abcam

Product datasheet

Goat Anti-Mouse IgG H&L (Alexa Fluor® 488) ab150113

★★★★★ 15 Abreviews 1170 References 画像数 8

製品の概要

製品名 Goat Anti-Mouse IgG H&L (Alexa Fluor® 488)

由来種Goatターゲット生物種Mouse

アプリケーション 適用あり: IHC-Fr, ICC/IF, ELISA, Flow Cyt, IHC-P

免疫原 Other Immunogen Type corresponding to Mouse IgG.

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

製品の特性

製品の状態 Liquid

保存方法 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.

ארעדעד Preservative: 0.02% Sodium azide

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

精製度 Immunogen affinity purified

特記事項(精製) This antibody was isolated by affinity chromatography using antigen coupled to agarose beads.

ポリ/モノ ポリクローナル

アイソタイプ lgG

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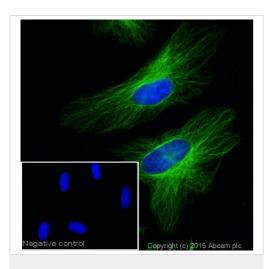
outlicensing@thermofisher.com.

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab150113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検討ください。

アプリケーション	Abreviews	特記事項
IHC-Fr	★★★★★ (3)	Use at an assay dependent concentration.
ICC/IF	**** <u>(6)</u>	1/200 - 1/1000.
ELISA		Use at an assay dependent concentration.
Flow Cyt		1/2000. ab170190 - Mouse monoclonal lgG1 (Alexa Fluor® 488), is suitable for use as an isotype control to complement this secondary antibody.
IHC-P	★★★★★ (3)	Use at an assay dependent concentration.

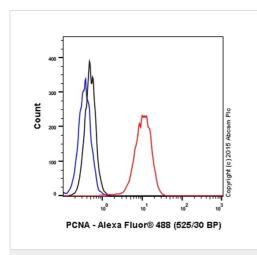
画像



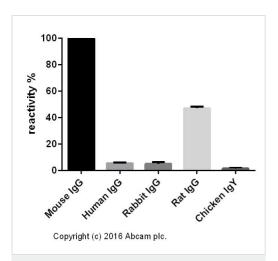
Immunocytochemistry/ Immunofluorescence - Goat
Anti-Mouse IgG H&L (Alexa Fluor® 488) (ab150113)

ICC/IF image of <u>ab7291</u> stained HeLa cells. The cells were 100% methanol fixed (5 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then incubated in 1% BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to block non-specific protein-protein interactions. The cells were then incubated with the antibody (<u>ab7291</u>, 1 μ g/ml) overnight at +4°C. The secondary antibody (green) was ab150113 Alexa Fluor[®] 488 goat anti-mouse lgG (H+L) used at 2 μ g/ml for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43 μ M.

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



Flow Cytometry - Goat Anti-Mouse IgG H&L (Alexa Fluor® 488) (ab150113)



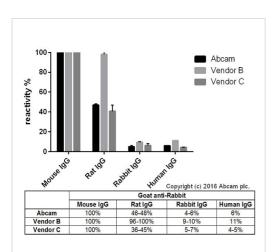
ELISA - Goat Anti-Mouse IgG H&L (Alexa Fluor® 488) (ab150113)

Overlay histogram showing HeLa cells stained with <u>ab29</u> (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween 20 for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (<u>ab29</u>, 0.1µg/1x10⁶ cells) for 30 min at 22°C. The secondary antibody used was Alexa Fluor[®] 488 goat anti-mouse lgG (H&L) (ab150113) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse lgG2a [ICIGG2A] (<u>ab91361</u>, 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.

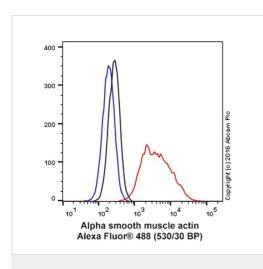
Cross-reactivity of the polyclonal secondary antibody <u>ab182017</u> was tested using a sandwich ELISA approach. The wells were coated with the indicated IgG standards at 1 μ g/ml (50 μ l/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. <u>ab182017</u> was then added starting at 1 μ g/ml and gradually diluted 1/4 (50 μ l/well), followed by incubation for 2h. For the detection Donkey anti-Goat IgG H&L (HRP) (<u>ab6885</u>) was used at 1/10,000 dilution (50 μ l/well), followed by incubation for 1h at RT.

For the batch tested, <u>ab182017</u> showed a cross-reactivity below 2% towards Chicken IgY, 6% towards Human IgG, 7% towards Rabbit IgG and 47% towards Rat IgG.

These data were developed using the unconjugated antibody (ab182017).



ELISA - Goat Anti-Mouse IgG H&L (Alexa Fluor® 488) (ab150113)



Flow Cytometry - Goat Anti-Mouse IgG H&L (Alexa Fluor® 488) (ab150113)

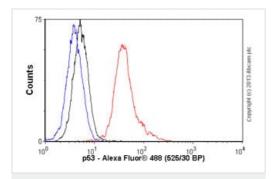
Cross-reactivity of Goat anti-Mouse IgG H&L (ab182017) and Goat anti-Mouse IgG H&L obtained from two different vendors was tested using a sandwich ELISA approach. The wells were coated with the indicated IgG standards (Rabbit, Human, Mouse and Rat) at 1 μ g/ml (50 μ l/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. Secondary antibodies were 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 Donkey anti-Goat IgG H&L (HRP) (ab6885) was used at 1/10,000 dilution (50 μ l/well), followed by incubation for 1h at RT. This data is from a representative dilution.

These data were developed using the unconjugated antibody (ab182017).

Overlay histogram showing SV40LT-SMC cells stained with ab7817 (red line). The cells were fixed with 4% formaldehyde (10 min) and then permeabilized with 0.1% PBS-Triton X-100 for 15 min. The cells were then incubated in 1x PBS / 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (ab7817, 0.1µg/1x10⁶ cells) for 30 min at 22°C. The secondary antibody used was Goat Anti-Mouse IgG H&L (Alexa Fluor[®] 488) (ab150113) at 1/2000 dilution for 30 min at 22°C lsotype control antibody (black line) was mouse IgG2a [18C8BC7AD10] (ab170191) used at the same concentration and conditions as the primary antibody. Unlabelled sample (blue line) was also used as a control.

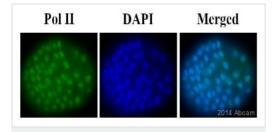
Acquisition of >5,000 events were collected using a 50 mW Blue laser (488nm) and 530/30 bandpass filter.

This antibody gave a positive signal in HeLa cells fixed with 80% methanol (5 min)/permeabilized with 0.1% PBS-Triton X-100 for 15 min used under the same conditions.



Flow Cytometry - Goat Anti-Mouse IgG H&L (Alexa Fluor® 488) (ab150113)

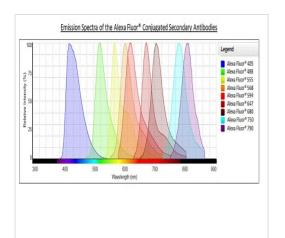
Overlay histogram showing HeLa cells stained with <u>ab26</u> (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (<u>ab26</u>, 1µg/1x10^6 cells) for 30 min at 22°C. The secondary antibody used was Alexa Fluor[®] 488 goat anti-mouse lgG (H&L) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse lgG1 [ICIGG1] (<u>ab91353</u>, 2µg/1x10^6 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. This antibody gave a positive signal in HeLa cells fixed with 4% paraformaldehyde (10 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



IHC - Wholemount - Goat Anti-Mouse IgG H&L (Alexa Fluor® 488) (ab150113)

This image is courtesy of an anonymous Abreview.

IHC - Wholemount of Caenorhabditis elegans embryo labelling RNA polymerase II CTD repeat YSPTSPS with <u>ab817</u>. Sample was incubated with primary antibody (1/100 in PBS + 3% BSA + 0.1% Triton-X 100) for 24 hours at 4°C . ab150113, an Alexa Fluor $^{\$}$ 488-conjugated goat anti-mouse IgG polyclonal (undiluted) was used as the secondary antibody.



Alexa Fluor® - Goat Anti-Mouse IgG H&L (Alexa Fluor® 488) (ab150113)

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