

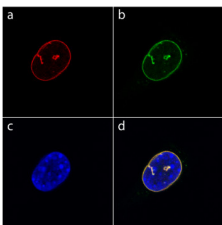
Product no **AS15 2987****GFP | Green Fluorescence Protein (affinity purified)****Product information**

Immunogen	Highly purified native GFP protein derived from <i>Aequorea victoria</i> , UniProt: P42212
Host	Rabbit
Clonality	Polyclonal
Purity	Immunogen affinity purified serum in PBS pH 7.4.
Format	Liquid
Quantity	100 µg at 1 mg/ml
Storage	Store at -20 °C. Avoid repeated freeze-thaw cycles. Please remember to spin the tubes briefly prior to opening them to avoid any losses that might occur from material adhering to the cap or sides of the tube.
Additional information	10 mM TRIS buffer pH 8,0 containing 0,02% sodium azide

Application information**Recommended dilution** | 1 : 5000-1 : 25 000 (ELISA), 1: 100 - 1 : 500 (IF), 1 : 2000-1 : 10 000 (WB)**Confirmed reactivity** | Native GFP, Recombinant GFP (E.coli), all variants of GFP, including EGFP**Additional information** | Minimal cross-reactivity with *E.coli* proteins.

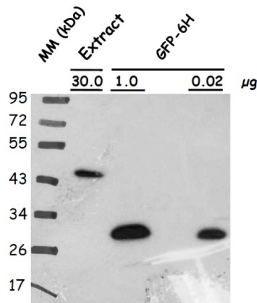
Antibody reactivity to mVenus or EBFP2 has not yet been determined.

- Selected references**
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- [Sun](#) et al. (2021) The epigenetic factor FVE orchestrates cytoplasmic SGS3-DRB4-DCL4 activities to promote transgene silencing in Arabidopsis. *Sci Adv.* 2021 Aug 4;7(32):eabf3898. doi: 10.1126/sciadv.abf3898. PMID: 34348894; PMCID: PMC8336953.
- [Stelate](#) et al. (2021) Correlative Light-Environmental Scanning Electron Microscopy of Plasma Membrane Efflux Carriers of Plant Hormone Auxin. *Biomolecules.* 2021 Sep 26;11(10):1407. doi: 10.3390/biom11101407. PMID: 34680040; PMCID: PMC8533460.
- [Wieczorek](#) et al. (2020) Development of a New Tomato Tarrado Virus-Based Vector Tagged with GFP for Monitoring Virus Movement in Plants. *Viruses.* 2020 Oct 20;12(10):1195. doi: 10.3390/v12101195. PMID: 33092281; PMCID: PMC7588970.
- [Kulich](#) et al. (2018). Deubiquitinase OTU5 affects Root Responses to Phosphate Starvation. *Plant Physiol.* 2018 Jan 4. pii: pp.01693.2017. doi: 10.1104/pp.17.01693.

Immunolocalization

Detection of YFP-tagged Lamin A in the nucleus of mouse fibroblasts using GFP | Green Fluorescence Protein (affinity purified) antibodies. Fixation and permeabilization was performed with methanol at -20 °C for 10 min. **a)** Indirect immunofluorescence labeling of YFP-lamin A with anti-GFP primary antibody (dilution 1: 500) as primary antibody and detected by TRITC-conjugated goat-anti-rabbit as secondary antibody. **b)** Green fluorescence image of YFP-tagged Lamin A, which is incorporated into the nuclear lamina and tubular structures that penetrate into the nucleus. **c)** DAPI staining of nuclear DNA. **d)** Merged images

western blot



Protein extracts were obtained from *Arabidopsis thaliana* seeds (producing atLEA4-5 fused to GFP; other lanes contain GFP-6H loaded in indicated amounts). Following extraction buffer was used: 0.7 M sucrose, 0.5 M Tris-base, 30 mM HCl, 50 mM EDTA, 0.1 M KCl, 2% β -mercaptoethanol, 12 mg/ml poly-vinyl-poly-pyrrolidone (PVPP). This buffer was complemented with 2mL of equilibrated phenol before extraction. Samples were centrifuged and the protein phase was recovered; the extracted proteins were precipitated with 0.1 M ammonium acetate dissolved in methanol, centrifuged and the pellet washed with cold (-20 °C) 80% acetone. The protein pellet was dissolved in SDS-solubilization buffer (1% CHES, 2% SDS, 2% β -mercaptoethanol, 10% glycerol). Thirty μ g of seed protein extract was denatured with Laemmli buffer at 95 °C for 5 min and proteins were separated on 15% SDS-PAGE and blotted 1.5 h to nitrocellulose membrane in a liquid transfer system. Blots were blocked with 2% skim milk ON at 4 °C with agitation. After rinsing with TBS, blots were incubated in the primary antibody at a dilution of 1:10 000 (anti-GFP) for 3h at 4 °C with agitation. The antibody solution was decanted and the blot was rinsed briefly, then washed once for 15 min. and 3 times for 5 min in TBS-T at 4 °C with agitation. Then, blots were incubated in secondary antibody (anti-rabbit IgG horse-radish peroxidase conjugate) diluted to 1:25 000 in for 1h at RT with agitation. The blot was washed as above and developed for 2 min with SuperSignal West Pico Chemiluminescent Substrate, Thermo Fisher Scientific. Exposure time was for 30 seconds.

Courtesy of Dr. Alejandra Covarrubias, UNAM, Mexico