



Goat anti-Mouse IgG2a Cross-Adsorbed Secondary Antibody, Alexa Fluor™ 555

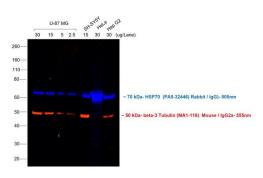
Product Details	
Size	500 μg
Species Reactivity	Mouse
Host/Isotype	Goat / IgG
Class	Polyclonal
Туре	Secondary Antibody
Conjugate	Alexa Fluor™ 555
Excitation/Emission Max	553/568 nm
Immunogen	Mouse IgG2a
Form	liquid
Concentration	2 mg/mL
Purification	purified
Storage buffer	PBS, pH 7.5
Contains	5mM sodium azide
Storage conditions	4° C, store in dark
RRID	AB_2535776

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

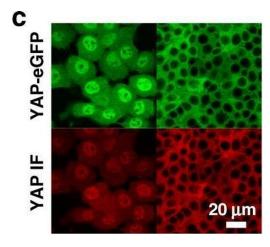
Product Specific Information

Product will be shipped at Room Temperature.

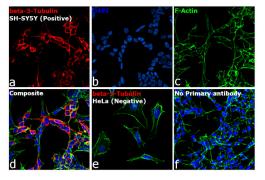
Product Images For Goat anti-Mouse IgG2a Cross-Adsorbed Secondary Antibody, Alexa Fluor™ 555



Mouse IgG2a Cross-Adsorbed Secondary Antibody (A-21137) in WB Multiplexed fluorescent western blot was performed using Goat anti-Mouse IgG2a Cross-Adsorbed Secondary Antibody, Alexa Fluor™ 555 (Product # A-21137). Whole cell extracts of U-87 MG (Lane 1, 2, 3, 4), SH-SY5Y (Lane 5), HeLa (Lane 6) and Hep G2 (Lane 7) 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 beta-3 Tubulin mouse IgG2a Monoclonal Antibody (Product # MA1-118) and HSP70 Polyclonal Antibody (Product # PA5-32446). Secondary antibodies (Product # A-21137, 1: 10,000) and (Product # A32808,1:20,000) were used for detection of beta-3 Tubulin and HSP70 respectively. Fluorescent detection was performed usingiBrightFL1500 (Product # A44115). The anti-mouse secondary antibody (Product # A-21137) specifically detects the mouse primary antibody but not the rabbit primary antibody.



Mouse IgG2a Cross-Adsorbed Secondary Antibody (A-21137) in ICC/IF Characterization of YAP and TEAD genome-knockin cell lines.a Cartoon of CRISPR-Cas9 based insertion of eGFP-P2A-puromycin cassette at 3' end of the YAP gene. b western blot against eGFP from MCF10AYAP-GFP-KI whole cell lysate. Single predominant band at the predicted YAP-eGFP fusion size 96 kDa. Repeated twice with similar results. c Correlation of the YAP nuclear/cytoplasmic ratio (N/C) signal for YAP-eGFP and anti-YAP immunofluorescence in sparse and dense cultures. Nsparse = 61 cells, Ndense = 155 cells from one experiment. Repeated twice with similar results. d Representative Immunofluorescence images of data from c. e MCF10AYAP-GFP-KI cells plated on soft (0.4 kPa) and stiff (60 kPa) acrylamide showing altered cell morphology and YAP localization. Cells were counterstained with the live-cell nuclear stain, SiR-DNA™. f Quantitative comparison of YAP-eGFP N/C on fibronectin coated acrylamide of varying stiffnesses. N0.4 kPa = 834, N1 kPa = 818, N6 kPa = 1386, N60 kPa = 647 cells. A one-sided Mann-Whitney U-test was used to test differences in YAP N/C. 0.4 kPa vs. 1 kPa: 5.8 x 1035, Ranksum = 570,253. 1 $kPa vs. 6 kPa: p = 5.3 \times 1097$, Ranksum = 600,652. 6 kPa vs. 60 kPa: P = 1.452x 1016, Ranksum = 1,308,747. Data compiled from two independent experiments for each condition. g Time-course of YAP N/C normalized to initial value following Latrunculin B treatment at 0.5 µg/mL (1... Image collected and cropped by CiteAb from the following publication (https://pubmed.ncbi.nlm.nih.gov /32917893), licensed under a CC BY license.



Mouse IgG2a Cross-Adsorbed Secondary Antibody (A-21137) in ICC/IF Immunofluorescence analysis of Goat anti-Mouse IgG2a Cross-Adsorbed Secondary Antibody, Alexa Fluor™ 555 (Product # A-21137) was performed using SH-SY5Y (positive model) and HeLa (negative model) cells stained with beta-3 Tubulin Monoclonal Antibody (Product # MA1-118). 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 2 µg/mL primary antibody for 3 hours at room temperature. Goat anti-Mouse IgG2a Cross-Adsorbed Secondary Antibody, Alexa Fluor™ 555 (Product # A-21137, 1:2000) in 0.1% BSA in PBS for 45 minutes at room temperature, was used for detection of beta-3 Tubulin in the cytoskeleton (Panel a: red). Nuclei (Panel b: blue) were stained with Hoechst33342 (Product # H1399). F-actin was stained with Alexa Fluor® 488 Phalloidin (Product # A12379, 1:300) (Panel c: green). Panel d represents the composite image. The specificity of the secondary antibody was proved by the absence of signal in HeLa (negative model for beta 3 Tubulin) due to no primary antibody binding (Panel e). Nonspecific staining was not observed with secondary antibody alone (panel f). The images were captured at 20X magnification.

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□ 108 References

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