# abcam

# Product datasheet

# Goat Anti-Mouse IgG H&L (Alexa Fluor® 647) ab150115

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### Overview

Product name Goat Anti-Mouse IgG H&L (Alexa Fluor® 647)

Host species Goat

Target species Mouse

**Specificity** ab150115 is specific to Mouse IgG.

ab150115 has less than 47% cross-reactivity with rat lgG.

**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® 647. Ex: 652nm, Em: 668nm

## **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 notes**This antibody was isolated by affinity chromatography using antigen coupled to agarose beads.

**Clonality** Polyclonal

**Isotype** IgG

**General notes**Alexa Fluor<sup>®</sup> is a registered trademark of Molecular Probes, Inc, a Thermo Fisher Scientific

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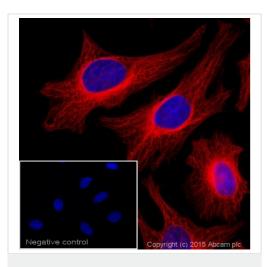
#### **Applications**

The Abpromise guarantee Our Abpromise guarantee covers the use of ab150115 in the following tested applications.

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

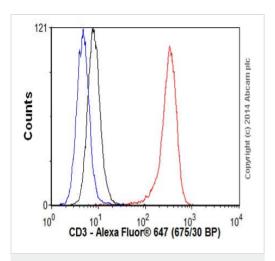
Application	Abreviews	Notes
IHC-Fr	<b>★★★★</b> <u>(2)</u>	Use at an assay dependent concentration.
ICC/IF	<b>★★★★★ (4)</b>	1/200 - 1/1000.
ELISA		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration.
Flow Cyt		1/2000 - 1/4000.  ab176103 - Mouse monoclonal lgG1 (Alexa Fluor® 647), is suitable for use as an isotype control to complement this secondary antibody.

#### **Images**

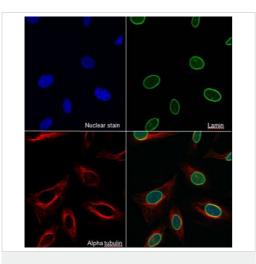


Immunocytochemistry/ Immunofluorescence - Goat Anti-Mouse IgG H&L (Alexa Fluor® 647) (ab150115) ICC/IF image of <u>ab7291</u> stained HeLa cells. The cells were 4% paraformaldehyde fixed (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then incubated in 1% BSA / 10% normal donkey serum / 0.3M glycine in 0.1% PBS-Tween for 1h to block non-specific protein-protein interactions. The cells were then incubated with the primary antibody (<u>ab7291</u>, 5 $\mu$ g/ml) overnight at +4°C. The secondary antibody (red) was ab150115 Alexa Fluor 647 goat anti-mouse IgG (H+L) used at 1 $\mu$ g/ml for 1h.DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43 $\mu$ M.

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



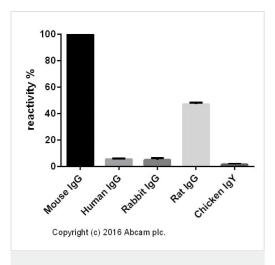
Flow Cytometry - Goat Anti-Mouse IgG H&L (Alexa Fluor® 647) (ab150115)



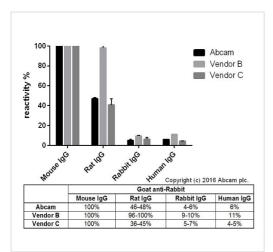
Immunocytochemistry/ Immunofluorescence - Goat
Anti-Mouse IgG H&L (Alexa Fluor® 647) (ab150115)

Overlay histogram showing Jurkat cells stained with <u>ab8090</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 goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (<u>ab8090</u>, 0.1 $\mu$ g/1x10<sup>6</sup> cells) for 30 min at 22°C. The secondary antibody Goat anti-mouse lgG H&L (Alexa Fluor<sup>®</sup> 647) (ab150115) was used at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse lgG2a [ICIGG2A] (<u>ab91361</u>, 0.1 $\mu$ g/1x10<sup>6</sup> 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 solid-state 25mW red diode laser (635nm) and 675/30 bandpass filter.

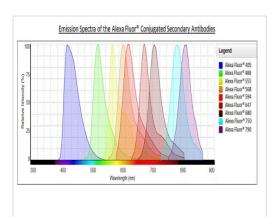
The cells were 100% methanol fixed (5 min) and then incubated in 1% BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab7291, 1µg/ml) and (ab16048, 1µg/ml) overnight at +4°C. The secondary antibodies were ab150115 Alexa Fluor<sup>®</sup> 647 (red) goat anti-mouse IgG (H+L) used at 2µg/ml for 1h and ab150077 Alexa Fluor<sup>®</sup> 488 (green) goat anti-rabbit IgG (H+L) used at 2µg/ml for 1h. DAPI was used to stain the cell nuclei.



ELISA - Goat Anti-Mouse IgG H&L (Alexa Fluor® 647) (ab150115)



ELISA - Goat Anti-Mouse IgG H&L (Alexa Fluor® 647) (ab150115)



Alexa Fluor® - Goat Anti-Mouse IgG H&L (Alexa Fluor® 647) (ab150115)

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  $\mu$ g/ml (50 $\mu$ 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  $\mu$ g/ml and gradually diluted 1/4 (50  $\mu$ 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  $\mu$ l/well), followed by incubation for 1h at RT.

Fot 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.

This data was developed using the unconjugated antibody (ab182017).

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  $\mu$ g/ml (50 $\mu$ 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  $\mu$ g/ml and gradually diluted 1/4 (50  $\mu$ 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  $\mu$ l/well), followed by incubation for 1h at RT. This data is from a representative dilution.

This data was developed using the unconjugated antibody (ab182017).

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