

### Goat Anti-Rabbit IgG H&L (Alexa Fluor® 594) ab150080

★★★★★ [3 Abreviews](#) [592 References](#) [画像数 6](#)

#### 製品の概要

製品名	Goat Anti-Rabbit IgG H&L (Alexa Fluor® 594)
由来種	Goat
ターゲット生物種	Rabbit
アプリケーション	<b>適用あり:</b> IHC-Fr, ICC/IF, ELISA, IHC-P, Flow Cyt
免疫原	Other Immunogen Type corresponding to Rabbit IgG.
標識	Alexa Fluor® 594. Ex: 590nm, Em: 617nm

#### 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.
バッファー	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)	The 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	IgG
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## アプリケーション

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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。

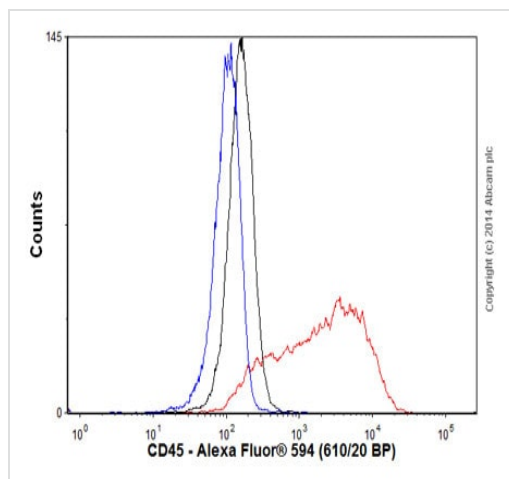
アプリケーション	Abreviews	特記事項
IHC-Fr	★★★★★ (1)	Use at an assay dependent concentration.
ICC/IF		1/200 - 1/1000.
ELISA		Use at an assay dependent concentration. Use at an assay dependent dilution
IHC-P	★★★★★ (1)	Use at an assay dependent concentration. Use at an assay dependent dilution
Flow Cyt		1/2000 - 1/4000.

## 画像



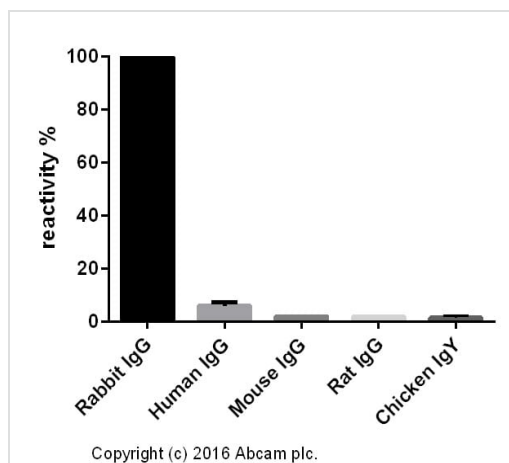
ICC/IF image of **ab6046** stained HeLa cells. The cells were 4% formaldehyde fixed (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then incubated in 1%BSA / 10% normal donkey serum / 0.3M glycine in 0.1% PBS-Tween for 1h to block non-specific protein-protein interactions. The cells were then incubated with the primary antibody (**ab6046**, 5µg/ml) overnight at +4°C. The secondary antibody (orange) was ab150080 Alexa Fluor® 4594 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.

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



Flow Cytometry - Goat Anti-Rabbit IgG H&L (Alexa Fluor® 594) (ab150080)

Overlay histogram showing Jurkat cells stained with **ab40763** (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 (**ab40763**, 1/1000 dilution) for 30 min at 22°C. The secondary antibody Goat anti-rabbit IgG H&L (Alexa Fluor® 594) (ab150080) was used at 1/4000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit IgG (monoclonal) (**ab172730**, 0.1µg/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 561nm laser and 610/20 bandpass filter.

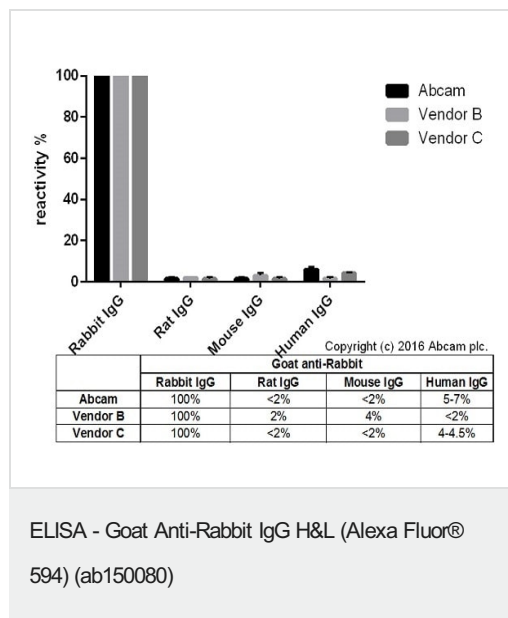


ELISA - Goat Anti-Rabbit IgG H&L (Alexa Fluor® 594) (ab150080)

Cross-reactivity of the polyclonal secondary antibody **ab182016** was tested using a sandwich ELISA approach. The wells were coated with the indicated IgG standards at 1 µg/ml (50 µl/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. **ab182016** was then added starting at 1 µg/ml and gradually diluted 1/4 (50 µl/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 µl/well), followed by incubation for 1h at RT.

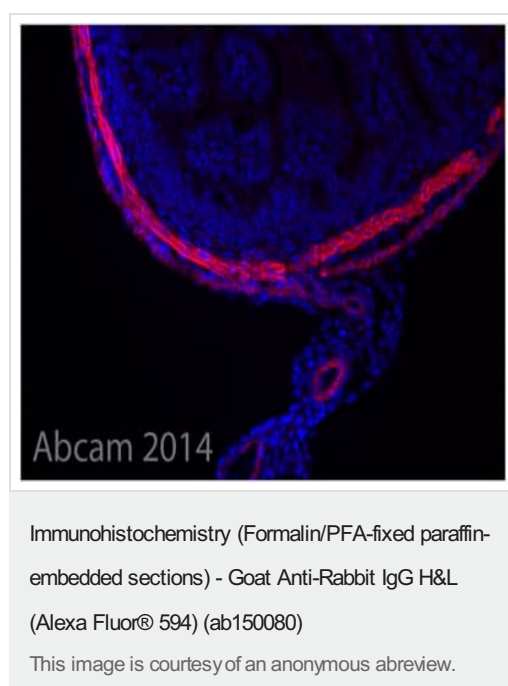
**For the batch tested, **ab182016** showed a cross-reactivity of 5-7% towards Human IgG and below 2% towards Mouse IgG, Rat IgG and Chicken IgY.**

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

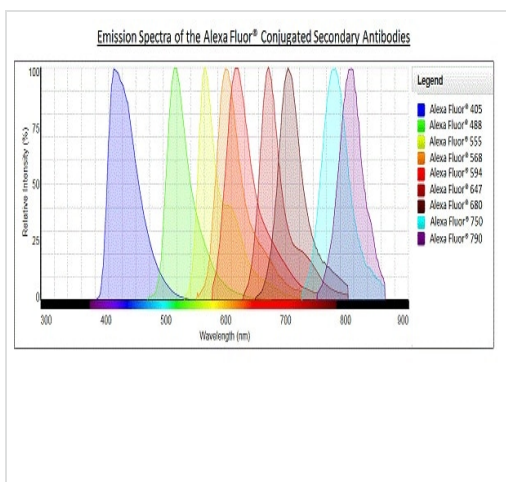


Cross-reactivity of Goat anti-Rabbit IgG H&L ([ab182016](#)) and Goat anti-Rabbit IgG H&L obtained from two different vendors was tested using a sandwich ELISA approach. The wells were coated with the indicated IgG standards (Rabbit, Human, Mouse and Rat) at 1 µg/ml (50 µl/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. Secondary antibodies were then added starting at 1 µg/ml and gradually diluted 1/4 (50 µl/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 µl/well), followed by incubation for 1h at RT. This data is from a representative dilution.

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



IHC-P image of alpha smooth muscle actin ([ab5694](#)) staining E16.5 mouse embryo gut. Paraformaldehyde fixed and paraffin embedded E16.5 mouse embryo gut sections were dewaxed and rehydrated before antigen retrieval (4 mins in a pressure cooker in 10mM Tris/0.4mM EDTA buffer pH 9.5). They were then incubated in 50mM NH4Cl for 30 minutes and washed/blocked in 3x 10 minute washes of PBS containing 1% BSA + 0.2% gelatine and 0.05% saponin. Sections were incubated overnight with a primary antibody against alpha smooth muscle actin ([ab5694](#)), diluted 1/250 in PBS containing 0.1% BSA and 0.3% triton. After 3 x 10 minute washes in of PBS containing 0.1% BSA, 0.2% gelatine and 0.05% saponin, the sections were incubated for 1 hr in the secondary antibody (ab150080, diluted 1/400, shown in red) and then the 3 washes repeated. Sections were mounted in Vectashield with DAPI (blue).



Alexa Fluor® - Goat Anti-Rabbit IgG H&L (Alexa Fluor® 594) (ab150080)

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