

This product is for research use only (not for diagnostic or therapeutic use)

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## Product no AS12 2113

## Anti-AHK2 | Histidine kinase 2

#### **Product information**

Immunogen KLH-conjugated synthetic peptide derived from Arabidopsis thaliana AHK2 protein sequence, TAIR: AT5G35750.

UniProt: Q9C5U2

**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.

## **Application information**

Recommended dilution 1:1000 (WB)

Expected | apparent 131 kDa

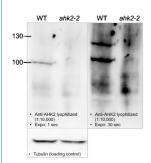
MW 131 KDa

Predicted reactivity Nicotiana benthamiana, Vitis vinifera

Species of your interest not listed? Contact us

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

# application information



The *Arabidopsis thaliana* wild-type (WS) and *ahk2-2* T-DNA insertion mutant plants were grown on MS/2 media for six days. Tissue was collected, frozen in liquid nitrogen, ground in 3 volumes of 2x SDS-PAGE loading buffer and heated at 95 C for 5 min. After 10 min centrifugation at RT, **20** µI of the supernatant was loaded per lane on a **7.5% TGX gel** (BioRad). After separation, proteins were blotted 1 hr onto a supported nitrocellulose membrane. Blots were blocked with 10% milk in PBS-T for 15 min at room temperature (RT) with agitation. Blot was incubated in the primary antibody diluted 1: 10, 000 in 1% milk in PBS-T for 16h at 4C with agitation. The antibody solution was decanted and the blot was washed 3 times for 15 min in PBS-T at RT with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1: 2000 in PBS-T for 1h at RT with agitation. The blot was washed as above and developed for 5 min with chemiluminescent detection reagent according to the manufacturer\s instructions. Chemiluminescent signal was captured using ChemiDoc equipped with a CCD camera.

Courtesy of Dr. Jasmina Kurepa, University of Kentucky College of Agriculture, Food and Environment, USA