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contact: support@agrisera.com

Agrisera AB | Box 57 | SE-91121 Vännas | Sweden | +46 035 33000 | www.agrisera.com

product AS12 2110 PIP2-1-7 | plasma membrane aquaporin isoforms 1-7 (C-terminal)

product information

Background	Plasma membrane aquaporin, PIP2;7 is water channel protein required for water transport across cell membrane. Alternative names: plasma membrane intrinsic protein 2-7, AtPIP2;7, plasma membrane intrinsic protein 3, salt stress-induced major intrinsic protein, PIP3a
Immunogen	<u>KLH</u> -conjugated synthetic peptide derived from <i>Zea mays</i> PIP2-7 C-terminal, <u>Q9ATM4</u> , conserved also in <i>Zea mays</i> PIP2-1, UniProt: <u>Q84RL7</u> , PIP2-2, UniProt: <u>Q9ATM8</u> , PIP2-3 (80 % conservation) UniProt: <u>Q9ATM7</u> , PIP2-4 (80 % conservation) UniProt: <u>Q9ATM6</u> , PIP2-5 (70 % conservation) UniProt: <u>Q9ATM5</u> , PIP2-6 (50 % conservation) UniProt: <u>Q9ATM5</u>
Host	Rabbit
Clonality	Polyclonal
Purity	Serum
Quantity	100 μl
Reconstitution	For reconstitution add 100 μ l of sterile water.
Storage	store at -20°C; make aliquots to avoid repeated freeze-thaw cycles. Please, remember to spin tubes briefly prior to opening them to avoid any losses that might occur from material adhering to the cap or sides of the tubes.
Tested applications	western blot (WB)
Related products	AS09 488 PIP2;1 aquaporin PIP2;1
	AS09 488 PIP2;1 aquaporin PIP2;1 AS09 490 PIP2;2 plasma membrane aquaporin 2b AS09 491 PIP2;1,PIP2;2,PIP2;3 plasma membrane intrinistic protein 2-1,2-2, 2-3
	AS09 491 PIP2;1,PIP2;2,PIP2;3 plasma membrane intrinistic protein 2-1,2-2, 2-3
Additional information	Protocol for isolation of plant plasma membrane proteins can be found here.
	This antibody has a potential to work in immunolocalization studies, as it is recognizing C-terminal part of the sequence.

Application information

Recommended dilution	1: 3000 with standard ECL (WB)
Expected apparent MW	30.7 30 kDa (<i>Zea mays</i>)
Confirmed reactivity	Lactuca sativa, Pisum sativum, Solanum lycopersicum, Zea mays
Predicted reactivity	Arabidopsis thaliana, Brassica oleracea, Cicer arietinum, Coffea arabica, Cucumis sativus, Fragaria chiloensis, Glycine max, Medicago trunculata, Mimosa pudica, Nicotiana tabacum, Olea europaea, Phaseolus vulgaris, Pisum sativum, Pyrus communis, Spinacia oleracea, Solanum lycopersicum, Solanum tuberosum, Triticum urartu, Vitis vinifera,monocots: Hordeum vulgare, Oryza sativa, Triticum aestivum, trees: Picea mariana, Populus trichocarpa
Not reactive in	no confirmed exceptions from predicted reactivity known in the moment
Additional information	detection pattern consists of di and monomer of PIP2-7
Selected references	to be added when available, antibody released in May 2012.



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application example



10 µg of total protein from *Zea mays* roots **(1)**, *Phaseolus vulgaris* leaves **(2)** or roots **(3)** extracted with a mixture of 250 mM sorbitol, 50 mM Tris–HCl (pH 8), 2 mM EDTA, and protease inhibitors [1 mM phenylmethylsulfonyl Xuoride, 1 mg ml⁻¹ each of leupeptin, aprotinin, antipain, chymostatin, and pepstatin were separated on 12 % SDS-PAGE and blotted 1h to **PVDF**. Blots were blocked with 5% milk in TBS-T for 2h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1: 3.000 for 1h at RT with agitation. The antibody solution was decanted and the blot was rinsed briefly twice, then washed four times for 5 min in TBS-T at RT with agitation. Blot was incubated in secondary antibody (goat anti-rabbit IgG horse radish peroxidase conjugated, from Agrisera <u>AS09 602</u>) diluted to 1:30 000 in TBS-T for 1h at RT with agitation. The blot was washed as above and developed for 5 min with ECL according to the manufacturers instructions. Exposure time was 60 seconds.

Courtesy of Dr. Ricardo Aroca, CSIC, Spain