

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

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### Product no AS16 4065

# NdhH | NAD(P)H-quinone oxidoreductase subunit H (chloroplastic)

#### **Product information**

Immunogen KLH-conjugated synthetic peptide derived from Arabidopsis thaliana NdhH protein sequence, UniProt: P56753-1, TAIR:

**Host** Rabbit

Clonality Polyclonal

**Purity** Immunogen affinity purified serum in PBS pH 7.4.

Format Lyophilized

Quantity 50 μg

**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.

## **Application information**

Recommended dilution 1:5000 (WB)

Expected | apparent

45 | 49 kDa

Confirmed reactivity | Arabidopsis thaliana, Zea mays

Predicted reactivity

Cannabis sativa, Hordeum vulgare, Microcystis aeruginosa PCC 7806, Physcomitrella patens, Synechocystis sp.

Species of your interest not listed? Contact us

Not reactive in Pisum sativum, Phaseolus vulgaris

**Additional information** This product can be sold containing proclin if requested

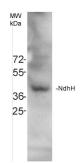
Selected references

Seiml-Buchinger et al. (2022) Ascorbate peroxidase postcold regulation of chloroplast NADPH dehydrogenase activity controls cold memory. Plant Physiol. 2022;190(3):1997-2016. doi:10.1093/plphys/kiac355

Wada et al. (2021) Identification of a Novel Mutation Exacerbated the PSI Photoinhibition in pgr5/pgrl1 Mutants; Caution for Overestimation of the Phenotypes in Arabidopsis pgr5-1 Mutant. Cells. 2021 Oct 26;10(11):2884. doi: 10.3390/cells10112884. PMID: 34831107; PMCID: PMC8616342.

Nikkanen et al. (2018). Regulation of chloroplast NADH dehydrogenase-like complex by NADPH-dependent thioredoxin system. CSH, BioRixiv, doi: https://doi.org/10.1101/261560.

#### **Application example**



Total proteins of Arabidopsis thaliana leaf were precipitated with 10% TCA, washed with acetone and solubilized in a buffer with 8M urea, 100 mM Tris-HCL pH 7.5, 1 mM EDTA, 2% SDS. 10 ug of TCA precipites were separated in a BioRad TGX 4-20% precast gels and blotted 1h to a PVDF membrane using Höefer semi-dry blotter. Blot was blocked with 8 % milk in TTBS for 30 min. at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1: 5 000 overnight in +4°C. The antibody solution was decanted and the blot was washed 3 x 5 min with TTBS at RT with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated, from Agrisera, AS09 602) diluted to 1:20 000 in 1% milk/TTBS for 2h at RT with agitation. The blot was washed 3x5 min in TTBS and 1x5 min in TBS at RT with agitation and developed for 5 min with ECL according to the manufacturer's instructions.



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