

This product is **for research use only** (not for diagnostic or therapeutic use)

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### Product no AS09 602-trial

# Goat anti-Rabbit IgG (H&L), HRP conjugated - trial sample

### **Product information**

**Immunogen** Purified Rabbit IgG, whole molecule,

Host Goat

Clonality Polyclonal

**Purity** Immunogen affinity purified using solid phase rabbit IgG.

Format Liquid

Quantity 10 μl

Storage Store liquid material at 2-8 °C up to 6 months.

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**Additional information** Concentration: 1.0 mg/ml.

Antibody is provided in: 10 mM Sodium Phosphate, 0.15 M Sodium Chloride, pH 7.2, 1% BSA (w/v), Protease IgG free, 0.1 % (v/v) Kathon CG.

Affinity purified antibody is > 95 % pure, according to SDS-PAGE.

## **Application information**

Recommended dilution 1:50 000 -1:90 000 (ELISA), 1:500 -1:5000 (IHC), 1:10 000 -1:50 000 (WB)

Confirmed reactivity Based on IEP, this antibody Reacts with: Rabbit IgG heavy chainslight chains on all Rabbit immunoglobulins

Not reactive in No confirmed exceptions from predicted reactivity are currently known

**Additional information** No reactivity is observed to non-immunoglobulin rabbit serum

Selected references

Migocka et al. (2018). Cucumber metal tolerance protein 7 (CsMTP7) is involved in the accumulation of Fe in mitochondria under Fe excess. Plant J. 2018 Jun 22. doi: 10.1111/tpj.14006.

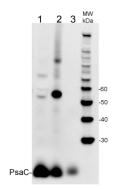
Tong et al. (2018). Delivery of siRNA in vitro and in vivo using PEI-capped porous silicon nanoparticles to silence MRP1 and inhibit proliferation in glioblastoma. J Nanobiotechnology. 2018 Apr 13;16(1):38. doi: 10.1186/s12951-018-0365-y. Nikkanen et al. (2018). Regulation of chloroplast NADH dehydrogenase-like complex by NADPH-dependent thioredoxin system. CSH, BioRixiv. doi.org/10.1101/261560.

<u>Gzyl</u> et al. (2017). Gamma-tubulin distribution and ultrastructural changes in root cells of soybean (Glycine max L.) seedlings under cadmium stress. Environmental and Experimental Botany, Vol 143, Nov 2017, Pages 82-90. <u>Kamies</u> et al. (2017). A Proteomic Approach to Investigate the Drought Response in the Orphan Crop Eragrostis tef. Proteomes. 2017 Nov 15;5(4). pii: E32. doi: 10.3390/proteomes5040032.

Niederhuber et al. (2017). Super-resolution microscopy of the β-carboxysome reveals a homogenous matrix. Mol Biol Cell. 2017 Aug 9. pii: mbc.E17-01-0069. doi: 10.1091/mbc.E17-01-0069.

This antibody is listed in first 7000 most published antibodies in the world by CiteAB report.

### application example



5 μg of total extract from (1) Hordeum vulgaretotal leaf, (2) Zea mays (3) Spinacia oleracea extracted with PEB (AS08 300) were separated on 4-12% NuPage (Invitrogen) LDS-PAGE and blotted 1h to PVDF. Blots were blocked immediately following transfer in 2% ECL Advance blocking



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reagent (GE Healthcare) in 20 mM Tris, 137 mM sodium chloride pH 7.6 with 0.1% (v/v) Tween-20 (TBS-T) for 1h at room temperature with agitation. Blots were incubated in the primary anti-PsaC antibody (AS04 042) at a dilution of 1: 10 000 for 1h at room temperature with agitation. The antibody solution was decanted and the blot was rinsed briefly twice, then washed once for 15 min and 3 times for 5 min in TBS-T at room temperature with agitation. Blots were incubated in secondary antibody (goat anti-rabbit IgG horse radish peroxidase conjugated, AGRISERA) diluted to 1:50 000 in 2% ECL Advance blocking solution for 1h at room temperature with agitation. The blots were washed as above and developed for 5 min with ECL Advance detection reagent according to the manufacturers instructions. Images of the blots were obtained using a CCD imager (FluorSMax, Bio-Rad) and Quantity One software (Bio-Rad). Exposure time was 30 seconds.

#### Comparison of Agrisera secondary antibody sensitivity



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10 μg of mitochondrial fraction from *Arabidopsis thaliana* (1,3) and *Arabidopsis thaliana* leaf extract (2,4) were separated on 10% gel and blotted on nitrocellulose membrane using wet transfer (0.22% CAPS, pH 11). Filters where blocked (1.5h) in 5% milk in TBST (1X TBS, 0,1% Tween 20), incubated with 1: 1000 anti-COXII antibodies (2h in TBST) followed by incubation with 1: 10 000 secondary anti-rabbit (1h) HRP-coupled antibodies from **Agrisera (left panel)** and **other manufacture (right panel)** and visualized with standard ECL on Kodak autoradiography film for 5 s. Antibody in left panel detects target protein also in total cell extract (2) and can be used in higher dilution than applied 1: 10 000.

Agrisera goat anti-rabbit HRP conjugated antibody (AS09 602) can be used at following dilutions: 1: 50 000 -1: 90 000 (ELISA), 1: 75 000 with enhanced ECL and 1: 25 000 with regular ECL (WB), 1: 500 -1: 5000 (IHC).