abcam

Product datasheet

Goat F(ab')2 Anti-Rat IgG H&L (Alexa Fluor® 555) preadsorbed ab150162

1 References 画像数 2

製品の概要

製品名 Goat F(ab')2 Anti-Rat lgG H&L (Alexa Fluor® 555) preadsorbed

由来種Goatターゲット生物種Rat

特異性 Reacts with the heavy and light chains of rat lgG, and with the light chains of rat lgM and lgA as

demonstrated by ELISA, FLISA, and flow cytometry. Goat F(ab')2 Anti-Rat IgG H&L (Alexa Fluor®

555) preadsorbed shows minimal cross reactivity with mouse immunoglobulins.

アプリケーション 適用あり: IHC-Fr, ICC/IF, ELISA, IHC-P, Flow Cyt

吸着処理血清

Human, Mouse more details

免疫原 Synthetic peptide corresponding to lgG. 標識 Alexa Fluor® 555. Ex: 555nm, Em: 565nm

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.

ארע"א Preservative: 0.02% Sodium azide

Constituents: 23% Glycerol (glycerin, glycerine), 68% PBS, 1% BSA

精製度 Immunogen affinity purified

特記事項(精製) Antiserum was solid phase adsorbed to ensure class specificity. Antiserum was cross adsorbed

using human and mouse immunosorbents to remove cross reactive antibodies. The antibody to rat lgG was isolated by affinity chromatography using antigen coupled to agarose beads. F(ab')2 fragments were generated using a pepsin digestion. Fc fragments and whole lgG molecules have

been removed.

ポリ/モノ ポリクローナル

アイソタイプ lgG

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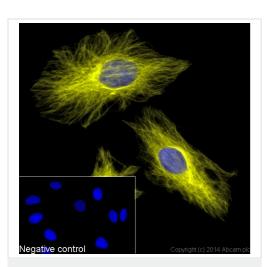
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アプリケーション

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アプリケーション	Abreviews	特記事項
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.

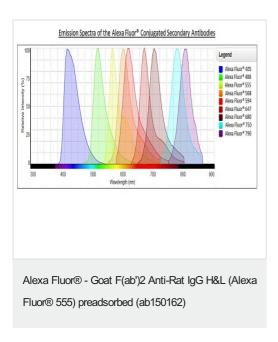
画像



Immunocytochemistry/ Immunofluorescence - Goat F(ab')2 Anti-Rat IgG H&L (Alexa Fluor® 555) preadsorbed (ab150162)

ICC/IF image of <u>ab6160</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>ab6160</u>, 1µg/ml) overnight at +4°C. The secondary antibody (yellow) was ab150162, Goat F(ab')2 Anti-Rat lgG H&L (Alexa Fluor® 555) preadsorbed, used at 1µg/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.



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