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Product no AS15 2868

RpoC1 | RNA polymerase beta' subunit (chloroplast)

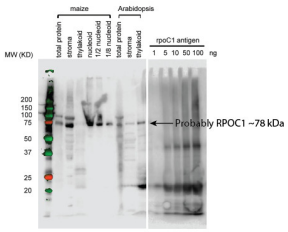
Product information

Immunogen	His-tagged, highly conserved fragment of *Zea mays* RpoC.
Host	Rabbit
Clonality	Polyclonal
Purity	Immunogen affinity purified serum in PBS pH 7.4.
Format	Lyophilized
Quantity	100 µg
Reconstitution	For reconstitution, add 50 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.

Application information

Recommended dilution	1 : 500-1 : 1000 (WB)
Expected | apparent MW	78 | 78 kDa
Confirmed reactivity	*Arabidopsis thaliana, Zea mays*
Predicted reactivity	*Alloteropsis semialata, Aristida purpurea, Chasmanthium latifolium, Chaetobromus involucratus subsp. dregeanus, Chionochloa macra, Coleataenia prionitis, Danthonia californica , Digitaria exilis, Echinochloa oryzicola, Eragrostis tef, Eremitis sp., Isachne distichophylla, Hilaria cenchroides , Hickelia madagascariensis, Microlaena stipoides, Miscanthus sinensis, Neololeba atra, Olmeca reflexa, Oryza sativa, Otatea acuminata, Panicum miliaceum , Pariana campestris, Phragmites australis, Potamophila parviflora, Puelia olyriformis, Rhynchoryza subulata, Saccharum hybrid, Saccharum officinarum, Setaria viridis, Sorghum bicolor, Sorghum timorense, Sporobolus maritimus, Stipa purpurea, Triticum aestivum, Zoysia macrantha* Species of your interest not listed? Contact us
Not reactive in	No confirmed exceptions from predicted reactivity are currently known
Additional information	There may be occur a cross reacting band at 100 kDa in maize samples

Application example



Stroma, thylakoid, nuclei and total protein from *Arabidopsis thaliana* and *Zea mays* were extracted with 50 mM HEPES-KOH pH 8, 330 mM sorbitol exaction buffer. Components were separated on 12% SDS-PAGE and blotted overnight to Nitrocellulose membrane using wet transfer. Blots were blocked with 5% non-fat milk for 1h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1: 500 for 2h at RT 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 secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1:10 000 in for 1h at RT with agitation. The blot was washed as above and developed for 5min with chemiluminescent detection reagent of extreme low femtogram range. Exposure time was 120 seconds.

Courtesy of Jie Shen and Dr. Alice Barkan, University of Oregon, USA