

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

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Product no AS22 4883

PetL | Cytochrome b6-f complex subunit 6

Product information

Immunogen KLH-conjugated peptide derived from Nicotiana tabacum PetL protein sequence, UniProt: P69401

Host Rabbit

Clonality Polyclonal

Purity Antigen affinity purified serum, in PBS pH 7.4

Format Lyophilized

Quantity 50 μg

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. 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 1:1000 (WB)

Expected | apparent

3,4 | kDa (due to N-terminal or C-terminal processing)

Confirmed reactivity Nicotiana tabacum

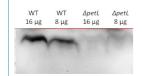
Predicted reactivity

Arabidopsis thaliana, Elsholtzia minima, Plantago ovata, Solanum aloysiifolium, Solanum lycopersicum, Solanum

tuberosum, 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 available in March 2024.



8-16 µg/well of thylakoids extracted from Nicotiana tabacum. Exact buffer components were: 50 mM Hepes-KOH pH 7.5, 100 mM sorbitol, 10 mM MgCl2 and denatured with 0.1 M Tris/HCl pH 6.8, 0.24% glycerol, 8% SDS, 2% b-mercaptoethanol, 0.2 mg/ml Coomassie Blue G250 at 95 °C 10 min. Samples were separated in the cold on 10 % SDS-tricine and blotted for overnight to PVDF, using wet transfer in the cold. Blot was blocked with 4 % milk for 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1: 500 for ON/4°C with agitation. The antibody solution was decanted and the blot was rinsed briefly twice, then washed once for 10 min and 3 times in TBS-T at RT with agitation. Blot was incubated in matching secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated, AS09 602) diluted to 1: 25 000 1h/RT for 1h/RT with agitation. The blot was washed as above and developed with a following chemiluminescent detection reagent: AgriseraBright, AS16 ECL-N. Exposure time was 25 seconds.