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## Product no AS23 4975

## Anti-PDHE1-B | Pyruvate dehydrogenase E1 component subunit beta-1, mitochondrial **Product information**

Immunogen KLH-conjugated unique peptide derived from Arabidopsis thaliana PDHE1-B protein sequence, UniProt: Q38799 TAIR:

**Host** Rabbit

Clonality Polyclonal

**Purity** Antigen affinity purified serum, in PBS pH 7.4

Format Lyophilized

Quantity 50 ug

**Reconstitution** For reconstitution, add 50 μl, of sterile or deionized water.

Store lyophilized/reconstituted at -20°C; once reconstituted make aliquots to avoid repeated freeze-thaw cycles. Storage Please, remember to spin tubes briefly prior to opening them to avoid any losses that might occur from lyophilized

material adhering to the cap or sides of the tubes.

## Application information

Recommended dilution 1:500 - 1:2000 (WB)

Expected | apparent

MW

39.2 | 37 kDa (signal peptide is removed)

Predicted reactivity

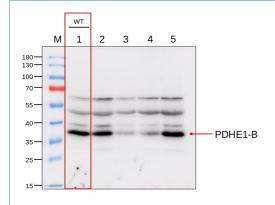
Arachis hypogaea, Brachypodium distachyon, Brassica napus, Cannabis sativa, Capsicum annuum, Citrus sp., Cucumis sativus, Glycine max, Gossypium sp., Hordeum vulgare, Malus domestica, Manihot esculenta, Medicago truncatula, Nicotiana tabacum, Oryza sativa, Pisum sativum, Populus sp., Solanum lycopersicum, Solanum tuberosum,

Sorghum bicolor, Spinacia oleracea, Theobroma cacao, Triticum sp., Vitis vinifera, Zea mays

Species of your interest not listed? Contact us

Not reactive in No confirmed exceptions from predicted reactivity are currently known

**Selected references** To be added when available. Antibody released in October 2024.



## Samples:

- 1 40 μg of Arabidopsis thaliana Col-0 whole leaf extract.
- 2 40 μg of Arabidopsis thaliana mutant iar4 (AT1G24180), Arabidopsis thaliana pyruvate dehydrogenase E1a-like subunit.
- 3 40 μg of Arabidopsis thaliana mutant iar4l-1 (AT1G59900), Arabidopsis thaliana pyruvate dehydrogenase E1a subunit.
- 4 40 μg of Arabidopsis thaliana mutant iar4l-2 (AT1G59900), Arabidopsis thaliana pyruvate dehydrogenase E1a subunit.
- 5 40 ug of Arabidopsis thaliana mutant SALK\_004367C (AT5G50850), Arabidopsis thaliana pyruvate dehydrogenase E1 component subunit beta-1. The T-DNA insertion was located in the 3'UTR region of AT5G50850, thus SALK 004367C might not be a knock-out mutant of mtPDH-E1b.

40 μg/well of total protein extracted freshly from Arabidopsis thaliana whole rosette leaves. Exact buffer components were: 50 mM Tris-HCl (pH 7.5), 150 mM NaCl, 0.5 mM EDTA, 10% (v/v) glycerol, 1% (v/v) Nonidet P-40 (NP-40), 1% (w/v) deoxycholate, 0.1% (w/v) SDS, 1 × Complete protease inhibitor cocktail (Roche), 1 mM PMSF, and denatured with 5xSDS sample buffer (300 mM Tris-HCL(pH 6.8), 10% SDS, 0.1% Bromophenol, 50% Glycerol, 500 mM DTT) at 95°C/5 min. Samples were separated in the RT on 10 % SDS-PAGE, and blotted for 7 min to nitrocellulose (pore size of 0.2 um), using: iBlotTM Dry Blotting System (Invitrogen) in the RT. Blot was blocked with 5% milk for: 3h/RT with



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agitation. Blot was incubated in the primary antibody at a dilution of 1: 500 for ON/4°C with agitation in 2% milk. The antibody solution was decanted, and the blot was rinsed briefly twice, then washed 5 times for 5 min in PBS-T at RT with agitation. Blot was incubated in matching secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1: 5000 in 2% milk for 1h/RT with agitation. The blot was washed as above and developed with a following chemiluminescent detection reagent: AS16 ECL-N-10 Agrisera Bright. Exposure time was 1 minute.

To increase signal/noise ratio, primary antibody can be used in a dilution of 1: 1000 or 1: 2000 1h/RT incubation.

Courtesy of Dr. Mengping Li, Department of Botany and Plant Biology, UNIGE, Switzerland