

Product datasheet for TA397870S

OriGene Technologies, Inc.

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Mouse IgG (H&L) Antibody ATTO 488 Conjugated Pre-Adsorbed

Product data:

Product Type: Secondary Antibodies

Product Name: Mouse IgG (H&L) Antibody ATTO 488 Conjugated Pre-Adsorbed

Applications: IF, WB

Recommended Dilution: WB: >1:10,000

IF: >1:5,000

FLISA: >1:20,000

Reactivity: Mouse **Host:** Goat

Immunogen: Mouse IgG whole molecule

Formulation: 0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2

Reconstitution Method: Restore with deionized water (or equivalent) - Reconstitution Volume: 100 μL

Concentration: 1.0 mg/mL - lot specific

Conjugation: ATTO 488

Storage: Store vial at 4° C prior to restoration. For extended storage aliquot contents and freeze at -

20° C or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. This product is stable for several weeks at 4° C as

an undiluted liquid. Dilute only prior to immediate use.

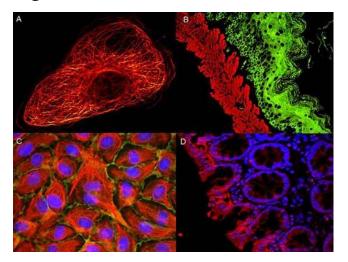
Note: Anti-Mouse IgG ATTO 488 Antibody has been tested by dot blot and western blot. Anti-Mouse

IgG (H&L) conjugated by ATTO 488 is designed for STED microscopy, FRET,

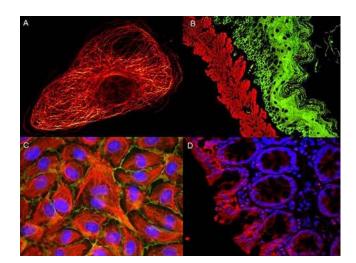
immunofluorescence microscopy, fluorescence based plate assays (FLISA) and fluorescent western blotting. This product is also suitable for multiplex analysis, including multicolor imaging, utilizing various commercial platforms. The emission spectra for this ATTO conjugate matches the principle output wavelengths of most common fluorescence instrumentation.



Product images:



ATTO ® dyes can be used for multicolor immunofluorescent detection with low background and high signal. Examples shown are: A. Tubulin in PtK2- male Rat Kangaroo Kidney Epithelial Cells was detected using ATTO 532 labeled secondary antibody. B. Muscle alphaactin was stained with a mouse primary antibody and ATTO 488 anti-mouse IgG (green) while Cytokeratin was stained with polyclonal rabbit anti-cytokeratin and ATTO 647N anti-rabbit IgG (red). C. HUVEC (Human umbilical vein endothelial cells were stained with anti- Vimentin-ATTO 532 (green), anti-E-Cadherin-ATTO 655 (red) and DAPI (blue). D. Rat colon sections were stained with Anti-Aquaporin 3-ATTO 594 antibody. Hoechst 33342 (blue) is used as counterstain. Images provided courtesy of Dr. Jörg Reichwein, ATTO-TEC GmbH



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