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## Product no AS12 1861

## ABI1 | Abscisic acid insensitive 1

## **Product information**

Immunogen KLH-conjugated peptide, derived from *Arabidopsis thaliana* ABI1 sequence UniProt: <u>P49597.</u> TAIR: <u>AT4G26080.</u> Chosen peptide is not present in AtABI2.

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

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 It is of crucial importance to chose a material in which ABI1 protein is highly expressed like seeds or senescent leaf.

This protein could not be detected using this antibody in plants grown under optimal (non stressed) conditions, The

antibody detects both, recombinant and endogenous ABI1 proteins.

## **Application information**

**Recommended dilution** 5 μg (IP for a 200 ul of a cell extract), 3 μg (WB)

Expected | apparent 47.5 kDa

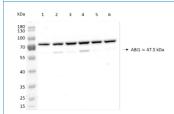
MW 47.5 KL

Not reactive in No confirmed exceptions from predicted reactivity are currently known

Additional information Important note: blocking with more than 3% skimmed milk will result in lack of signal for this antibody

Selected references Mitula et al. (2015). Arabidopsis ABA-Activated Kinase MAPKKK18 is Regulated by Protein Phosphatase 2C ABI1 and the Ubiquitin-Proteasome Pathway. Plant Cell Physiol. 2015 Dec;56(12):2351-67. doi: 10.1093/pcp/pcv146. Epub 2015

Oct 6.



Samples: 1 - 50 μg of *Arabidopsis thaliana* Col0 mock-treated (MG132 50 μM, 6 hours)

- 2 50 μg of Arabidopsis thaliana Col0 ABA-treated (MG132 50μM + ABA 50 μM, 6 hours)
- 3 50 µg of Arabidopsis thaliana ost1(snrk2.6) mock-treated (MG132 5 0µM, 6 hours)
- 4 50 μg of Arabidopsis thaliana ost1(snrk2.6) ABA-treated (MG132 50 μM + ABA 50μM 6, hours)
- 5 50 μg of Arabidopsis thaliana abi1-2 mock-treated (MG132 50 μM, 6 hours)
- 6 50 μg of Arabidopsis thaliana abi1-2 ABA-treated (MG132 50 μM + ABA 50μM, 6 hours)

50 μg/well of total protein extracted freshly from *Arabidopsis thaliana ro*ots with: 150 mM NaCl, 50mM Tris-HCL pH 8, 1% Triton X-100, anti-proteases cocktail (Complete mini EDTA free, "ROCHE") (1 tablet for 10ml), 3 mM DTT, 50 mM MG132, or 50 mM ABA; and denatured with exact buffer components at 95 ° C/5 min. Samples were separated on 10% SDS-PAGE and blotted overnight (ON) to PVDF (Inmobilon®-FL) (pore size of 0.45 μm), using: wet transfer. Blot was blocked with 3% milk for: 6h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1: 10 000 in TBS-T 1X for ON/4 °C with agitation. The antibody solution was decanted, and the blot was rinsed briefly twice, then washed 3 times for 5 min in TBS-T at RT with agitation. Blot was incubated in matching secondary antibody (Goat anti-rabbit IgG HRP conjugated, AS09 602, Agroisera) diluted to 1: 10 000 in for 1h/RT with agitation. The blot was washed as above and developed with a following chemiluminescent detection reagent: Agrisera ECL SuperBright (AS16 ECL-S-10) supplied by Agrisera. Exposure time was 10 seconds.

Courtesy of Drs. Javier Ocaña, Alberto Coego and Pedro L. Rodriguez, CSIC, Spain