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Product no AS11 1804

Anti-RPB1 | RNA polymerase II subunit B1

Product information

Immunogen KLH-conjugated synthetic peptide derived from known higher plant polymerase II sequences, including Arabidopsis

thaliana P18616, At4g35800

Host Rabbit

Clonality Polyclonal

Purity Immunogen affinity purified serum in PBS pH 7.4.

Format Lyophilized

Quantity 50 ug

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 Storage 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 Purified IgG was lyophilized from PBS pH 7,4 + 0,02 % sodium azide, Therefore only distilled water is required for its

re-constitution

Application information

Recommended dilution 1,2 or 5 μg (ChIP, IP)

Expected | apparent

204 kDa

Predicted reactivity Amygdalus persica, Asparagus officinalis, Brassica rapa, Capsella rubella, Glycine max, Hordeum vulgare, Solanum

lycopersicum, Oryza glaberrima, Panicum italicum, Persea americana, Pisum sativum, Ricinus communis, Secale cereale, Solanum tuberosum, Sorghum vulgare, Triticum aestivum, Vitis vinifera

Species of your interest not listed? Contact us

Not reactive in Physcomitrium patens

Additional information This antibody is not suitable for western blot in denatured conditions

Crawford et al. (2024). The Mediator kinase module enhances polymerase activity to regulate transcriptional memory Selected references

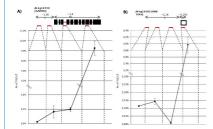
after heat stress in Arabidopsis. EMBO J. 2024 Feb;43(3):437-461. Wang et al. (2020). Close arrangement of CARK3 and PMEIL affects ABA-mediated pollen sterility in Arabidopsis

thaliana. Plant Cell Environ. 2020 Aug 20.doi: 10.1111/pce.13871.

Godoy Herz et al. (2019). Light Regulates Plant Alternative Splicing through the Control of Transcriptional Elongation. Mol Cell. 2019 Jan 15. pii: S1097-2765(18)31035-9. doi: 10.1016/j.molcel.2018.12.005.

Dolata et al. (2015). NTR1 is required for transcription elongation checkpoints at alternative exons in Arabidopsis. EMBO J. 2015 Jan 7. pii: e201489478.

Application example



Chromatin immunoprecipitation (ChIP). Chromatin immunoprecipitation was performed as described (Bowler et al. (2004), Plant Journal) with modifications. Sonic and IP buffer were prepared as described (Kaufmann et al. (2010)), Nature Protocols). Chromatin pellet was sonicated at 4 °C with a Diagenode Bioruptor set at high intensity for 10 min (30 sec on, 30 sec off intervals) to obtain 250-500 bp DNA fragments. Chromatin (20 µg) was immunoprecipitated with a RNA Pol II antibody (2µg/IP) and Dynabeads®Protein G (Life Technologies). Reverse corsslinking and elution of DNA fragments were performed using Chelex (Biorad) at 95°C, 10 min as described (Rowley et al. (2013), Methods). Sample without antibody was used to determine the nonspecific background pulled-down directly by the beads. Relative abundance of regions of interest in



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immunoprecipitated DNA was measured by real-time PCR. Red lines indicates amplified region. Values on the charts are shown as the mean \pm SD level (% of input) of RNA Pol II on GAPDH gene (A) and MIR168a (B) gene. Data comes from three independent experiments.

Courtesy of MSc Jakub Dolata, Adam Mickiewicz University, Poland