



Goat anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor™ 635

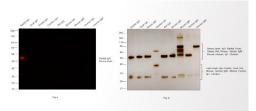
| Product Details | |
|----------------------------|--|
| Size | 1 mg |
| Species Reactivity | Rabbit |
| Host/Isotype | Goat / IgG |
| Class | Polyclonal |
| Туре | Secondary Antibody |
| Conjugate | Alexa Fluor™ 635 |
| Excitation/Emission Max | 633/648 nm |
| Immunogen | Gamma Immunoglobins Heavy and Light chains |
| Form | Liquid |
| Concentration | 2 mg/mL |
| Purification | purified |
| Storage buffer | PBS, pH 7.5 |
| Contains | 5mM sodium azide |
| Storage conditions | 4°C, store in dark |
| RRID | AB_2536187 |

| Applications | Tested Dilution | Publications |
|---|------------------|---------------|
| Western Blot (WB) | 1:5,000-1:10,000 | - |
| Immunohistochemistry (IHC) | 1-10 μg/mL | - |
| Immunohistochemistry (Frozen) (IHC (F)) | - | 0 Publication |
| Immunocytochemistry (ICC/IF) | 4 μg/mL | - |
| Miscellaneous PubMed (Misc) | - | 0 Publication |

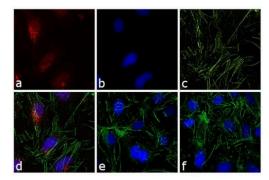
Product Specific Information

Product will be shipped at Room Temperature.

Product Images For Goat anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor™ 635



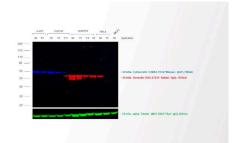
Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-31577) Specificity of secondary antibody was demonstrated by specific detection of the target immunoglobulin. Antibody specificity was demonstrated by specific detection of Rabbit IgG. A band at ~55 kDa corresponding to Rabbit IgG Heavy Chain was observed in Rabbit IgG but not in other species using Goat anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor[™] 635 (Product # A-31577) in Western Blot. {RE}



Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-31577) in ICC/IF

Immunofluorescence analysis of Goat anti-Rabbit IgG (H+L) Secondary Antibody, Alexa Fluor 635 was performed using HepG2 cells stained with alpha-1 antitrypsin Rabbit Polyclonal Primary Antibody (Product # PA5-16661). The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with 0.1% TritonTM X-100 for 10 minutes, blocked with 1% BSA for 1 hour and labeled with Rabbit primary antibody (1:250 dilution) 3 hours at room temperature. Goat anti-Rabbit IgG (H+L) Secondary Antibody, Alexa Fluor 635 (Product # A-31577) was used at a concentration of 4 µg/mL in phosphate buffered saline containing 0.2 % BSA for 45 minutes at room temperature, for detection of alpha-1 antitrypsin 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.

Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody (A-31577) in WB



Multiplexed fluorescent western blot was performed using Goat anti-Rabbit IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor 635 (Product # A-31577). Whole cell extracts of A-431 (Lane 1, 2), HaCaT (Lane 3, 4, 5), SH-SY5Y (Lane 6, 7, 8), HeLa (Lane 9, 10) and MCF7 (Lane 11) were electrophoresed usingNuPAGE™ 4-12% Bis-Tris Protein Gel (Product # NP03222BOX). Resolved proteins were transferred onto anitrocellulose membrane (Product # IB23001) byiBlot® 2 Dry BlottingSystem (Product # IB21001). The blot was probed with Cytokeratin 5 Monoclonal Antibody (3E2F1) (Product # MA5-15347), Vimentin Polyclonal Antibody (Product # PA5-27231) and alpha Tubulin Monoclonal Antibody (YL1/2) (Product # MA1-80017). Secondary antibodies (Product # A-11357, 1:20,000), (Product # A-31577, 1:10,000) and (Product # A48269, 1:10,000) were used for detection of Cytokeratin 5, Vimentin and alpha Tubulin respectively. Fluorescent detection was performed usingiBright™FL1500 (Product # A44115). The anti-rabbit secondary antibody (Product # A-31577) specifically detects the rabbit primary antibody.

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□ 48 References

Chronic administration of prebiotics and probiotics prevent pathophysiological hallmarks of Alzheimer's disease in the cortex of APP/PS1 mice. Front Pharmacol (2025)

Chronic administration of prebiotics and probiotics ameliorates pathophysiological hallmarks of Alzheimer's disease in a APP/PS1 transgenic mouse model. Front Pharmacol (2024)

Increased Homer Activity and NMJ Localization in the Vestibular Lesion het-/- Mouse soleus Muscle. Int J Mol Sci (2024)

Optogenetic Determination of Dynamic and Cell-Type-Specific Inhibitory Reversal Potentials. J Neurosci (2024)

New orphan disease therapies from the proteome of industrial plasma processing waste- a treatment for aceruloplasminemia. Commun Biol (2024)

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