

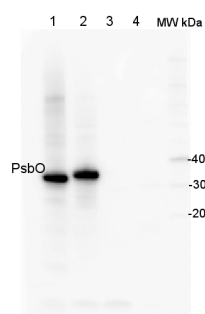
Product no **AS05 092****PsbO | 33 kDa of the oxygen evolving complex (OEC) of PSII (anti-peptide)****Product information**

|                       |   |
|-----------------------|---|
| <b>Immunogen</b>      | N-terminally located peptide chosen from <i>Arabidopsis thaliana</i> PsbO1 and PsbO2 isoforms. UniProt: <a href="#">P23321</a> , PsbO1, TAIR: <a href="#">At5g66570</a> ; UniProt: <a href="#">A0A178VBH5</a> , TAIR: <a href="#">At3g50820</a>                               |
| <b>Host</b>           | Rabbit  |
| <b>Clonality</b>      | Polyclonal  |
| <b>Purity</b>         | Serum   |
| <b>Format</b>         | Lyophilized   |
| <b>Quantity</b>       | 100 µl  |
| <b>Reconstitution</b> | For reconstitution add 100 µl of sterile water  |
| <b>Storage</b>        | 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** Loading based on 50-100 ng of chlorophyll is enough to obtain good signal with this antibody

**Application information**

|                               |   |
|-------------------------------|---|
| <b>Recommended dilution</b>   | 1 : 1000 (WB)   |
| <b>Expected   apparent MW</b> | 33 kDa  |
| <b>Confirmed reactivity</b>   | <i>Arabidopsis thaliana</i> , <i>Cucumis sativus</i> , <i>Hordeum vulgare</i> , <i>Manihot esculenta</i> , <i>Nicotiana tabacum</i> , <i>Pisum sativum</i> , <i>Sinapis alba</i> , <i>Triticum aestivum</i> , <i>Zea mays</i>   |
| <b>Predicted reactivity</b>   | <i>Brassica oleracea</i> , <i>Pisum sativum</i> , <i>Populus tremula</i> , <i>Picea sitchensis</i> , <i>Vitis vinifera</i><br>Species of your interest not listed? <a href="#">Contact us</a>   |
| <b>Not reactive in</b>        | <i>Chlamydomonas reinhardtii</i> , <i>Synechococcus</i> sp. PCC 7942  |
| <b>Additional information</b> | Good signal is obtained with this antibody with a load from 0,5 chlorophyll µg/well   |
| <b>Selected references</b>    | <p><a href="#">Mazur</a> et al. (2021) The SnRK2.10 kinase mitigates the adverse effects of salinity by protecting photosynthetic machinery. <i>Plant Physiol.</i> 2021 Dec 4;187(4):2785-2802. doi: 10.1093/plphys/kiab438. PMID: 34632500; PMCID: PMC8644180.</p> <p><a href="#">Toubiana</a> et al. (2020). Correlation-based Network Analysis Combined With Machine Learning Techniques Highlight the Role of the GABA Shunt in <i>Brachypodium Sylvaticum</i> Freezing Tolerance. <i>Sci Rep</i>, 10 (1), 4489</p> <p><a href="#">Wang</a> et al. (2019). YR36/WKS1-mediated Phosphorylation of PsbO, an Extrinsic Member of Photosystem II, Inhibits Photosynthesis and Confers Stripe Rust Resistance in Wheat. <i>Mol Plant.</i> 2019 Oct 14. pii: S1674-2052(19)30330-2. doi: 10.1016/j.molp.2019.10.005.</p> <p><a href="#">An</a> et al. (2019). Protein cross-interactions for efficient photosynthesis in the cassava cultivar SC205 relative to its wild species. <i>J Agric Food Chem.</i> 2019 Jul 19. doi: 10.1021/acs.jafc.9b00046.</p> <p><a href="#">Rozp?dek</a> et al. (2018). Acclimation of the photosynthetic apparatus and alterations in sugar metabolism in response to inoculation with endophytic fungi. <i>Plant Cell Environ.</i> 2018 Dec 5. doi: 10.1111/pce.13485.</p> |

**Application example**

**2 µg of total protein** from (1) *Arabidopsis thaliana* leaf, (2) *Hordeum vulgare* leaf, (3) *Chlamydomonas reinhardtii* total cell, (4) *Synechococcus* sp. 7942 total cell were all extracted with PEB (**AS08 300**) and 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) diluted to 1:50 000 in 2% blocking solution for 1h at room temperature with agitation. The blots were washed as above and developed for 5 min with chemiluminescence 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).