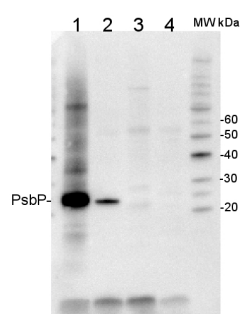


Product no **AS06 167****PsbP | 23 kDa protein of the oxygen evolving complex (OEC) of PSII (anti-peptide)****Product information**

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| Immunogen | KLH-conjugated synthetic peptide derived from PsbP protein of <i>Arabidopsis thaliana</i> Q42029 , At1g06680 |
| Host | Rabbit |
| Clonality | Polyclonal |
| Purity | Serum |
| Format | Lyophilized |
| Quantity | 100 µl |
| Reconstitution | For reconstitution add 100 µl of sterile water |
| Storage | 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 containing ProClin if requested |

Application information

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| Recommended dilution | 1 : 2000 (WB) |
| Expected apparent MW | 28 23 kDa (<i>Arabidopsis thaliana</i>) |
| Confirmed reactivity | <i>Arabidopsis thaliana</i> , <i>Hordeum vulgare</i> , <i>Nicotiana tabacum</i> , <i>Picea abies</i> , <i>Pisum sativum</i> , <i>Physcomitrium patens</i> |
| Predicted reactivity | <i>Oryza sativa</i> , <i>Solanum tuberosum</i> , <i>Triticum aestivum</i> , <i>Picea sitchensis</i> , <i>Populus balsamifera</i> |
| | Species of your interest not listed? Contact us |
| Not reactive in | <i>Chlamydomonas reinhardtii</i> , <i>Synechococcus</i> sp. PCC 7942, <i>Zostera marina</i> |
| Additional information | If you work with <i>Chlamydomonas reinhardtii</i> , please use following PsbP antibody: AS06 142-23 |
| Selected references | <p>Mazur 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>Tamburino et al. (2017). Chloroplast proteome response to drought stress and recovery in tomato (<i>Solanum lycopersicum</i> L.). <i>BMC Plant Biol.</i> 2017 Feb 10;17(1):40. doi: 10.1186/s12870-017-0971-0.</p> <p>Pavlovic et al. (2016). Light-induced gradual activation of photosystem II in dark-grown Norway spruce seedlings. <i>Biochim Biophys Acta.</i> 2016 Feb 18. pii: S0005-2728(16)30028-7. doi: 10.1016/j.bbabo.2016.02.009.</p> <p>Albanese et al. (2016). Isolation of novel PSII-LHCII megacomplexes from pea plants characterized by a combination of proteomics and electron microscopy. <i>Photosynth Res.</i> 2016 Jan 9.</p> <p>Grassi et al. (2012). Early events in plastid protein degradation in stay-green <i>Arabidopsis</i> reveal differential regulation beyond the retention of LHCII and chlorophyll. <i>J. Proteome Res.</i> October 2.</p> <p>Lang et al. (2011). Simultaneous isolation of pure and intact chloroplasts and mitochondria from moss as the basis for sub-cellular proteomics. <i>Plant Cell Rep.</i> 2011 Feb;30(2):205-15. doi: 10.1007/s00299-010-0935-4.</p> |

Application example

2 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, 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: 50 000 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 1:10 000 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 to the manufacturers instructions. Images of the blots were obtained using a CCD imager (FluorSMax, Bio-Rad) and Quantity One software (Bio-Rad).