

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

contact: support@agrisera.com

Agrisera AB | Box 57 | SE-91121 Vännäs | Sweden | +46 (0)935 33 000 | www.agrisera.com

Product no AS05 071

Anti-AtpH | ATP synthase subunit c (chloroplastic)

Product information

Immunogen Purified c subunit from Spinacia oleracea UniProt: P69447.

Host Rabbit

Clonality Polyclonal

Purity Serum

Format Lyophilized

Quantity 200 μl

Reconstitution For reconstitution add 200 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.

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

Application information

Recommended dilution 1 : 1000-1 : 10 000 (WB)

Expected | apparent

8 kDa (for Arabidopsis thaliana)

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

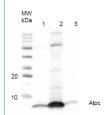
Note that increased incubation at 95°C (20-30 min) prior to loading is recommended to break the multimeric c-mer structure, detection of partial ring structures (e,g, 5 or 6 subunits) may occur

Selected references

Nath et al. (2016). A Nitrogen-Fixing Subunit Essential for Accumulating 4Fe-4S-Containing Photosystem I Core 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.