

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 AS12 2115

Anti-RPL37 | Ribosomal protein L37 (cytoplasmic)

Product information

Immunogen Recombinant, full length RPL37 of Chlamydomonas reinhardtii, UniProt: A8IBG1, expressed in E.coli

Host Rabbit

Clonality Polyclonal

Purity Serum

Format Liquid

Quantity 50 μl

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

10.5 kDa

Application information

Recommended dilution 1:10 000 (WB)

Expected | apparent

Confirmed reactivity Chlamydomonas reinhardtii

Predicted reactivity

Arabidopsis thaliana, Glycine max, Solanum lycopersicum, Oryza sativa, Ostreococcus lucimarinus, Pinus sp., Ricinus communis, Micromonas sp., Volvox carteri

Species of your interest not listed? Contact us

Not reactive in

No confirmed exceptions from predicted reactivity are currently known

Selected references

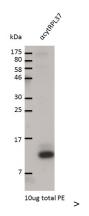
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Couse et al. (2017). Autophagic flux is required for the synthesis of triacylglycerols and ribosomal protein turnover in Chlamydomonas. J Exp Bot. 2017 Oct 19. doi: 10.1093/jxb/erx372.

Couso et al. (2017). Autophagic flux is required for the synthesis of triacylglycerols and ribosomal protein turnover in Chlamydomonas. J Exp Bot. 2017 Oct 19. doi: 10.1093/jxb/erx372.

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Application example



10 µg of total protein from Chlamydomonas reinhardtii extracted with standard lysis buffer (Tris-HCl pH 6.8 50mM, 10mM EDTA, 2% SDS, Sigma protease inhibitors) were separated on 15 % SDS-PAGE and blotted to nitrocellulose membrane using a wet transfer cell. Blot was blocked in TBS-T containing 5% non-fat dry milk for 1h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1: 10 000 in TBS-T containing 5% non-fat dry milk for 1h at RT with agitation. The antibody solution was decanted and the blot was washed 3 times for 15 min in TBS-T containing 1% non-fat dry milk with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG



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horse radish peroxidase conjugated) diluted to 1:10 000 in TBS-T containing 1% non-fat dry milk for 1h at RT with agitation. The blot was washed as above and developed for 1 min with ECL according to the manufacturers instructions. Exposure time was 30 seconds.

Courtesy of Dr. Silvia Ramundo, University of Geneva, Switzerland