abcam

Product datasheet

Donkey Anti-Rabbit IgG H&L (Alexa Fluor® 488) ab150073

★★★★★ 7 Abreviews 474 References 5 图像

概述

宿主 Donkey **靶标种属** Rabbit

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

免疫原 The details of the immunogen for this antibody are not available.

偶联物 Alexa Fluor® 488. Ex: 495nm, Em: 519nm

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.

Avoid freeze / thaw cycle. Stable for 12 months at -20°C. Store In the Dark.

存储溶液 Preservative: 0.02% Sodium azide

Constituents: 23% Glycerol (glycerin, glycerine), PBS, 1% BSA

纯**度** Immunogen affinity purified

纯**化**说明 This antibody was isolated by affinity chromatography using antigen coupled to agarose beads.

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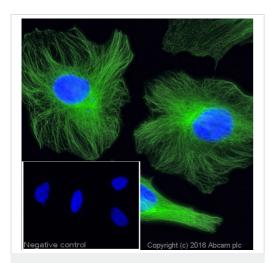
应用

The Abpromise guarantee Abpromise™承诺保证使用ab150073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
ICC/IF	★★★★★ (2)	1/200 - 1/1000.
Flow Cyt	**** (1)	1/2000 - 1/4000.
IHC-P		Use at an assay dependent concentration.
ELISA		Use at an assay dependent concentration.
IHC-Fr		Use at an assay dependent concentration.

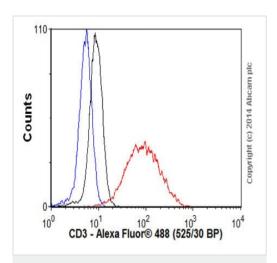
图片



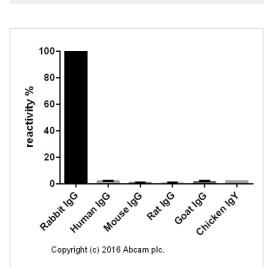
Immunocytochemistry/ Immunofluorescence -Donkey Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150073) ICC/IF image of <u>ab6046</u> stained HeLa cells. The cells were 4% formaldehyde fixed (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then incubated in 1% BSA / 10% normal donkey serum / 0.3M glycine in 0.1% PBS-Tween for 1h to block nonspecific protein-protein interactions. The cells were then incubated with the antibody (<u>ab6046</u>, 1µg/ml) overnight at +4°C. The secondary antibody (green) was ab150073 Alexa Fluor[®] 488 donkey anti-rabbit IgG (H+L) used at 1µg/ml for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.

The negative control (inset) is a secondary-only assay to demonstrate low non-specific binding of the secondary antibody.

This product also gave a positive signal under the same testing conditions in HeLa cells fixed with 4% formaldehyde (10min).



Flow Cytometry - Donkey Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150073)



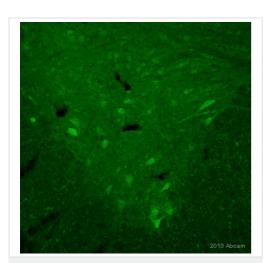
ELISA - Donkey Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150073)

Overlay histogram showing Jurkat cells stained with <u>ab16669</u> (red line). The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal donkey serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (<u>ab16669</u>, 1/1000 dilution) for 30 min at 22°C. The secondary antibody Donkey anti-rabbit lgG H&L (Alexa Fluor® 488) (ab150073) was used at 1/4000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit lgG (monclonal) (0.1µg/1x10⁶ cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.

Cross-reactivity of the polyclonal secondary antibody <u>ab182020</u> was tested using a sandwich ELISA approach. The wells were coated with the indicated lgG standards at 1 μ g/ml (50 μ l/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. <u>ab182020</u> was then added starting at 1 μ g/ml and gradually diluted 1/4 (50 μ l/well), followed by incubation for 2h. For the detection Goat anti-Donkey lgG H&L (HRP) (<u>ab6988</u>) was used at 1/20,000 dilution (50 μ l/well), followed by incubation for 1h at RT.

For the batch tested, <u>ab182020</u> showed a cross-reactivity below 2% towards human IgG, mouse IgG, rat IgG, goat IgG and chicken IgY.

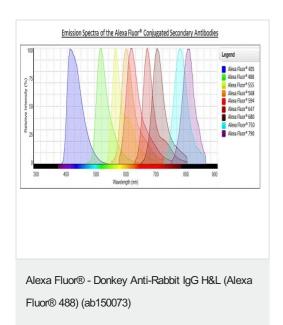
This data was developed using the unconjugated antibody (ab182020).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Donkey Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150073)

This image is courtesy of an Abreview submitted by Laura Comley.

<u>ab104603</u> staining DYNLL1 in mouse cervical spinal cord tissue sections by Immunohistochemistry (IHC-P - paraformaldehydefixed, paraffin-embedded sections). Tissue was fixed with paraformaldehyde, permeabilized with 0.1% Triton X-100, and blocked with 10% serum for 1 hour at 21°C. Samples were incubated with primary antibody (2 μ g/ml in PBS/10% serum/0.1% Triton X-100) for 16 hours at 4°C. An Alexa Fluor[®] 488-conjugated Donkey anti-rabbit IgG H&L (ab150073) (1/500) was used as the secondary antibody.



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