



Goat anti-Rabbit IgG (H+L) Cross-Adsorbed Secondary Antibody, **HRP**

Product Details		
Size	1.5 mL	
Species Reactivity	Rabbit	
Host/Isotype	Goat / IgG	
Class	Polyclonal	
Туре	Secondary Antibody	
Conjugate	HRP	
Form	Lyophilized	
Concentration	0.8 mg/mL	
Purification	Affinity chromatography	
Storage buffer	PBS, pH 7.6, with 15mg/mL BSA	
Contains	no preservative	
Storage conditions	4° C	
RRID	AB_228338	

Applications	Tested Dilution	Publications
Western Blot (WB)	1:10,000-1:200,000	0 Publication
Immunohistochemistry (IHC)	1:500-1:5,000	0 Publication
Immunocytochemistry (ICC/IF)	1:500-1:5,000	-
Miscellaneous PubMed (Misc)	-	0 Publication

Product Specific Information

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

This antibody has been successfully used in Western blot, and ICC applications.

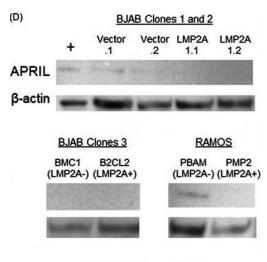
Antibody Specificity: This antibody reacts with the heavy chains of rabbit IgG and with the light chains common to most rabbit immunoglobulins. No antibody was detected against non-immunoglobulin serum proteins. The product has been tested by ELISA and/or solid-phase adsorbed to ensure minimal cross-reaction with human serum proteins. However, this antibody may cross-react with immunoglobulins from other species. This antibody may cross-react with SuperBlock® Blocking Buffers.

Restoration and Storage: Store product at 4°C until opened. Restore with 1.5 mL distilled water (0.8 mg/mL after restoration). Centrifuge product if it is not completely clear after standing for 1-2 hours at room temperature. To judge clarity, draw product into a pasteur pipette. Product may be stored for several weeks at 4°C as undiluted liquid. After dilution, do not use for more than one day.

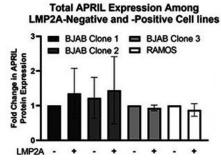
To extend the shelf-life of this product, add an equal volume of glycerol to make a final concentration of approximately 50% glycerol and store at -20°C.

Country of Origin: USA

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

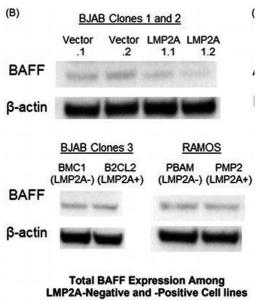


Rabbit IgG (H+L) Cross-Adsorbed Secondary Antibody (31462) in WB LMP2A does not affect the production BAFF or APRIL levels in B cell lymphoma cells. LMP2A-negative (BJAB Vector.1, BJAB Vector.2, BJAB BCM1, RAMOS-PBAM) and LMP2A-positive (BJAB LMP2A1.1, BJAB LMP2A1.2, BJAB B2CL2, RAMOS-PMP2) B cell lines were incubated for 48 h before supernatants were isolated and analyzed by ELISA for soluble BAFF (A) or APRIL (C) protein lysates were isolated from LMP2A-negative and -positive B cell lines and Western Blot was used to determine total BAFF (B) or APRIL (D) Experiments are either a combination of three experiments for (A, C) or representative of two experiments for (B, D) and a bar graph demonstrating the results of combining the representative Western blots. All data was analyzed statistically as described in the Section 2 and was found to not reach a p < .05. BAFF, B cell Activating Factor of the tumor necrosis factor family; LMP2A, Latent Membrane Protein 2A. Image collected and cropped by CiteAb from the following publication (https://pubmed.ncbi.nlm.nih.gov/36301035), licensed under a CC BY license.



HepG2 Jurkat 160 110 80 -60 -50 40 CacyBP 30 ~30 kDa 20 15 GAPDH 1:10,000 1:100,000 1:200,000 (Fig. 1) (Fig. 2) (Fig. 3)

Rabbit IgG (H+L) Cross-Adsorbed Secondary Antibody (31462) in WB
Western blot analysis was performed on whole cell extracts (30 µg lysate) of Hep
G2 (Lane 1) and Jurkat (Lane 2). The blots were probed with Anti-CacyBP Rabbit
Polyclonal Antibody (Product # 720326, 1 µg/mL) and detected by
chemiluminescence using Goat anti-Rabbit IgG (H+L) Cross-Adsorbed
Secondary Antibody, HRP (Product # 31462) at dilutions 1:10,000 (Fig. 1), 1:
100,000 (Fig. 2) and 1:200,000 (Fig. 3). A 30 kDa band corresponding to CacyBp
was observed. Known quantity of protein samples were electrophoresed using
Novex® NuPAGE®12 % Bis-Tris gel (Product # NP0342BOX), XCell SureLock™
Electrophoresis System (Product # El0002) and Novex® Sharp Pre-Stained
Protein Standard (Product # LC5800). Resolved proteins were then transferred
onto a nitrocellulose membrane with iBlot® 2 Dry Blotting System (Product #
IB21001). The membrane was probed with the relevant primary and secondary
antibody after blocking with 5 % skimmed milk. Chemiluminescent detection was
performed using Pierce™ ECL Western Blotting Substrate (Product # 32106).



■ BJAB Clone 1 ■ BJAB Clone 3 ■ BJAB Clone 2 ■ RAMOS Rabbit IgG (H+L) Cross-Adsorbed Secondary Antibody (31462) in WB LMP2A does not affect the production BAFF or APRIL levels in B cell lymphoma cells. LMP2A-negative (BJAB Vector.1, BJAB Vector.2, BJAB BCM1, RAMOS-PBAM) and LMP2A-positive (BJAB LMP2A1.1, BJAB LMP2A1.2, BJAB B2CL2, RAMOS-PMP2) B cell lines were incubated for 48 h before supernatants were isolated and analyzed by ELISA for soluble BAFF (A) or APRIL (C) protein lysates were isolated from LMP2A-negative and -positive B cell lines and Western Blot was used to determine total BAFF (B) or APRIL (D) Experiments are either a combination of three experiments for (A, C) or representative of two experiments for (B, D) and a bar graph demonstrating the results of combining the representative Western blots. All data was analyzed statistically as described in the Section 2 and was found to not reach a p < .05. BAFF, B cell Activating Factor of the tumor necrosis factor family; LMP2A, Latent Membrane Protein 2A. Image collected and cropped by CiteAb from the following publication (https://pubmed.ncbi.nlm.nih.gov/36301035), licensed under a CC BY license.

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□ 181 References

old Change in BAFF Protein Expression

LMP2A

Monitoring APOBEC3A protein levels in human cancer cells. Methods Cell Biol (2024)

Immunogenicity and Tolerability of a SARS-CoV-2 TNX-1800, a Live Recombinant Poxvirus Vaccine Candidate, in Syrian Hamsters and New Zealand White Rabbits. Viruses (2023)

A Combinatorial Regulatory Platform Determines Expression of RNA Polymerase III Subunit RPC7 (POLR3G) in Cancer. Cancers (Basel) (2023)

Astrocytic A2A receptors silencing negatively impacts hippocampal synaptic plasticity and memory of adult mice. Glia (2023)

TRAF7 is an essential regulator of blood vessel integrity during mouse embryonic and neonatal development. iScience (2023)

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