

Product datasheet

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

[70 References](#) [5 Images](#)

Overview

Product name	Goat Anti-Mouse IgG H&L (DyLight® 488) preadsorbed
Host species	Goat
Target species	Mouse
Specificity	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.
Tested applications	Suitable for: WB, IHC-P, ICC/IF, Flow Cyt
Minimal cross-reactivity	Chicken, Cow, Goat, Horse, Human, Pig, Rabbit, Rat more details
Conjugation	DyLight® 488. Ex: 493nm, Em: 518nm

Properties

Form	Liquid
Storage instructions	Shipped at 4°C. Store at +4°C.
Storage buffer	pH: 6.8 Preservative: 0.09% Sodium azide Constituents: 0.2% BSA, PBS
Purity	Immunogen affinity purified
Purification notes	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.
Clonality	Polyclonal
Isotype	IgG

Applications

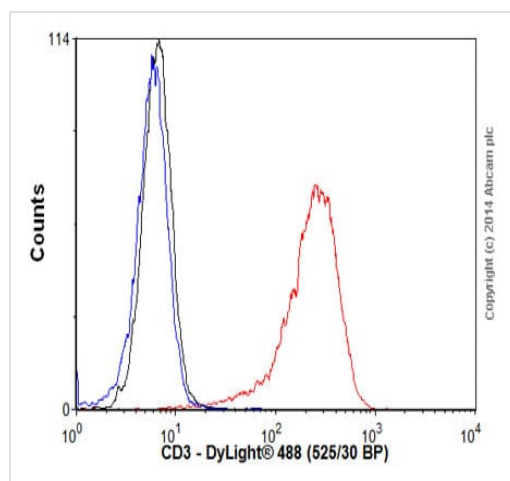
The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab96879 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

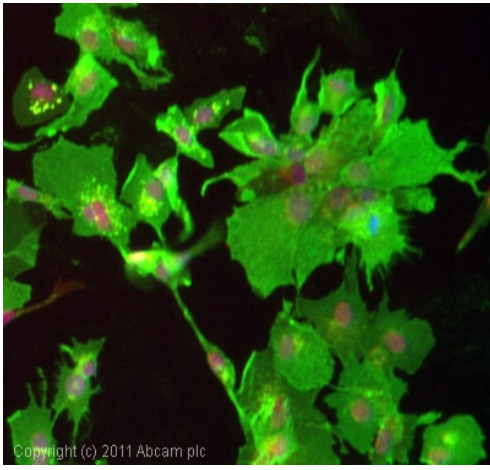
Application	Abreviews	Notes
WB		1/1000 - 1/20000. Predicted molecular weight: 36 kDa. 5% non-fat dry milk in PBST or TBST is recommended for blocking and incubation of antibodies. BSA is not recommended.
IHC-P		1/50 - 1/500.
ICC/IF		1/50 - 1/500.
Flow Cyt		1/1000 - 1/2000.

Images



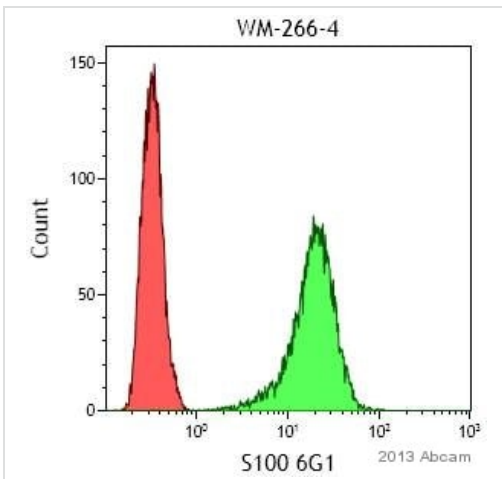
Flow Cytometry - Goat Anti-Mouse IgG H&L
(DyLight® 488) preadsorbed (ab96879)

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/1x10⁶ cells) for 30 min at 22°C. The secondary antibody Goat anti-mouse IgG H&L (DyLight® 488, preadsorbed) (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/1x10⁶ 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)

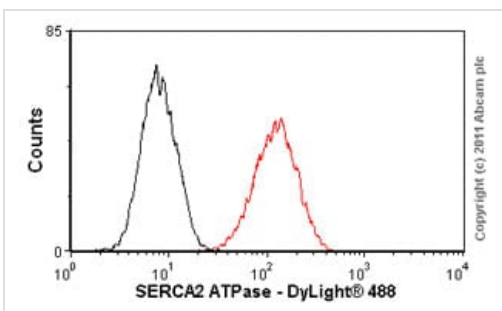
ICC/IF image of **ab40084** stained HepG2 cells. The cells were 4% PFA fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (**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)

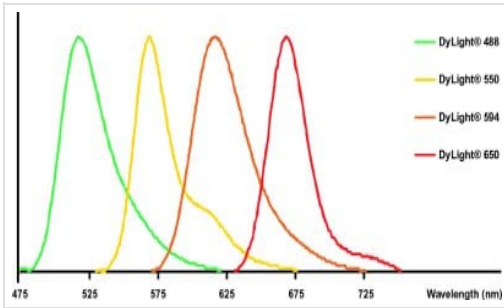
This image is courtesy of an anonymous Abreview.

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.



Emission spectra of DyLight[®] fluorochromes available in our catalog.

Line colors represent the approximate visible colors of the wavelength maxima.

Goat Anti-Mouse IgG H&L (DyLight[®] 488)
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