abcam

Product datasheet

Goat Anti-Rabbit IgG H&L (Alexa Fluor® 647) ab150079

★★★★★ 1 Abreviews 403 References 4 图像

概述

产品名称 山羊抗兔lgG H&L (Alexa Fluor® 647)

宿主 Goat **靶标种属** Rabbit

经测试应用 适用于: IHC-Fr, ICC/IF, ELISA, IHC-P, Flow Cyt

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

偶联物 Alexa Fluor® 647. Ex: 652nm, Em: 668nm

性能

形式 Liquid

存放说明 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.

存储溶液 Preservative: 0.02% Sodium azide

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

纯**度** Immunogen affinity purified

纯**化**说明 This antibody was isolated by affinity chromatography using antigen coupled to agarose beads.

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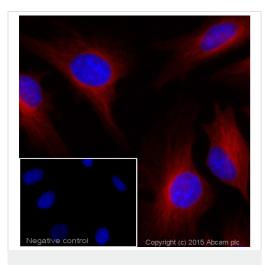
应用

The Abpromise guarantee Abpromise™承诺保证使用ab150079于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
IHC-Fr	★★★ ☆☆ <u>(1)</u>	Use at an assay dependent concentration.
ICC/IF		1/200 - 1/1000.
ELISA		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration.
Flow Cyt		1/2000.

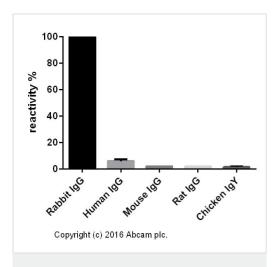
图片



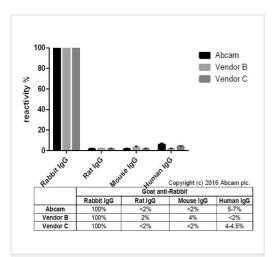
Immunocytochemistry/ Immunofluorescence - Goat
Anti-Rabbit IgG H&L (Alexa Fluor® 647) (ab150079)

ICC/IF image of <u>ab6046</u> in 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>ab6046</u>, 2 μ g/ml) overnight at +4°C. The secondary antibody ab150079 (shown in red) was used at 1 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.



ELISA - Goat Anti-Rabbit IgG H&L (Alexa Fluor® 647) (ab150079)



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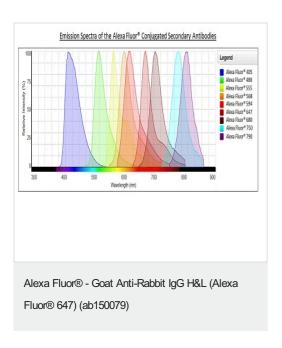
Cross-reactivity of the polyclonal secondary antibody <u>ab182016</u> was tested using a sandwich ELISA approach. The wells were coated with the indicated IgG standards at 1 μ g/ml (50 μ l/well) and incubated overnight at 4°C, followed by a 5% BSA blocking step for 2h at RT. <u>ab182016</u> was then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

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).



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