

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

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Product no AS05 071 Anti-AtpH | ATP synthase subunit c (chloroplastic)

Product information

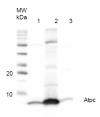
Immunogen	Purified c subunit from <i>Spinacia oleracea</i> UniProt: <u>P69447.</u>
Host	Rabbit
Clonality	Polyclonal
Purity	Serum
Format	Lyophilized
Quantity	200 µl
Reconstitution	For reconstitution add 200 µl of sterile water
Storage	Store lyophilized/reconstituted at -20°C; once reconstituted make aliquots 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	
Recommended dilution	1 : 1000-1 : 10 000 (WB)
Expected apparent MW	8 kDa (for Arabidopsis thaliana)
Confirmed reactivity	Arabidopsis thaliana, Nicotiana benthamina, Spinacia oleracea, Thermosynechococcus elongatus
Predicted reactivity	Algae, Cannabis sativa, Cyclotella cryptica, Glycine max, Oryza sativa, Physcomitrium patens, Phaeodactylum tricornutum, Pisum sativum, Populus alba, Pinus thunbergii, Thalassiosira pseudonana, Zea mays, Vitis vinifera
	Species of your interest not listed? Contact us

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

Additional information Note that increased incubation at 95°C (20-30 min) prior to loading is recommended to break the multimeric c-mer structure, detection of partial ring structures (e,g, 5 or 6 subunits) may occur

Nath et al. (2016). A Nitrogen-Fixing Subunit Essential for Accumulating 4Fe-4S-Containing Photosystem I Core Selected references Proteins. Plant Physiol. 2016 Dec;172(4):2459-2470. Epub 2016 Oct 26. Lawrence et al. (2010). Recombinant production and purification of the subunit c of chloroplast ATP synthase. Protein Expression and Purification 76: 15-24.

Application example



0.7 µg of purified ATP-synthase complex from Spinacia oleracea (CF0F1) (1), 15.3 µg of purified ATP-synthase complex (CF0F1) from Nicotiana benthamiana (2) and 48.6 µg of Thermosynechococcus elongatus thylakoid preparation (3) were separated on 12% polyacrylamide gel and blotted on PVDF membrane. Filters where blocked (0.5h), incubated with 1: 1000 anti-AtpH antibodies (1h), followed by incubation with 1: 5 000 secondary anti-rabbit antibody (1.25h), coupled to HRP and visualized with chemiluminescence detection reagent.

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