abcam

Product datasheet

Goat Anti-Rabbit IgG H&L (Alexa Fluor® 555) ab150078

**** 2 Abreviews 236 References 6 Images

Overview

Product name Goat Anti-Rabbit lgG H&L (Alexa Fluor® 555)

Host species Goat

Target species Rabbit

Tested applications Suitable for: IHC-Fr, ICC/IF, ELISA, IHC-P, Flow Cyt

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

Conjugation Alexa Fluor® 555. Ex: 555nm, Em: 565nm

Properties

Form Liquid

Storage instructions 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.

Storage buffer Preservative: 0.02% Sodium azide

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

Purity Immunogen affinity purified

Purification notesThe antibody was isolated by affinity chromatography using antigen coupled to agarose beads.

Clonality Polyclonal

Isotype IgG

General notesAlexa Fluor[®] is a registered trademark of Molecular Probes, Inc, a Thermo Fisher Scientific

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1

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Applications

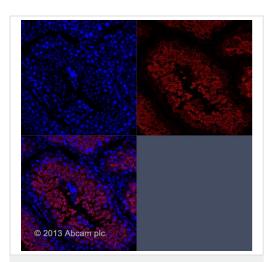
The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab150078 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-Fr		Use at an assay dependent concentration.
ICC/IF	**** (1)	1/200 - 1/1000.
ELISA		Use at an assay dependent concentration. Use at an assay dependent dilution
IHC-P	**** <u>(1)</u>	Use at an assay dependent concentration. Use at an assay dependent dilution
Flow Cyt		1/2000.

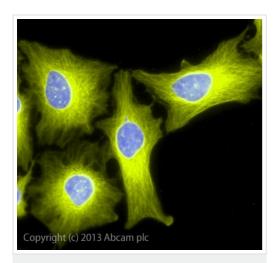
Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Goat Anti-Rabbit IgG H&L (Alexa Fluor® 555) (ab150078)

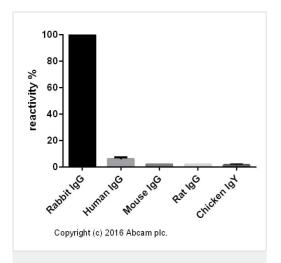
This image is courtesy of an abreview by Zachary Yu-Ching Lin

This data was done using a primary antibody to Calmegin (EPR11832) <u>ab171971</u>. The primary antibody concentration was 1: 100 and the secondary antibody concentration was 1:500.



Immunocytochemistry/ Immunofluorescence - Goat
Anti-Rabbit IgG H&L (Alexa Fluor® 555) (ab150078)

ICC/IF image of <u>ab6046</u> stained HeLa cells. The cells were 100% methanol fixed (5 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (<u>ab6046</u>, 1 μ g/ml) overnight at +4°C. The secondary antibody (yellow) was ab150078 Alexa Fluor[®] 555 goat anti-rabbit 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.

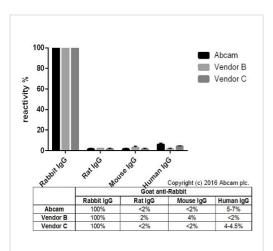


ELISA - Goat Anti-Rabbit IgG H&L (Alexa Fluor® 555) (ab150078)

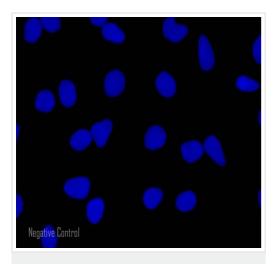
Cross-reactivity of the polyclonal secondary antibody <u>ab182016</u> was tested using a sandwich ELISA approach. The wells were coated with the indicated IgG standards at 1 μ g/ml (50 μ l/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. <u>ab182016</u> was then 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>ab182016</u> showed a cross-reactivity of 5-7% towards Human IgG and below 2% towards Mouse IgG, Rat IgG and Chicken IgY.

This data was developed using the unconjugated antibody (ab182016).



ELISA - Goat Anti-Rabbit IgG H&L (Alexa Fluor® 555) (ab150078)

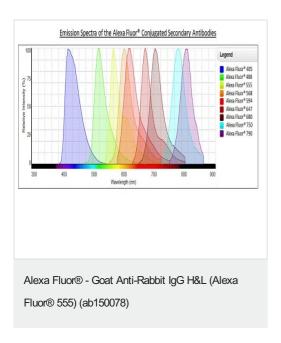


Immunocytochemistry/ Immunofluorescence - Goat
Anti-Rabbit IgG H&L (Alexa Fluor® 555) (ab150078)

Cross-reactivity of Goat anti-Rabbit IgG H&L (ab182016) and Goat anti-Rabbit 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.

This data was developed using the unconjugated antibody (ab182016).

HeLa cells showing negative staining by ICC/IF using only secondary antibody. The cells were 100% methanol fixed (5 min) and then incubated in 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The secondary antibody (yellow) was ab150078 Alexa Fluor $^{\rm B}$ 555 goat anti-rabbit IgG (H+L) used at 2µg/ml for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.



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