abcam

Product datasheet

Donkey Anti-Rabbit IgG H&L (Alexa Fluor® 594) ab150076

143 References 3 图像

概述

宿主 Donkey **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® 594. Ex: 590nm, Em: 617nm

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

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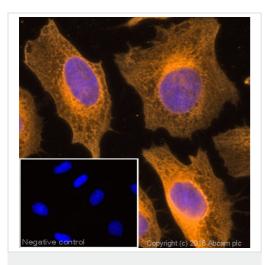
应用

The Abpromise guarantee Abpromise™承诺保证使用ab150076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		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		Use at an assay dependent concentration. Use at an assay dependent dilution
Flow Cyt		1/2000.

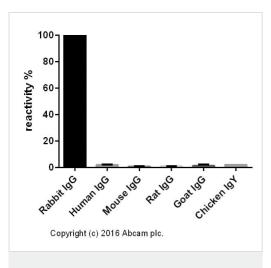
图片



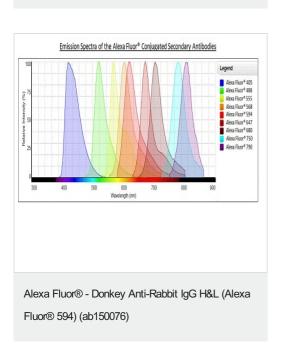
Immunocytochemistry/ Immunofluorescence Donkey Anti-Rabbit IgG H&L (Alexa Fluor® 594)
(ab150076)

ICC/IF image of <u>ab6046</u> 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 goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to block nonspecific 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 (orange) was ab150076 Alexa Fluor® 594 donkey 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.

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



ELISA - Donkey Anti-Rabbit IgG H&L (Alexa Fluor® 594) (ab150076)



Cross-reactivity of the polyclonal secondary antibody <u>ab182020</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>ab182020</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 Goat anti-Donkey IgG H&L (HRP) (<u>ab6988</u>) was used at 1/20,000 dilution (50 μ l/well), followed by incubation for 1h at RT.

For the batch tested, <u>ab182020</u> showed a cross-reactivity below 2% towards human IgG, mouse IgG, rat IgG, goat IgG and chicken IgY.

This data was developed using the unconjugated antibody (ab182020).

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