



Anti-GFP (CHICKEN) Antibody - 600-901-215S

Code: 600-901-215S

Size: 25 µL

Product Description: Anti-GFP (CHICKEN) Antibody - 600-901-215S

Concentration: 1.0 mg/ml by UV absorbance at 280 nm

PhysicalState: Liquid (sterile filtered)

Label	Unconjugated
Host	Chicken
Species Reactivity	wt, rGFP, eGFP
Buffer	0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2
Stabilizer	None
Preservative	0.01% (w/v) Sodium Azide
Storage Condition	Store GFP Antibody at -20° C or below prior to opening. This vial contains a relatively low volume of reagent (25 µL). To minimize loss of volume dilute 1:10 by adding 225 µL of the buffer stated above directly to the vial. Recap, mix thoroughly and briefly centrifuge to collect the volume at the bottom of the vial. Use this intermediate dilution when calculating final dilutions as recommended below. Store the vial at -20°C or below after dilution. Avoid cycles of freezing and thawing.
Synonyms	chicken anti-GFP antibody, chicken Anti-Green Fluorescent Protein Antibody, GFP antibody, Green Fluorescent Protein antibody, EGFP, enhanced Green Fluorescent Protein, Aequorea victoria, Jellyfish antibody
Application Note	Anti-GFP is designed to detect GFP and its variants. This antibody can be used to detect GFP by ELISA (sandwich or capture) for the direct binding of antigen and recognizes wild type, recombinant and enhanced forms of GFP. Biotin conjugated polyclonal anti-GFP used in a sandwich ELISA is well suited to titrate GFP in solution using this antibody in combination with Rockland's monoclonal anti-GFP (600-301-215) using either form of the antibody as the capture or detection antibodies. However, use the monoclonal form only for the detection of wild type or recombinant GFP as this form does not sufficiently detect 'enhanced' GFP. The detection antibody is typically conjugated to biotin and subsequently reacted with streptavidin conjugated HRP (code # S000-03). Fluorochrome conjugated Polyclonal anti-GFP can be used to detect GFP by immunofluorescence microscopy in prokaryotic (E.coli) and eukaryotic (CHO cells) expression systems and can detect GFP containing inserts. Significant amplification of signal is achieved using fluorochrome conjugated polyclonal anti-GFP relative to the fluorescence of GFP alone. For immunoblotting use either alkaline phosphatase or peroxidase conjugated polyclonal anti-GFP to detect GFP or GFP containing proteins on western blots.
Background	Chicken Anti-GFP is designed for immunofluorescence microscopy, fluorescence based plate assays (FLISA) and fluorescent western blotting. GFP Antibody is also suitable for multiplex analysis, including multicolor imaging, utilizing various commercial platforms. Chicken IgY lacks the classic "Fc" domain and does not bind to mammalian IgG Fc receptors. Thus resulting in lower backgrounds for western blotting, ELISA and Immunohistochemistry.
Purity And Specificity	Anti-GFP Antibody IgY purification was prepared from egg yolks by immunoaffinity chromatography using Green Fluorescent Protein (Aequorea victoria) coupled to agarose beads followed by solid phase adsorption(s) to remove any unwanted reactivities. Assay by immunoelectrophoresis resulted in a single precipitin arc against anti-Chicken Serum and purified and partially purified Green Fluorescent Protein (Aequorea victoria). No reaction was observed against Human, Mouse or Rat serum proteins.
Assay Dilutions	User Optimized
ELISA	1:2,000 - 1:12,000
Western Blot	1:200 - 1:1,000
Immunohistochemistry	1:200 - 1:1,000
IF Microscopy	User Optimized
Other Assays	User Optimized
Expiration	Expiration date is three (3) months from date of opening.
Immunogen	The immunogen is a Green Fluorescent Protein (GFP) fusion protein corresponding to the full length amino acid sequence (246aa) derived from the jellyfish Aequorea victoria.

Specific Reference

Tseng C-Y, Firestein BL. The role of PSD-95 and cypin in morphological changes in dendrites following sublethal NMDA exposure. *J. Neurosci.* 2011 Oct 26;31(43):15468–15480.

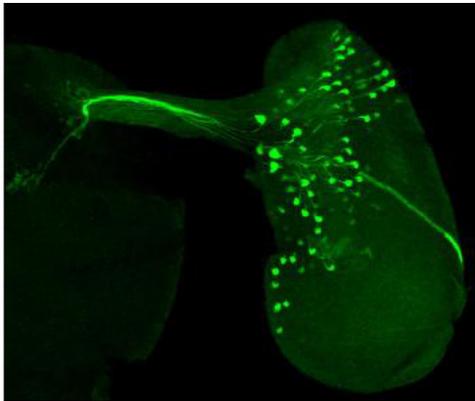
He M. (2012) miRNA Tagging and Affinity-purification (miRAP). *Bio Protoc.* 2012 Oct 5;2(19). pii: e265.

Related Products

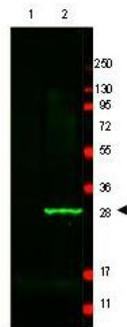
200-301-268	Anti-AKT pS473 (MOUSE) Monoclonal Antibody - 200-301-268
610-4302	Anti-MOUSE IgG (H&L) (RABBIT) Antibody Peroxidase Conjugated - 610-4302
611-1302	Anti-RABBIT IgG (H&L) (GOAT) Antibody Peroxidase Conjugated - 611-1302
B304	NORMAL GOAT SERUM (NGS) - B304

Images

- 1 Immunofluorescence Microscopy of chicken anti-GFP antibody. Tissue: KruppelGAL4 driver line in *Drosophila* eye disc. Fixation: 0.5% PFA. Antigen retrieval: not required. Primary antibody: anti-GFP antibody diluted 1:500 for 2 hr at RT. Secondary antibody: Alexa™488 conjugated anti-Chicken IgG at 1:300 for 1 hr at RT. Blocking: 5% NGS in PBS with 0.1% Triton X-100 for 15 min. Staining: recombinant tau-myc-GFP protein as green fluorescent signal.



- 2 Western Blot of anti-GFP antibody. Lane 1: mouse spleen lysate. Lane 2: mouse spleen lysate spiked with 50ng of WT-GFP. Load: 20 µg per lane. Primary antibody: Anti-GFP antibody at 2 µg/ml for 2 hr at room temperature. Secondary antibody: IRDye™800 Conjugated Affinity Purified anti-Chicken IgG [H&L] [Goat] MX10 (p/n 603-132-126) at 1:20,000 for 45 min at RT. Block: 5% BSA in PBS 2 hr at room temperature. Predicted/Observed size: 27 kDa for GFP epitope tag. Other band(s): none.



- 3 Western Blot of Anti-GFP (CHICKEN) antibody. Lane 1: MW. Lane 2: GFP. Load: 0.05 µg. Primary antibody: Anti-GFP (CHICKEN) antibody (p/n 600-901-215) was used at 1:1000 overnight at 4°C. Secondary antibody: Anti-Chicken IgG (GOAT) peroxidase conjugated antibody (p/n 603-103-126) secondary antibody was used at 1:40,000 in Blocking Buffer for Fluorescent Western Blotting (p/n MB-070). Block: 1% BSA-TTBS (p/n MB-013, diluted to 1X) 30 min at 20°C. Predicted/Observed size: 27 kDa for GFP. Other band(s): none.

100

80

60

50

40

30

20



Disclaimer

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