Human 交差が予測される動物種: Chimpanzee, Rhesus monkey, Gorilla, Orangutan 
免疫原	Synthetic peptide, corresponding to a region between amino acids 443-493 of Human C14orf133 (NP_071350.2).
ポジティブ・コントロール	HeLa, 293T, Jurkat and NIH3T3 whole cell lysates.
特記事項	<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&As</p>

製品の特性

製品の状態	Liquid
保存方法	Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
バッファー	pH: 7 Preservative: 0.09% Sodium azide Constituent: 99% Tris citrate/phosphate
精製度	pH 7 to 8 Immunogen affinity purified
特記事項 (精製)	ab125084 was affinity purified using an epitope specific to C14orf133 immobilized on solid support.
ポリ/モノ	ポリクローナル

アイソタイプ

IgG

アプリケーション

The Abpromise guarantee

Abpromise保証は、次のテスト済みアプリケーションにおけるab125084の使用に適用されます

アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご確認ください。

アプリケーション	Abreviews	特記事項
WB		1/2000 - 1/10000. Predicted molecular weight: 57 kDa.
IP		Use at 2-10 µg/mg of lysate.

ターゲット情報

機能

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.

関連疾患

Defects 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.

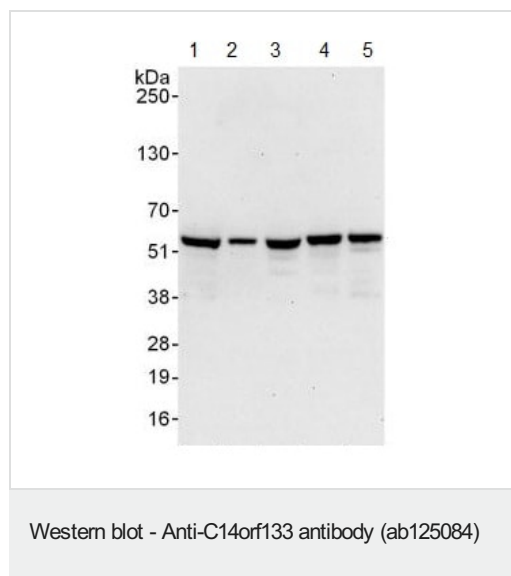
配列類似性

Belongs to the VIPAR family.

細胞内局在

Early endosome. Recycling endosome. Late endosome.

画像



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

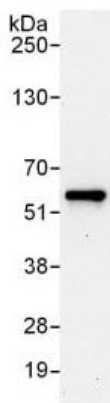
Lane 4 : Jurkat whole cell lysate at 50 µg

Lane 5 : NIH3T3 whole cell lysate at 50 µg

Developed using the ECL technique.

Predicted band size: 57 kDa

Exposure time: 30 seconds



Immunoprecipitation - Anti-C14orf133 antibody
(ab125084)

Detection of C14orf133 by Western Blot of Immunoprecipitate.

ab125084 at 1 µg/ml staining C14orf133 in HeLa whole cell lysate immunoprecipitated using ab125084 at 6 µg/mg lysate (1 mg/IP; 20% of IP loaded/lane).

Detection: Chemiluminescence with exposure time of 30 seconds.

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