abcam

Product datasheet

Anti-CD3G antibody [EPR4517] ab134096



重组 RabMAb

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概述

产品名称 Anti-CD3G抗体[EPR4517]

描述 兔单克隆抗体[EPR4517] to CD3G

宿主 Rabbit

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

种属反应性 与反应: Mouse, Human

免疫原 Synthetic peptide within Human CD3G (C terminal). The exact sequence is proprietary.

Database link: P09693

阳性对照 WB: Jurkat and mouse thymus lysates. IHC-P: Human tonsil, human colon and mouse spleen

tissues. ICC/IF: Jurkat cells. IP: Jurkat cell lysate. Flow Cyt (intra): Jurkat 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**.

Rat: We have preliminary internal testing data to indicate this antibody may not react with this

species. Please contact us for more information.

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Stable for 12 months at -20°C.

存储溶液 pH: 7.2

Preservative: 0.01% Sodium azide

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

纯度 Protein A purified

克隆 单克隆

克隆编号 EPR4517

同种型 IgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab134096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		1/20000 - 1/200000. Predicted molecular weight: 20 kDa. For unpurified use at 1/10000 - 1/50000.
IP		1/40. For unpurified use at 1/10 - 1/100.
IHC-P		1/1000. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. For unpurified use at 1/250 - 1/500.
Flow Cyt (Intra)		1/70.
ICC/IF		1/1000. For unpurified use at 1/500 - 1/1000.

靶标

功能 The CD3 complex mediates signal transduction.

序列相似性 Contains 1 lg-like (immunoglobulin-like) domain.

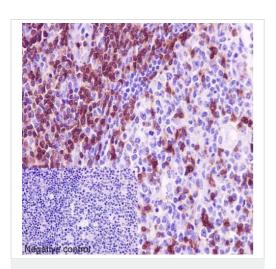
Contains 1 ITAM domain.

结**构域** A di-leucine motif and a tyrosine-based motif are individually sufficient to induce both endocytosis

and delivery to lysosomes.

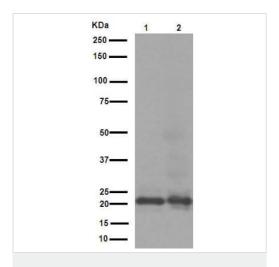
细**胞定位** Membrane.

图片



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD3G antibody
[EPR4517] (ab134096)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human tonsil tissue labelling CD3G with purified ab134096 at 1/1000. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit lgG was used as the secondary antibody. Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.



Western blot - Anti-CD3G antibody [EPR4517] (ab134096)

All lanes : Anti-CD3G antibody [EPR4517] (ab134096) at 1/50000 dilution (purified)

Lane 1: Jurkat cell lysate

Lane 2: Human fetal thymus lysate

Lysates/proteins at 20 µg per lane.

Secondary

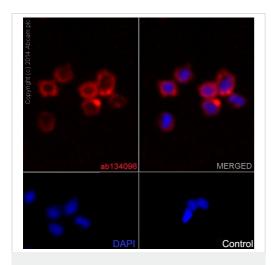
All lanes : Peroxidase-conjugated goat anti-rabbit $\lg G \, (H+L)$ at

1/1000 dilution

Predicted band size: 20 kDa **Observed band size:** 18-28 kDa

Blocking buffer and concentration: 5% NFDM/TBST.

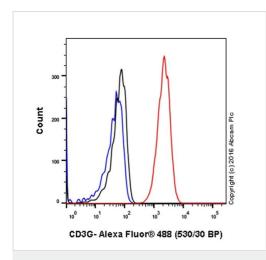
Diluting buffer and concentration: 5% NFDM /TBST.



Immunocytochemistry/ Immunofluorescence - Anti-CD3G antibody [EPR4517] (ab134096)

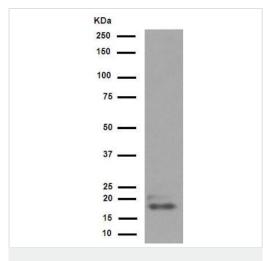
Immunocytochemistry/Immunofluorescence analysis of Jurkat cells labelling CD3G with purified ab134096 at 1/1000. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. An Alexa Fluor $^{\!(\!8\!)}$ 555-conjugated goat anti-rabbit lgG (1/500) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain.

Control: primary antibody (1/1000) and secondary antibody, <u>**ab150113**</u>, an Alexa Fluor[®] 488-conjugated goat anti-mouse lgG (1/500).



Flow Cytometry (Intracellular) - Anti-CD3G antibody [EPR4517] (ab134096)

Intracellular Flow Cytometry analysis of Jurkat (human acute T cell leukemia) cells labeling CD3G with purified ab134096 at 1/70 dilution (10ug/mL) (red). Cells were fixed with 4% paraformaldehyde and permeabilised with 90% methanol. A Goat anti rabbit lgG (Alexa Fluor[®]488) (1/2000 dilution) was used as the secondary antibody. Rabbit monoclonal lgG (Black) was used as the isotype control, cells without incubation with primary antibody and secondary antibody (Blue) was used as the unlabeled control.



Western blot - Anti-CD3G antibody [EPR4517] (ab134096)

Anti-CD3G antibody [EPR4517] (ab134096) at 1/50000 dilution (purified) + Mouse thymus tissue lysate at 20 µg

Secondary

Peroxidase-conjugated goat anti-rabbit lgG (H+L) at 1/1000 dilution

Predicted band size: 20 kDa **Observed band size:** 18-28 kDa

Blocking buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM /TBST.

KDa 1 2

250—
150—
100—
75—
50—
37—
25—
20—
15—
10—

Western blot - Anti-CD3G antibody [EPR4517] (ab134096)

All lanes : Anti-CD3G antibody [EPR4517] (ab134096) at 1/10000 dilution (unpurified)

Lane 1: Jurkat cell lysate

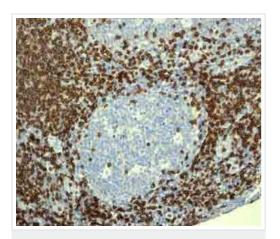
Lane 2: Mouse thymus tissue lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes: HRP-conjugated goat anti-rabbit lgG at 1/2000 dilution

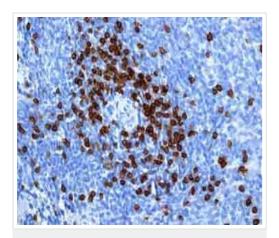
Predicted band size: 20 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD3G antibody
[EPR4517] (ab134096)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human tonsil tissue labelling CD3G with unpurified ab134096.

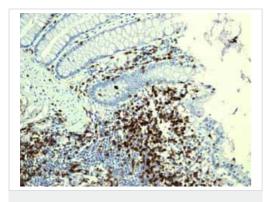
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD3G antibody
[EPR4517] (ab134096)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse spleen tissue labelling CD3G with unpurified ab134096.

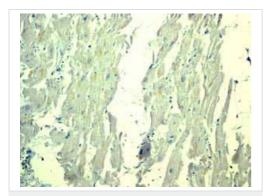
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD3G antibody
[EPR4517] (ab134096)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human colon (lymphocytes) tissue labelling CD3G with unpurified ab134096.

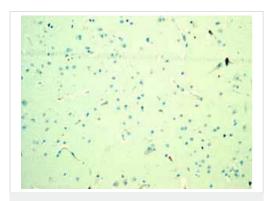
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD3G antibody
[EPR4517] (ab134096)

Immunohistochemistry (Formalin/PFA-fixed parffin-embedded sections) analysis of human skeletal muscle tissue. Unpurified **ab32362** shows negative staining.

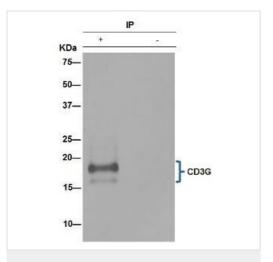
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD3G antibody
[EPR4517] (ab134096)

Immunohistochemistry (Formalin/PFA-fixed parffin-embedded sections) analysis of human brain tissue. Unpurified <u>ab32362</u> shows negative staining.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

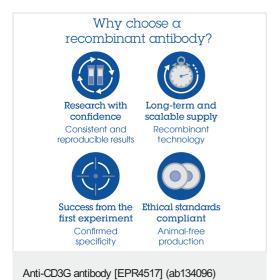


Immunoprecipitation - Anti-CD3G antibody [EPR4517] (ab134096)

ab134096 (purified) at 1/40 immunoprecipitating CD3G in Jurkat cell lysate (Lane 1). Lane 2 - PBS. For western blotting, a HRP-conjugated anti-rabbit lgG, specific to the non-reduced form of lgG was used as the secondary antibody (1/1500).

Blocking buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM /TBST.



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