abcam

Product datasheet

Donkey Anti-Goat IgG H&L (Alexa Fluor® 647) ab150131

89 References 2 Images

Overview

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

Host species Donkey

Target species Goat

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

Immunogen Full length protein corresponding to Goat lgG.

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

Clonality Polyclonal

Isotype IgG

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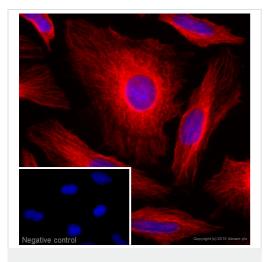
Applications

The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab150131 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.
ICC/IF		1/200 - 1/1000.
IHC-P		Use at an assay dependent concentration.
Flow Cyt		1/2000.
ELISA		Use at an assay dependent concentration.

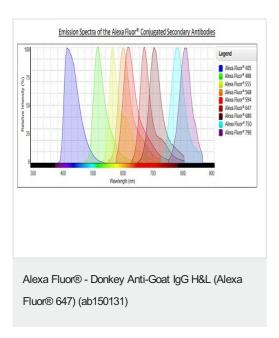
Images



Immunocytochemistry/ Immunofluorescence -Donkey Anti-Goat IgG H&L (Alexa Fluor® 647) (ab150131)

ICC/IF image of <u>ab7291</u> stained 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 / 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>ab7291</u>, 1 μ g/ml) overnight at +4°C. Ab98800, goat anti-mouse IgG, was then added as a secondary bridging antibody, at 1/250 dilution for 1h. Ab150131 Alexa Fluor® 647 donkey anti-goat IgG (H+L) (shown in red) was then used at 1 μ g/ml for 1h as a tertiary antibody. 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.



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