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Product no AS16 4082

RPL33 | 50S ribosomal protein L33 (chloroplastic)

Product information

Immunogen KLH-conjugated peptide, derived from Arabidopsis thaliana RPL33 UniProt: P56796, TAIR: AtCq00640

Host Rabbit

Clonality Polyclonal

Purity Immunogen affinity purified serum in PBS pH 7.4.

Format Lyophilized

Quantity 50 μg

Reconstitution for reconstitution add 50 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.

Application information

Recommended dilution 1: 2000 - 1: 5000 (WB)

Expected | apparent

7,6 | 10 kDa

Confirmed reactivity Arabidopsis thaliana, Hordeum vulgare, Zea mays

Predicted reactivity

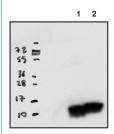
Acacia restiacea, Berberis bealei, Blossfeldia liliputana, Boehmeria spicata, Braya humilis, Calochortus albus, Citrullus colocynthis, Corymbia gummifera, Croomia heterosepala, Croton texensis, Cucurbita pepo, Dodonaea viscosa, Eucalyptus salmonophloia, Fragaria ananassa, Gossypium populifolium, Hippophae rhamnoides, Holodiscus discolor, Khaya senegalensis, Lessertia frutescens, Morus alba var. atropurpurea, Natsiatum herpeticum, Oryza sativa, Pachycladon cheesemanii , Pereskia sacharosa, Polylepis sp., Potentilla lancinata, Ranzania japonica, Ribes fasciculatum var. chinense, Rubus niveus, Solanum lycopersicum, Solanum tuburosum, Sonneratia alba, Trifolium semipilosum, Typha domingensis, Urophysa rockii, Vicia faba , Vicia sepium, Ximenia americana

Species of your interest not listed? Contact us

Not reactive in cyanobacteria

Additional information Ccross reacting bands may appear when total cell extract is analysed,

Application example



10 μg of stromal protein from Arabidopsis thaliana (1) and Zea mays (2) denatured denatured at 95°C 10 minutes were separated on 4-20% SDS-PAGE gradient gel and blotted 1h to PVDF using wet transfer (80V 1 h 30 minutes). Blot was blocked with 5 % milk in TBS-T 1h/RT. Blot was incubated in the primary antibody at a dilution of 1: 2 000 in TBS-T or 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 Agrisera matching secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated AS09 602) diluted to 1:20 000 in for 1h/RT with agitation. The blot was washed as above and developed with chemiluminescent detection reagent and touch exposure was applied.

Courtesy of Dr. Louis-Valentin Mereignier, CNRS, France