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Product datasheet

Goat Anti-Rat IgG H&L (Alexa Fluor® 594) ab150160

★★★★★ 1 Abreviews 89 References 画像数 3

製品の概要

製品名 Goat Anti-Rat lgG H&L (Alexa Fluor® 594)

由来種 Goat **ターゲット生物種** Rat

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

免疫原 Synthetic peptide corresponding to lgG.

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.

ארעדעד 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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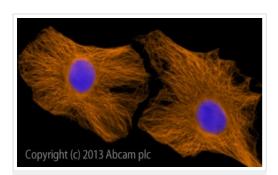
outlicensing@thermofisher.com.

アプリケーション

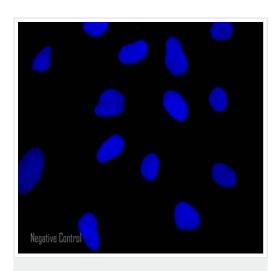
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アプリケーション	Abreviews	特記事項
IHC-Fr		Use at an assay dependent concentration.
ICC/IF		1/200 - 1/1000.
ELISA		Use at an assay dependent concentration. Use at an assay dependent dilution
IHC-P		Use at an assay dependent concentration. Use at an assay dependent dilution
Flow Cyt		1/2000.
IHC-FoFr		Use at an assay dependent concentration. PubMed: 28446880

画像

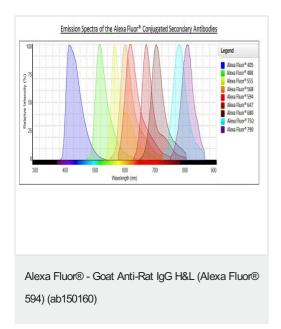


Immunocytochemistry/ Immunofluorescence - Goat Anti-Rat IgG H&L (Alexa Fluor® 594) (ab150160) ICC/IF image of <u>ab6160</u> stained HeLa cells. The cells were 100% methanol fixed (5 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (<u>ab6160</u>, 2 μ g/ml) overnight at +4°C. The secondary antibody (orange) was ab150160 Alexa Fluor® 594 goat anti-rat lgG (H+L) 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.



Immunocytochemistry/ Immunofluorescence - Goat Anti-Rat IgG H&L (Alexa Fluor® 594) (ab150160)

HeLa cells showing negative staining by ICC/IF using only secondary antibody. 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 secondary antibody (orange) was ab150160 Alexa Fluor® 594 goat anti-rat lgG (H+L) 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.



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