abcam

Product datasheet

Anti-CrkL antibody [Y244] ab32018





RabMAb

4 References 7 图像

概述

产品名称 Anti-CrkL抗体[Y244]

宿主 Rabbit

特异性 ab32018 recognises CrkL (Crk-like)protein.

The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for

mouse and rat.

经测试应用 适用于: WB, IHC-P, IP, Flow Cyt, ICC/IF

种属反应性 与反应: Mouse, Rat, Human

免疫原 Synthetic peptide within Human CrkL aa 250-350 (C terminal). The exact sequence is proprietary.

阳性对照 WB: Wild-type HeLa, CRKL knockout HeLa and K-562 cells. K-562 whole cell lysate. Mouse and

rat thymus lysate. IHC-P: Human colon tissue; ICC/IF: K-562 cells; Flow Cyt: K-562 cells.

常规说明 This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**[®] **patents**.

性能

形式 Liquid

存放说明 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

存储溶液 pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

纯**度** Protein A purified

克隆 单克隆

1

克隆编号 Y244

同种型 IgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab32018于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		1/1000. Detects a band of approximately 36 kDa (predicted molecular weight: 33 kDa).
IHC-P		1/100. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for mouse and rat.
IP		1/50.
Flow Cyt		1/20.
ICC/IF		1/50.

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蚆	47	К

功能 May mediate the transduction of intracellular signals.

序列相似性 Belongs to the CRK family.

Contains 1 SH2 domain.
Contains 2 SH3 domains.

图片



Western blot - Anti-CrkL antibody [Y244] (ab32018)

All lanes: Anti-CrkL antibody [Y244] (ab32018) at 1/1000 dilution (Purified)

Lane 1: K-562 (Human chronic myelogenous leukemia

lymphoblast) whole cell lysate

Lane 2: Mouse thymus lysate

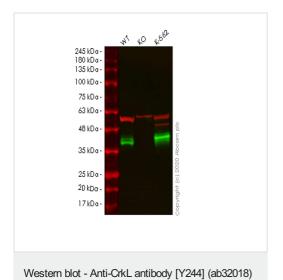
Lane 3: Rat thymus lysate

Secondary

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000

dilution

Predicted band size: 33 kDa



All lanes: Anti-CrkL antibody [Y244] (ab32018) at 1/1000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: CRKL knockout HeLa cell lysate

Lane 3: K-562 cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes: Goat anti-Rabbit IgG H&L (IRDye® 800CW)

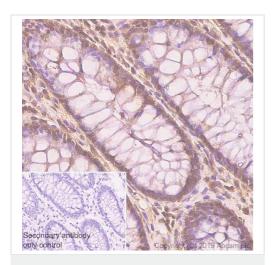
preadsorbed (ab216773) at 1/10000 dilution

Predicted band size: 33 kDa Observed band size: 37 kDa

Lanes 1-3: Merged signal (red and green). Green - ab32018 observed at 37 kDa. Red - loading control ab7291 observed at 50 kDa.

ab32018 Anti-CrkL antibody [Y244] was shown to specifically react with CrkL in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab265993 (knockout cell lysate ab257397) was used. Wild-type and CrkL knockout samples were subjected to SDS-PAGE. ab32018 and Anti-alpha Tubulin antibody [DM1A] -Loading Control (ab7291) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye®

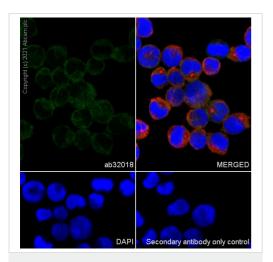
680RD) preadsorbed (<u>ab216776</u>) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CrkL antibody [Y244] (ab32018)

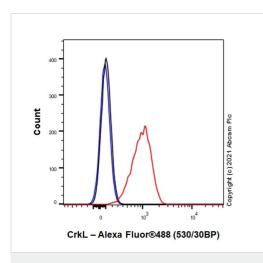
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human colon tissue sections labeling CrkL with purified ab32018 at 1/100 dilution (2.24 µg/mL). Heat mediated antigen retrieval was performed using Heat mediated antigen retrieval using Bond™ Epitope Retrieval Solution 2 (pH 9.0). Tissue was counterstained with Hematoxylin. Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) secondary antibody was used at 1/0 dilution. PBS instead of the primary antibody was used as the negative control.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument.



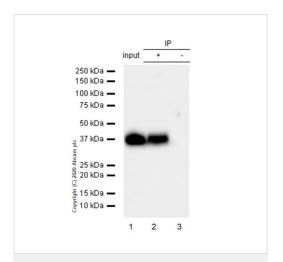
Immunocytochemistry/ Immunofluorescence - Anti-CrkL antibody [Y244] (ab32018)

Immunocytochemistry analysis of K-562 (Human chronic myelogenous leukemia lymphoblast) cells labeling CrkL with purified ab32018 at 1/50 dilution (4.5 μ g/mL). Cells were fixed in 100% Methanol and permeabilized with 0.1% tritonX-100. Cells were counterstained with Ab195889 Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594) 1/200 (2.5 μ g/mL). Goat anti rabbit lgG (Alexa Fluor®488, **ab150077**) was used as the secondary antibody at 1/1000 (2 μ g/mL) dilution. DAPI (blue) was used as nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



Flow Cytometry - Anti-CrkL antibody [Y244] (ab32018)

Flow Cytometry analysis of K-562 (Human chronic myelogenous leukemia lymphoblast) cells labelling CrkL with purified ab32018 at 1/20 dilution (10 μ g/mL) (Red). Cells were fixed with 4% Paraformaldehyde and permeabilised with 90% Methanol. A Goat anti rabbit lgG (Alexa Fluor® 488, <u>ab150077</u>) secondary antibody was used at 1/2000. Isotype control - Rabbit monoclonal lgG (Black). Unlabelled control - Cell without incubation with primary antibody and secondary antibody (Blue).



Immunoprecipitation - Anti-CrkL antibody [Y244] (ab32018)

Purified ab32018 at 1/50 dilution (2 μ g) immunoprecipitating CrkL in K-562 whole cell lysate.

Lane 1 (input): K-562 (Human chronic myelogenous leukemia lymphoblast) whole cell lysate $10\mu g$

Lane 2 (+): ab32018 + K-562 whole cell lysate.

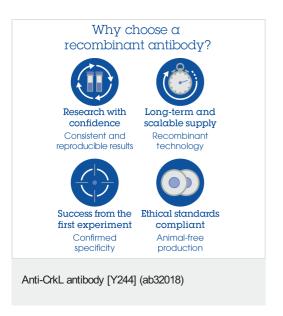
Lane 3 (-): Rabbit monoclonal lgG ($\underline{ab172730}$) instead of ab32018 in K-562 whole cell lysate.

VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) (1/1000 dilution) was used for Western blotting.

Blocking Buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM/TBST.

Observed band size: 37 kDa



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