

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

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Product no AS14 2783

Anti-SnRK2,2, SnRK2,3, SnRK2,6 | Ser/Thr-protein kinase SnRK

Product information

KLH-conjugated synthetic peptide derived from Arabidopsis thaliana SnRK2.2 UniProt: Q39192, TAIR: At3q50500,

SnRK2.3, UniProt: Q39193 TAIR: At5g66880, SnRK2.6 UniProt: Q940H6, TAIR: At4g33950

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.

Additional information This product can be sold containing proclin if requested

Application information

Recommended dilution 1:1000 (WB)

Expected | apparent

41 kDa

Predicted reactivity Brassica oleracea, Glycine max, Medicago sativa, Morsu notabilis, Oryza sativa, Phaseolus vulgaris, Populus sp., Triticum sp., Vicia faba

Species of your interest not listed? Contact us

Not reactive in Chlamydomonas reinhardtii, Mesotaenium sp., Zygnema sp.

Selected references Lu et al.(2024). N-glycosylation of SnRK2s affects NADPH maintenance in peroxisomes during prolonged ABA

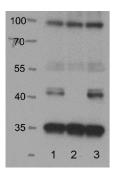
signalling. Nat Commun. 2024 Aug 5;15(1):6630. doi: 10.1038/s41467-024-50720-3.

Belda-Palazón et al. (2020) A dual function of SnRK2 kinases in the regulation of SnRK1 and plant growth. Nat Plants. 2020 Nov;6(11):1345-1353. doi: 10.1038/s41477-020-00778-w. Epub 2020 Oct 19. PMID: 33077877.

Wawer et al. (2018) mRNA Decapping and 5'-3' Decay Contribute to the Regulation of ABA Signaling in Arabidopsis

thaliana. Front Plant Sci. 2018 Mar 12;9:312. doi: 10.3389/fpls.2018.00312.

application example



20 µg of total protein from two-week-old seedling of Arabidopsis thaliana Col-0 (1), snrk2.2/3/6 (2), snrk1/4/5/7/8/9/10 (3) extracted with kinase buffer were separated on 10 % SDS-PAGE and blotted 30 min to PVDF using semi-dry. Blots were blocked with 5% skim milk in TBST for 1h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1: 5000 in 0.5% TBST for overnight at 4 oC with agitation. The antibody solution was decanted and the blot was washed 3 times for 10 min in TBS-T at RT with agitation. Blot was incubated in secondary antibody (anti-rabbit IgG goat peroxidase conjugated) diluted to 1:5 000 in 0.5 % TBST for 1h at RT with agitation. The blot was washed as above and developed for 5 min with ECL according to the manufacturer's instructions. Exposure time was 30 seconds.



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