Goat anti-Human IgM Secondary Antibody

Product Details	
Size	2 mg
Species Reactivity	Human
Host/Isotype	Goat / IgG
Class	Polyclonal
Type	Secondary Antibody
Conjugate	Unconjugated
Form	Liquid
Concentration	2.32 mg/mL
Purification	Antigen affinity chromatography
Storage buffer	PBS, pH 7.6
Contains	no preservative
Storage conditions	4° C
RRID	AB_228279

Applications	Tested Dilution	Publications
Western Blot (WB)	1:2000	-
Immunohistochemistry (IHC)	Assay-dependent	-
Immunocytochemistry (ICC/IF)	Assay-dependent	-
Flow Cytometry (Flow)	Assay-dependent	-
Immunoprecipitation (IP)	Assay-dependent	-

Product Specific Information

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

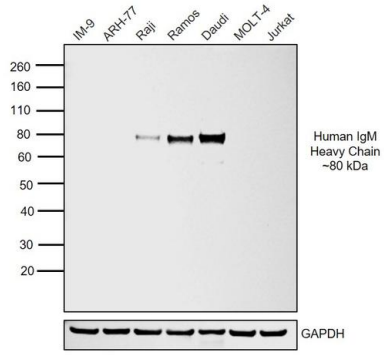
Product # 31136 has been successfully used in Western blot, IF, ICC, IHC, IP and FACS applications.

Product # 31136 reacts with the Fc5µ portion of human IgM heavy chain. This antibody does not react against normal human IgG or IgA, or against non-immunoglobulin serum proteins. However, this antibody may cross-react with IgM from other species.

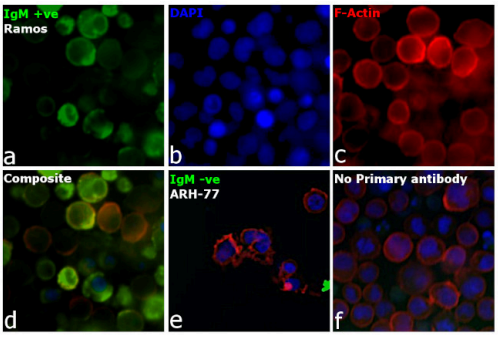
Store product at 4°C until opened. After opening, dilute only enough antibody for a single day's use. Store remainder at 4°C under sterile conditions.

Country of origin: USA

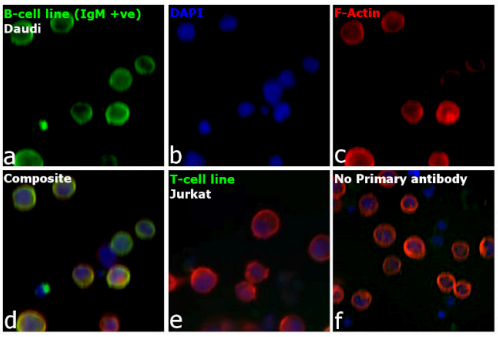
Product Images For Goat anti-Human IgM Secondary Antibody



Human IgM Secondary Antibody (31136) in WB
Western blot was performed using Goat anti-Human IgM Secondary Antibody (Product # 31136) and an ~80 kDa band corresponding to Human IgM Heavy Chain was observed in Raji, Ramos and Daudi but not in IM-9, ARH-77, MOLT-4 and Jurkat8203. Whole cell extracts (30 µg) of IM-9 (Lane 1), ARH-77 (Lane 2), Raji (Lane 3), Ramos (Lane 4), Daudi (Lane 5), MOLT-4 (Lane 6) and Jurkat (Lane 7) were electrophoresed using NuPAGE™ 4-12% Bis-Tris Protein Gel (Product # NP0322BOX). Resolved proteins were then transferred onto a nitrocellulose membrane (Product # IB23001) by iBlot® 2 Dry Blotting System (Product # IB21001). The blot was probed with Product # 31136 (1:2000 dilution) and detected by chemiluminescence with Rabbit anti-Goat IgG Heavy Chain Superclonal™ Recombinant Secondary Antibody, HRP Conjugate (Product # A27014, 1:20,000 dilution) using the iBright FL1500 (Product # A44115). Chemiluminescentdetection was performed using Novex® ECL Chemiluminescent Substrate Reagent Kit (Product # WP20005). Raji, Ramos and Daudi are known to express IgM whereas IM-9 and ARH-77 express IgG and are negative for IgM. MOLT-4 and Jurkat, being T-cell lines, do not express immunoglobulins (DOI:10.1002/eji.1830100305; 10.3791/3573; 10.1016/0022-1759(94)00286-6; PMID: 566614).



Human IgM Secondary Antibody (31136) in ICC/IF
Immunofluorescence analysis of Goat anti-Human IgM Secondary Antibody was performed using log phaseRamos cells (IgM producing B-cell line). The cells were fixed with 4% paraformaldehyde for 10 minutes, permeabilized with0.1% Triton™ X-100 for 15minutes, andblocked with 2% BSA for 1 hour at room temperature. The cells were labeled withGoat anti-Human IgM Secondary Antibody (Product # 31136) at 1:250 dilution in 0.1% BSA, incubated at 4degreecelsiusovernight and then with Donkey anti-Goat IgG (H+L) Highly Cross-Adsorbed Secondary Antibody, Alexa Fluor Plus 488 (Product # A32814) at a dilution of 1:2000 dilution for 45 minutes at room temperature (Panel a: green). Nuclei(Panel b: blue) were stained with Hoechst 33342 (Product # H1399). F-actin (Panel c: red) was stained with Alexa Fluor™ Plus 647 Phalloidin (Product #A30107, 1:2000 dilution). Panel d represents the merged image showing cytoplasmic (plasma membraneandgolgi-body like) localization. Panel e represents ARH-77 cells (IgM non-producing B-cell line) which is a negative model forIgM expression. Panel f represents control cells with no primary antibody to assess background. The images werecaptured at 40X magnification inCellInsightCX7 LZR High-Content Screening (HCS) Platform (Product # CX7A1110LZR) (DOI:10.1002/eji.1830100305; 10.3791/3573; 10.1016/0022-1759(94)00286-6; PMID: 566614).



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1 Reference

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