

Sheep anti-Rabbit IgG (H+L) Secondary Antibody, HRP

Product Details	
Size	1 mg
Species Reactivity	Rabbit
Host/Isotype	Sheep / IgG
Class	Polyclonal
Туре	Secondary Antibody
Conjugate	HRP
Immunogen	Gamma Immunoglobins Heavy and Light chains
Form	Lyophilized
Concentration	1 mg/mL
Purification	Affinity chromatography
Storage buffer	PBS, pH 7.2, with 1% BSA
Contains	0.1% ProClin 150
Storage conditions	Store at 4°C short term. For long term storage, store at -20°C, avoiding freeze/thaw cycles.
RRID	AB_2534843

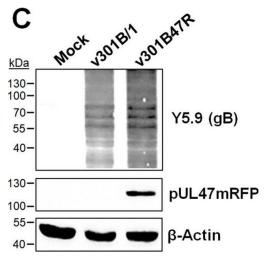
Applications	Tested Dilution	Publications
Western Blot (WB)	1:2,000-1:10,000	0 Publication
Immunohistochemistry (IHC)	1:500-1:20,000	-
ELISA (ELISA)	1:500-1:20,000	-

Product Specific Information

The sensitivity of each lot of antibody is confirmed using ELISA. The specificity of each lot of antibody is confirmed by immunoelectrophoresis (IEP).

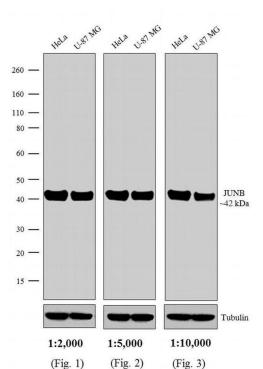
Rehydrate with 1.1 mL of deionized water and let stand 30 minutes at room temperature to dissolve. (Product has been overfilled to ensure complete recovery.) Centrifuge to remove any particulates. Prepare fresh working dilution daily. Store lyophilized material at 2-8 °C. For long term storage after reconstitution, dilute with 50% glycerol and store at -20 °C as a liquid. Based on Immunoelectrophoresis, no reactivity is observed to: non-immunoglobulin rabbit serum proteins.

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



Rabbit IgG (H+L) Secondary Antibody (A16172) in WB

Replication and fluorescent protein expression in tissue culture cells, (A) Mean plaque areas (n = 25) of viruses reconstituted from r301B/1 and r301B47R were measured and the results are shown as box & whisker plots. There were no significant differences in plaque sizes between the two viruses using Student's t tests. (B) Representative plaques for v301B/1 and v301B47R are shown. Plaques were stained with polyclonal chicken anti-GaHV-3 antibody and goat anti-chicken-IgY Alexa488 (green) was used as secondary antibody to identify plaques. Fluorescent expression of mRFP (red) was directly visualized and cells were counterstained with Hoechst 33342 to visualize nuclei. (C) Western blotting for pUL47mRFP using anti-mRFP antibody. The anti-GaHV-3 antibody Y5.931 was used to show the relative level of infection in the cultures. Antibody against chicken -actin is shown as a loading control. (D) Fluorescent protein expression in feather follicles (FFs) infected with v301B47R at 21 dpi. FFs were also stained with anti-HVT L78.2 or -GaHV-3 Y5.9 plus anti-mouse IgG-Alexa488 (green) and images were collected with a fluorescent stereoscope. (E) Percent of chickens positive for pUL47mRFP in experimental and naturally (contact) infected chickens over 60 days. ND not done. Image collected and cropped by CiteAb from the following publication (https://pubmed.ncbi.nlm.nih.gov/33833367), licensed under a CC BY license.



Rabbit IgG (H+L) Secondary Antibody (A16172) in WB

Western blot analysis was performed on nuclear enriched extracts (30 μg lysate) of HeLa (Lane 1) and U-87 MG (Lane 2). The blots were probed with Anti-JUNB Recombinant Rabbit Monoclonal Antibody (Product # 701702, 2 μg/mL) and detected by chemiluminescence using Sheep Anti-Rabbit IgG (H+L) Secondary Antibody, HRP (Product # A16172) at dilutions 1:2,000 (Fig. 1), 1:5,000 (Fig. 2) and 1:10,000 (Fig. 3). A 42 kDa band corresponding to JUNB was observed. Known quantity of protein samples were electrophoresed using Novex® NuPAGE®12 % Bis-Tris gel (Product # NP0342BOX), XCell SureLockTM Electrophoresis System (Product # EI0002) 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 PierceTM ECL Western blotting Substrate (Product # 32106).

□3 References

Mitochondrial Transcription Factor A Regulates Foxp3+ T Cell Mediated Maintenance of CD4+ T Cell Landscapes and Immunological Aging bioRxiv (2022)

The requirement of glycoprotein C (gC) for interindividual spread is a conserved function of gC for avian herpesviruses. Sci Rep (2021)

Effects of FSS on the expression and localization of the core proteins in two Wnt signaling pathways, and their association with ciliogenesis. Int J Mol Med (2018)

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