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product AS13 2640 ACT | Actin

product information

Background		Actin is a highly conserved protein and an essential component of cell cytoskeleton and plays an important role in cytoplasmic streaming, cell shape determination, cell division, organelle movement and extension growth. Preferentially expressed in young and expanding tissues, floral organ primordia, developing seeds and emerging inflorescence.	
Immunogen		ca. 100 amino acids of recombinant actin conserved more than 80 % in <i>Arabidopsis thaliana:</i> actin-1 <u>P0CJ46</u> <u>AT2G37620</u> , actin-2 <u>Q96292</u> <u>AT3G18780</u> , actin-3 <u>P0CJ47</u> <u>AT3G53750</u> , actin-4 <u>P53494</u> <u>AT5G59370</u> , actin-5 <u>Q8RYC2</u> At2g42100, actin-7 <u>P53492</u> At5g09810, actin-8 <u>Q96293</u> <u>AT1G49240</u> , actin-11 <u>P53496</u> , <u>AT3G12110</u> ,actin-12 <u>P53497</u> <u>AT3G46520</u>	
Host		Rabbit	
Clonality		Polyclonal	
Purity		Serum	
Format		Lyophilized	
Quantity		50 μl	
Reconstitution		For reconstitution add 50 μ l of sterile water.	
Storage		Store lyophilized/reconstituted at -20 °C; once reconstituted 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 lyophilized material adhering to the cap or sides of the tubes.	
Tested applications		immunofluorescence (IF), Western blot (WB)	
Related products		AS10 702 Anti-Actin-11, monclonal mouse antibody AS10 681 Anti-tubulin beta chain, rabbit antibody AS10 680 Anti-tubulin alpha chain, rabbit antibody AS16 3140 Actin-2, mouse monoclonal antibody AS16 3141 Actin-8, mouse monoclonal antibody AS16 3139 Actin-1, monoclonal antibody Plant protein extraction buffer Secondary antibodies	
Additional information	ł	Antibody available in 2 various pack sizes: 50, 100 and 150 ul - plasse, inquire	
		Antibody available in 5 valious pack sizes. 50, 100 and 150 μ - picase, <u>inquire</u> .	
Application inform	Application information		
Recommended dilution		1 : 250 (IF), 1 : 3000-1 : 5000 (WB)	
Expected apparent MW		41.6 45 kDa	
Confirmed reactivity		Agostis stolonifera cv. 'Penncross', Arabidopsis thaliana, Cynara cardunculus, Glycine max, Hordeum vulgare, Solanum tuberosum, Zea mays	
Predicted reactivity		Agropyron cristatum, Beta vulgaris, Betula luminifera, Brassica napus, Brassica rapa subsp. pekinensis, Capsella	

Agropyron cristatum, Beta Vuigaris, Betula luminirera, Brassica napus, Brassica rapa subsp. pekinensis, Capselia rubella, Castanea sativa, Chorispora bungeana, Cucumis sativus, Cyanidioschyzon merolae strain 10D, Glycine max, Glycine soja, Halogeton glomeratus, Medicago truncatula, Malus domestica, Nicotiana tabacum, Oryza sativa, Pisum sativun, Solanum lysopersicum, Solanum tuberosum, Phaseolus vulgaris, Picea abies, Picea sitchensis, Prunus avium, Ricinus communis, Rubus plicatus, Theobroma cacao, Triticum aestivum, Vicia faba

Not reactive in Chlamydomonas reinhardtii (too high background for this species)

Additional information This product can be sold containing ProClin if requested



<u>Jespersen</u> et al. (2017). Metabolic Effects of Acibenzolar-S-Methyl for Improving Heat or Drought Stress in Creeping Bentgrass. Front Plant Sci. 2017 Jul 11;8:1224. doi: 10.3389/fpls.2017.01224. eCollection 2017. (western blot, Agostis stolonifera cv. 'Penncross') <u>Qiu</u> et al. (2015). Soy 14-3-3 protein SGF14c, a new regulator of tolerance to salt–alkali stress. Plant Biotechnology Reports pp 1-9. <u>Shaw</u> et al. (2015). -aminobutyric acid mediated drought stress alleviation in maize (Zea mays L.). Environ Sci

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Buxa et al. (2015). Phytoplasma infection in tomato is associated with re-organization of plasma membrane, ER stacks, and actin filaments in sieve elements. ront Plant Sci. 2015; 6: 650. Published online 2015 Aug 19. Zheng et al. (2014). iTRAQ-based quantitative proteomics analysis revealed alterations of carbohydrate metabolism pathways and mitochondrial proteins in a male sterile cybrid pummelo. J Proteome Res. 2014 May 13.

Application example



15 µg of total protein extracted with PEB (<u>AS08 300</u>) from leaf tissue of (1) *Arabidopsis thaliana*, (2) *Hordeum vulgare*, (3) *Zea mays* were separated on 4-12% NuPage (Invitrogen) LDS-PAGE and blotted 1h to PVDF. Filters were blocked 1h with 2% low-fat **milk powder** in TBS-T (0.1% TWEEN 20) and probed with **anti-actin** (AS13 2640, 1:2500, 1h) and secondary anti-rabbit (1:10 000, 1 h) antibody (HRP conjugated, recommended secondary antibody <u>AS09 602</u>) in TBS-T containing 2% low fat milk powder. Antibody incubations were followed by washings in TBS-T (15, +5, +5, thin). All steps were performed at RT with agitation. Signal was detected with ECL Advance (GE Healthcare) using a Fuji LAS-3000 CCD (300s, standard sensitivity). Exposure time was 2 min.



Actin cytoskeleton in 5 days old *Arabidopsis thaliana* seedlings. Actin signal shown in green, PIN1 in red and DAPI in blue. The material has been fixed in 2 % formaldehyde for 45 minutes. Tissue cleaning has been performed before immunolocalization. Rabbit anti-actin primary antibody was diluted in 1:250 and anti-rabbit Alexa 488 and Alexa 555 were both diluted in 1:500 (Invitrogen). Scale bar - 20 µm.

Courtesy: Dr. Taras Pasternak, Freiburg University, Germany



Proteins were extracted from tuber flesh of Russet Burbank potato (*Solanum tuberosum*) with 0.1 M Tris HCl (pH=8.0), 5% sucrose (m/v), 2% (m/v) SDS, protease inhibitors (PMSF 1mM). Samples were heated 95°C 5 min, and 10 µg of total protein was resolved in 12% SDS PAGE and blotted to PVDF membrane for 1h-1.5h using tank transfer. Blots were blocked with a skimmed milk 4% (m/v) in T-TBS (1.5h) at RT with agitation. Primary antibodies (AS13 2640) were applied overnight +4°C in dilution 1:5000 with agitation. After washing with T-TBS 2-3 times, membrane was incubated with secondary antibodies (Goat Anti-Rabbit HRP conjugate, Transgen biotech HS101) 1:10000 for 1 hour at RT. Blot was washed as above and developed with ECL (Clarity Western ECL Substrate, BioRad, 170-5060) for 5 – 10 minutes. Exposure time – 20.395 seconds.

Courtesy of lauhenia Isayenka, University of Sherbrooke, Canada

This product IS FOR RESEARCH USE ONLY.



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