# abcam

# Product datasheet

# Goat Anti-Mouse IgG H&L (Alexa Fluor® 568) ab175473

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#### Overview

Product name Goat Anti-Mouse IgG H&L (Alexa Fluor® 568)

Host species Goat

Target species Mouse

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

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

**Conjugation** Alexa Fluor® 568. Ex: 578nm, Em: 603nm

#### **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**

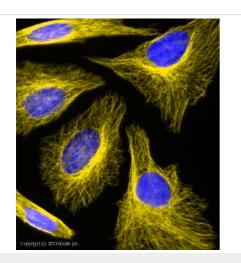
The Abpromise guarantee Our Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab175473 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.
Flow Cyt		1/2000.
ICC/IF	<b>★★★★★ (1)</b>	1/200 - 1/1000.
ELISA		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration.

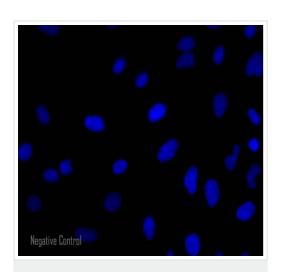
## **Images**



Immunocytochemistry/ Immunofluorescence - Goat
Anti-Mouse IgG H&L (Alexa Fluor® 568) (ab175473)

ICC/IF image of <u>ab7291</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>ab7291</u>, 1 $\mu$ g/ml) overnight at +4°C. The secondary antibody (yellow) was ab175473 Alexa Fluor<sup>®</sup> 568 Goat anti-Mouse IgG (H&L) used at 2 $\mu$ g/ml for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43 $\mu$ M.

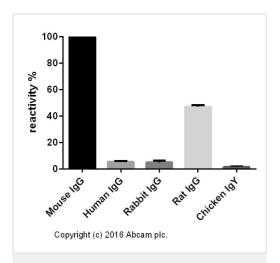
Alexa Fluor<sup>®</sup> 568 is pseudocolored in yellow.



Immunocytochemistry/ Immunofluorescence - Goat
Anti-Mouse IgG H&L (Alexa Fluor® 568) (ab175473)

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 ab175473 Alexa Fluor  $^{\mbox{\scriptsize $6$}}$  568 Goat anti-Mouse IgG (H&L) used at 2µg/ml for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.

Alexa Fluor® 568 is pseudocolored in yellow.

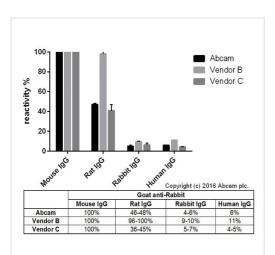


ELISA - Goat Anti-Mouse IgG H&L (Alexa Fluor® 568) (ab175473)

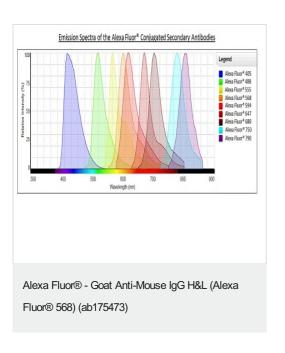
Cross-reactivity of the polyclonal secondary antibody <u>ab182017</u> was tested using a sandwich ELISA approach. The wells were coated with the indicated IgG standards at 1  $\mu$ g/ml (50 $\mu$ l/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. <u>ab182017</u> 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).



ELISA - Goat Anti-Mouse IgG H&L (Alexa Fluor® 568) (ab175473)



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).

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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