

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

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## Product no AS08 296

# Anti-GLN2 | GS2, chloroplastic form of glutamine synthetase

#### **Product information**

Immunogen

KLH-conjugated synthetic peptide which is a part of part of the glutamine synthetase/guanido kinase superfamily catalytic region chosen from various available sequences, including Arabidopsis thaliana GLN2, UniProt: Q43127, TAIR: <u>AT5G35630</u>

**Host** Rabbit

Clonality Polyclonal

Purity Serum

Format Lyophilized

Quantity 50 μl

**Reconstitution** For reconstitution add 50 µl of sterile water

Store lyophilized/reconstituted at -20 °C; once reconstituted make aliquots to 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 This product can be sold contacining proclin if requested

# Application information

**Recommended dilution** 1:5000 on 0.5-5 μg protein/lane detection (WB)

Expected | apparent

MW

47 | 44-45 kDa

Confirmed reactivity

Arabidopsis thaliana, Nicotiana tabacum, Oryza sativa, Pisum sativum, Spinacia oleracea

Predicted reactivity

Brassica napus, Diplotaxis tenuifolia, Eruca versicaria, Glycine max, Hordeum vulgare, Medicago truncatula, Pinus sylvestris, Phaseolus vulgaris, Physcomitrium patens, Populus sp., Triticum aestivum, Zea mays

Species of your interest not listed? Contact us

Not reactive in Diatoms

Selected references

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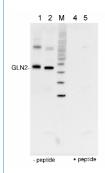
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## Application example





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0.5 μg of protein from Arabidopsis thaliana total leaf fraction (1), 5 μg of protein from *Spinacia oleracea* chlorplast enriched fraction (2), molecular weight markers (MagicMark<sup>TM</sup>,Invitrogen) (M), the same samples as in 1 and 2 but after peptide neutralisation assay, e.g. incubation of the antibody with 100 mM excess of peptide used to elicit andt-GLN2 antibody (4,5), extracted with PEB (AS08 300), were separated on 4-12% NuPage (Invitrogen) LDS-PAGE and blotted 1h to PVDF (Millipore). Filters were blocked 1h with 2% low-fat milk powder in TBS-T (0.1% TWEEN 20) and probed with anti-GLN2 antibody (AS08 296, 1:5 000, 1h) and secondary anti-rabbit (1:20000, 1 h) antibody (HRP conjugated) in TBS-T containing 2% low fat milk powder. Antibody incubations were followed by washings in TBS-T. All steps were performed at RT with agitation. Blots were developed for 5 min with chemiluminescent detection reagent according the manufacturers instructions. Images of the blots were obtained using a CCD imager (FluorSMax, Bio-Rad) and Quantity One software (Bio-Rad).