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Product no AS11 1787

## Anti-PsbC | CP43 protein of PSII

## **Product information**

Immunogen KLH-conjugated synthetic peptide chosen from known sequences of PsbC including Arabidopsis thaliana PsbC,

UniProt: <u>P56778</u>, TAIR: <u>AtCg00280</u>

**Host** Rabbit

Clonality Polyclonal

**Purity** Serum

Format Lyophilized

Quantity 50 ul

**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 Storage 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 Contains 0,01% ProClin

## **Application information**

Recommended dilution 1:3 000 (WB)

Expected | apparent

45 | 43 kDa

Confirmed reactivity

Arabidopsis thaliana, Chlamydomonas reinhardtii, Chlorella sorokiniana, Chlorella vulgaris, Chromochloris zofingiensis, Echinochloa crus-galli, Hordeum vulgare, Oryza sativa, Panax ginseng, Physcomitrium patens, Pisum sativum, Phaseolus vulgaris, Pyropia yezoensis, Synochococcus sp. PCC7002, Synechocystis sp. PCC6803, Tillandsia flabellate, Triticum aestivum, Triticale, Zea mays, Verbascum lychnitis, Vigna radiate

Predicted reactivity

Asimina parviflora, Borago officinalis, Cannabis sativa, Carthamus persicus, Casimirella guaranitica, Catalpa bungei, Calatola mollis, Citron x limon, Cunninghamia lanceolata, Deeringothamnus rugelii, Gonystylus bancanus, Ipomopsis aggregata, Leretia cordata, Lobatiriccardia lobata, Myricaria germanica, Nostoc sp. PCC7120, Nannochloropsis sp., Natsiatum herpeticum, Nicotiana benthamiana, Nothapodytes montana, Nerium oleander, Ottoschulzia rhodoxylon, Oxandra lanceolata, Solanum tuberosum, Oryza sativa, Panax quinquefolius, Prosopidastrum angusticarpum, Prosopis glandulosa, Rollinia mucosa, Rosmarinus officinalis, Saxifraga rivularis, Spinacia oleracea, Zelkova serrata, Zinnia violacea, Vachellia caven, Vitis vinifera, Zosteria marina, Xerocladia viridiramis

Species of your interest not listed? Contact us

Not reactive in Diatoms

Additional information In C4 plants like Echinochloa crus-galli and Zea mays antibody detects 2 bands.

Selected references

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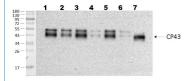
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Okegawa et al (2021) Maintaining the Chloroplast Redox Balance Through the PGR5-Dependent Pathway and the Trx System is Required for Light-Dependent Activation of Photosynthetic Reactions. Plant Cell Physiol. 2021 Oct 8:pcab148. doi: 10.1093/pcp/pcab148. Epub ahead of print. PMID: 34623443.



**5 μg of total protein** from (1) *Arabidopsis thaliana* leaf extracted with **Protein Extraction Buffer**, PEB (**AS08 300**), (**2**) *Hordeum vulgare* leaf extracted with PEB, (**3**) *Chlamydomonas reinhardtii* total cell extracted with PEB, (**4**) *Synechococcus* sp. 7942 total cell extracted with PEB, extracted with PEB were separated on **4-12%** NuPage (Invitrogen) **LDS-PAGE** and blotted 1h to **PVDF**. Blots were blocked immediately following transfer in 2% blocking reagent 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 antibody 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 (anti-rabbit IgG horse radish peroxidase conjugated, recommended secondary antibody **AS09 602**) diluted to 1:25 000 in 2% blocking solution for 1h at room temperature with agitation. The blots were washed as above and 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). Exposure time was 75 seconds.



1.5 µg of chlorophyll from thylakoids of various treatments of *Echinochloa crus-galli* (1-2), *Zea mays* (3-5), *Pisum sativum* (6-7), extracted with 0.4 M sorbitol, 50 mM Hepes NaOH, pH 7.8, 10 mM NaCl, 5 mM MgCl2 and 2 mM EDTA. Samples were denatured with Laemmli buffer at 75 °C for 5 min and were separated on 12% SDS-PAGE and blotted 30 min to PVDF using wet transfer. Blot was blocked with 5% fatty acid free milk for 1h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1: 3 000 overnight at 4 °C with agitation in 1% milk in TBS-T. The antibody solution was decanted and the blot was washed 4 times for 5 min in TBS-T at RT with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated, <u>AS09 602</u>, Agrisera ) diluted to 1:25 000 in 1 % milk in TBS-T for 1h at RT with agitation. The blot was washed 5 times for 5 min in TBS-T and 2 times for 5 min in TBS, and developed for 1 min with 1.25 mM luminol, 0.198 mM coumaric acid and 0.009% H2O2 in 0.1 M Tris- HCl, pH 8.5. Exposure time in ChemiDoc System was 240 seconds.

Courtesy of Dr. Wiola Wasilewska, Warsaw University, Poland