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Product no AS23 4933

Anti-RbcS I Ribulose bisphosphate carboxylase small subunit (cyanobacterial)

Product information

Immunogen KLH-conjugated peptide derived from protein sequence of RbcS from Synechococcus sp. (strain WH7803), UniProt:

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 1:5000 (WB)

Expected | apparent

12.9 kDa

Confirmed reactivity

Cyanobium sp. PCC 7001, Halothiobacillus neapolitanus, Spinacia oleracea (purified RbcS), Synechocystis sp. PCC6803, Synechococcus sp. PCC 7002 Synechococcus elongatus PCC 7942, Synechococcus sp. PCC 11901

Predicted reactivity

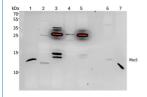
Cyanobacteria

Species of your interest not listed? Contact us

Not reactive in Chlamydomonas reinhardtii, Euglena gracilis

Additional information Antibody reactivity to RbcS in various cyanobacterial strains has to be individually optimized.

Selected references To be added when available, antibody available in November 2025.



samples:

- 1 29 μg of *Synechocystis* sp. PCC6803 whole cell extract.
- 2 14 µg of Cyanobium sp. PCC 7001 whole cell extract.
- 3 19 µg of Synechococcus sp. PCC 7002 whole cell extract.
- 4 14 μg of Synechococcus elongatus PCC 7942 whole cell extract.
- 5 14 ug of Synechococcus sp. PCC 11901 whole cell extract.
- 6 Halothiobacillus neapolitanus whole cell extract.
- 7 6.8 µg of purified Rubisco from spinach leaves.

Note: Protein load/well of respective extract is not the same, therefore differences in intensity of visualized RbcS bands.

About 20 µg/well of total protein extracted freshly from respective cyanobacteria, spinach or Halothiobacillus neapolitanus were extracted with 50 mM Tris-HCl, pH 8.0, 20 mM MgCl2, 20 mM NaHCO3, 0.2 mM EDTA, and denatured with exact buffer components at 95 °C for 10 min. Samples were separated in the cold on 17% SDS-PAGE and blotted for 1 h to PVDF or (pore size of 0.2 µm), using wet transfer in the cold. Blot was blocked with 5 % milk for: 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1: 5 000 for ON/4 °C with agitation. The antibody solution was decanted, and the blot was rinsed briefly twice, then washed once for 15 min and 3 times for 5 min in TBS-T at RT with agitation. Blot was incubated in matching secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1:5 000 in TBS for 1 h/RT with agitation. The blot was washed as above and developed with a following chemiluminescent detection reagent. Exposure time was 1 minute.



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