Goat anti-Mouse IgG (H+L) Cross-Adsorbed Secondary Antibody, Rhodamine

Product Details

Size	1.5 mg
Species Reactivity	Mouse
Host/Isotype	Goat / IgG
Class	Polyclonal
Туре	Secondary Antibody
Conjugate	Rhodamine
Excitation/Emission Max	573/591 nm
Form	Lyophilized
Concentration	1.5 mg/mL
Purification	Affinity chromatography
Storage buffer	PBS, pH 7.6, with 15mg/mL BSA
Contains	0.05% sodium azide
Storage conditions	4° C
RRID	AB_228303

Applications	Tested Dilution	Publications
Immunohistochemistry (IHC)	1:50 - 1:200	-
Immunocytochemistry (ICC/IF)	1.5 µg/mL	-
Flow Cytometry (Flow)	1:50 - 1:200	-
Immunoprecipitation (IP)	1:50 - 1:200	-

Product Specific Information

Concentration may vary slightly from lot-to-lot, see lot-specific datasheet for exact concentration.

Product # 31661 has been successfully used in Western blot, IF, ICC, IHC, IP and FACS applications.

Product # 31661 reacts with the heavy chains of mouse IgG and with the light chains common to most mouse immunoglobulins, but does not react against non-immunoglobulin serum proteins. The antibody has been tested by ELISA and /or solid-phase adsorbed to ensure minimal cross-reaction with human, bovine and horse serum proteins. However, this antibody may cross-react with immunoglobulins from other species.

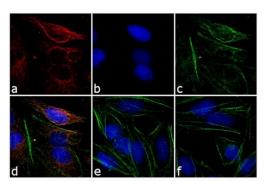
Store product protected from light at 4°C until opened. To extend the shelf-life of this product, add an equal volume of glycerol to make a final concentration of approximately 50% glycerol and store at -20°C. Rhodamine Amax= 550 nm; Emax= 570 nm. Fluorophore/Protein Absorbance Ratio: A550/A280 = 0.52 (lot-dependent).

Reconstitute with 1.1 mL of distilled water (1.5 mg/mL after restoration).

Country of Origin: USA

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Product Images For Goat anti-Mouse IgG (H+L) Cross-Adsorbed Secondary Antibody, Rhodamine



Mouse IgG (H+L) Cross-Adsorbed Secondary Antibody (31661) in ICC/IF Immunofluorescence analysis of Goat anti-Mouse IgG (H+L) Cross Adsorbed Secondary Antibody, Rhodamine conjugate was performed using MCF-7 cells stained with alpha Tubulin (23610501) Mouse Monoclonal Primary Antibody (Product # A11126). The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% Triton™ X-100 for 10 minutes, blocked with 1% BSA for 1 hour and labeled with Mouse primary antibody (1:250 dilution) for 3 hours at room temperature. Goat anti-Mouse IgG (H+L) Cross Adsorbed Secondary Antibody, Rhodamine conjugate (Product # 31661) was used at a concentration of 2 µg/mL in phosphate buffered saline containing 0.2 % BSA for 45 minutes at room temperature, for detection of alpha Tubulin in the cytoplasm (Panel a: red). Nuclei (Panel b: blue) were stained with DAPI in SlowFade® Gold Antifade Mountant (Product # S36938). F-actin was stained with Alexa Fluor® 488 Phalloidin (Product # A12379, 1:300) (Panel c: green). Panel d represents the composite image. No nonspecific staining was observed with the secondary antibody alone (panel f), or with an isotype control (panel e). The images were captured at 60X magnification.

□ 1 Reference

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Mechanism of restriction of normal and cystic fibrosis transmembrane conductance regulator-deficient human tracheal gland cells to adenovirus infection and ad-mediated gene transfer. Am J Respir Cell Mol Biol (2002)

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