abcam

Product datasheet

Goat Anti-Mouse IgG H&L (DyLight® 488) preadsorbed ab96879

70 References 5 图像

概述

产品名称	**山羊抗小鼠IgG H&L (DyLight® 488)预吸附二抗**
宿主	Goat
靶标种属	Mouse
特异性	By immunoelectrophoresis and ELISA this antibody reacts specifically with Mouse IgG and with light chains common to other Mouse immunoglobulins. No antibody was detected against non immunoglobulin serum proteins. Reduced cross-reactivity to bovine, chicken, goat, horse, human, pig, rabbit and rat IgG was detected.
经测试应用	**适用于:** WB, IHC-P, ICC/IF, Flow Cyt
最小交叉反应	Chicken, Cow, Goat, Horse, Human, Pig, Rabbit, Rat more details
偶联物	DyLight® 488. Ex: 493nm, Em: 518nm

性能

形式	Liquid
存放说明	Shipped at 4°C. Store at +4°C.
存储溶液	pH: 6.8 Preservative: 0.09% Sodium azide Constituents: 0.2% BSA, PBS
纯度	Immunogen affinity purified
纯化说明	Antiserum was cross adsorbed using bovine, chicken, horse, human, pig, rabbit and rat immunosorbents to remove cross reactive antibodies. This antibody was isolated by affinity chromatography using antigen coupled to agarose beads and conjugated to DyLight® 488.
克隆	**多克隆**
同种型	IgG

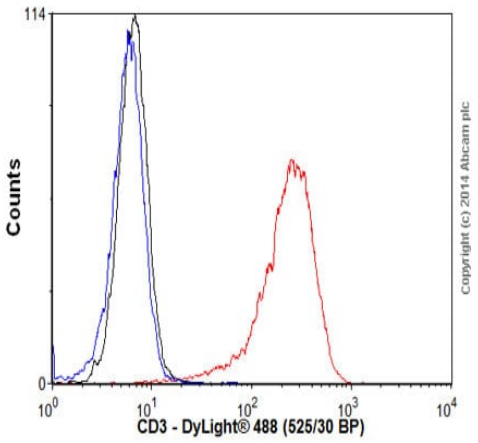
应用

The Abpromise guarantee	**Abpromise™承诺保证使用ab96879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。

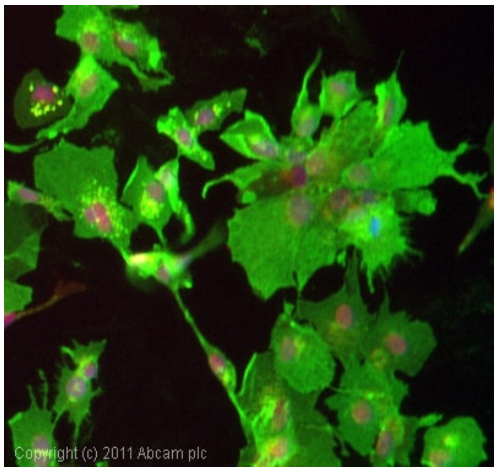
应用	Ab评论	说明
WB		1/1000 - 1/20000. Predicted molecular weight: 36 kDa. 5% non-fat dry milk in PBST or TBST is recommended for blocking and incubation of antibodies. BSA is not recommended.
IHC-P		1/50 - 1/500.
ICC/IF		1/50 - 1/500.
Flow Cyt		1/1000 - 1/2000.

图片



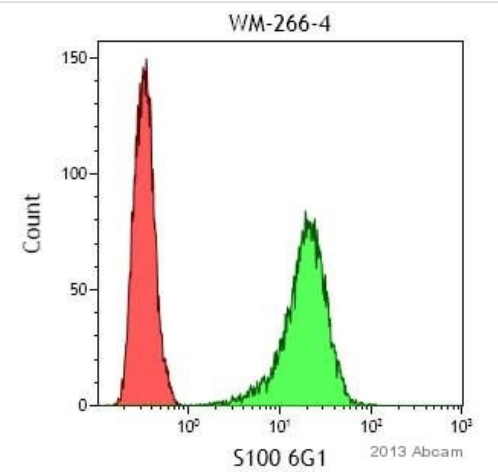
Flow Cytometry - Goat Anti-Mouse IgG H&L (DyLight® 488) preadsorbed (ab96879)

Overlay histogram showing Jurkat cells stained with **ab8090** (red line). The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (**ab8090**, 0.01µg/1x10^6 cells) for 30 min at 22°C. The secondary antibody Goat anti-mouse IgG H&L (DyLight® 488, pre-adsorbed) (ab96879) was used at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse IgG2a [ICIGG2A] (**ab91361**, 0.01µg/1x10^6 cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.



Immunocytochemistry/ Immunofluorescence - Goat Anti-Mouse IgG H&L (DyLight® 488) preadsorbed (ab96879)

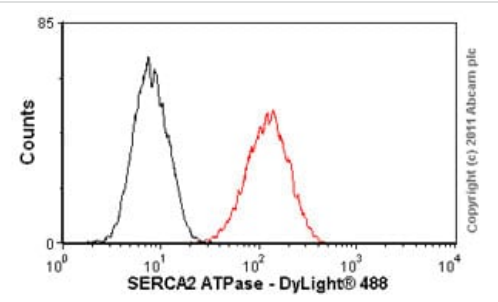
ICC/IF image of **ab40084** stained HepG2 cells. The cells were 4% PFA fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (**ab40084**, 5µg/ml) overnight at +4°C. The secondary antibody (green) was DyLight® 488 goat anti-mouse IgG - H&L, pre-adsorbed (ab96879) used at a 1/250 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.



Flow Cytometry - Goat Anti-Mouse IgG H&L (DyLight® 488) preadsorbed (ab96879)

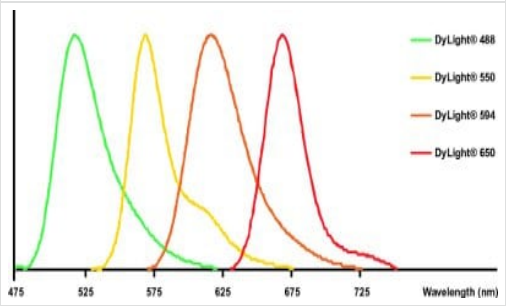
This image is courtesy of an anonymous Abreview.

ab85137 staining S100 in a human melanoma cell line by Flow Cytometry. The cells were harvested using EDTA and washed in PBS. The sample was incubated with the primary antibody (1/100 in PBS) for 15 minutes at room temperature. A DyLight® 488-conjugated goat anti-mouse IgG H&L (ab96879) (1/100) was used as the secondary antibody.



Flow Cytometry - Goat Anti-Mouse IgG H&L (DyLight® 488) preadsorbed (ab96879)

Overlay histogram showing HepG2 (Human liver hepatocellular carcinoma cell line) cells stained with **ab2861** (red line).
The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (**ab2861**, 1µg/1x106 cells) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-mouse IgG (H+L) ab96879 at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse IgG2a [ICIGG2A] (**ab91361**, 1µg/1x106 cells) used under the same conditions. Acquisition of >5,000 events was performed.



Goat Anti-Mouse IgG H&L (DyLight® 488) preadsorbed (ab96879)

Emission spectra of DyLight® fluorochromes available in our catalog.
Line colors represent the approximate visible colors of the wavelength maxima.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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