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# Product no AS08 371 Anti-HSP70 | Heat shock protein 70 (cytoplasmic)

### **Product information**

| Immunogen      | KLH-conjugated synthetic peptide conserved in known higher plant HSC70 proteins including three isoforms of<br><i>Arabidopsis thaliana</i> HSC70-1 UniProt: <u>F4KCE5</u> , HSC70-2 UniProt: <u>A0A178UTH3</u> and HSC70-3 Uniprot: <u>O65719</u> as<br>well as heat shock inducible Hsp70 of <i>Arabidopsis thaliana</i> TAIR: AT3g12580/T2E22_110 and At1g16030 and<br>AT3g12580/T2E22_110 |
|----------------|--|
| Host           | Rabbit   |
| Clonality      | Polyclonal   |
| Purity         | Serum  |
| Format         | Lyophilized  |
| Quantity       | 50 μl  |
| Reconstitution | For reconstitution add 50 $\mu$ 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 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**

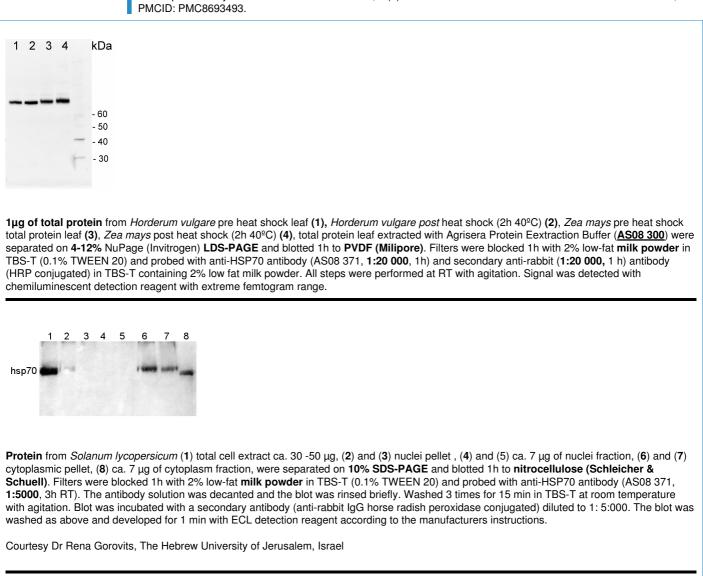
| Recommended dilution      | 1 : 3000 - 1: 10 000, 5 μg protein/well (WB), 2-3 μl/protein extract of concentration 3-5 mg/ml   |
|---------------------------|---|
| Expected   apparent<br>MW | 70 kDa  |
| Confirmed reactivity      | Arabidopsis thaliana, Brassica juncea, Brassica napus, Brassica oleracea, Chlamydomonas reinhardtii,<br>Chlamydomonas sp. UWO241, Cucumis sativus, E. teft, Fagopyrum esculentum, Glycine max, Hordeum vulgare,<br>Medicago sativa, Oryza sativa, Salicornia sp., Silene vulgaris, Solanum lycopersicum, Picrorhiza kurroa, Pinus strobus,<br>Polyscias elegans, Zea mays, algae: Desmodesmus subspicatus, Gracilaria vermiculophylla, phycobiont: Trebouxia<br>TR1 and TR9, Plasmodium falciparum, Setaria italica, Triticum aestivum, Ulva prolifera, Vicia faba  |
| Predicted reactivity      | Ageratina adenophora, Allium sativum, Arabis alpina, Arachis diogoi, Arundo donax, Brassica napus, brassica rapa<br>subsp. pekinensis, Camellia sinensis, Chlorella vulgaris, Citrus clementina, Citrus sinensis, Coffea arabica, Eriobotrya<br>japonica, Gossypium arboretum, Glycine soja, Helianthus annuus, Hordeum vulgare var. distichum, Lotus japonicus,<br>Medicago sativa, medicago truncatula, Musa acuminata subsp. malaccensis, Nannochloropsis gaditana, Nicotiana<br>tabacum, Nicotiana bethamiana, Olea europea, Phaseolus vulgaris, Physcomitrium patens, Pinus taeda, Pisum<br>sativum, Populus balsamifera, Populus trichocarpa, Saccharum sp, Salix gilgiana, Saussurea medusa,Solanum<br>tuberosum, Solanum commersonii, Spinacia oleracea, Tragopogon dubius, Tragopogon porrifolius, Triticum aestivum,<br>Vitis vinifera, Volvox sp.<br>Species of your interest not listed? <u>Contact us</u>  |
| Not reactive in           | Centella asiatica, Polyscias elegans  |
| Additional information    | Can be sold containing 0.1% ProClin if requested  |
|                           | This antibody can be used as a marker of cytoplasmic fraction in tomato (Anfoka et al. 2015).   |
|                           | Applied primary antibody dilution in western blot depends upon sensitivity of detection reagents (pico or femtogram for chemiluminescent detection).  |
|                           | Immunoprecipitation protocol using Agrisera anti-Hsp70 cytosolic antibodies, see tab: protocols.  |
| Selected references       | Tanaka et al. (2024). Heat shock protein 70 is associated with duration of cell proliferation in early pod development of soybean. Commun Biol. 2024 Jun 21;7(1):755. doi: 10.1038/s42003-024-06443-8.<br><u>Vuković</u> et al. (2024). Heat Priming Modifies Heat Stress Response in BPM1-Overexpressing Arabidopsis thaliana (L.)<br>Heynh. J Olant Growth Regulation. https://doi.org/10.1007/s00344-024-11337-4.<br><u>Llamas</u> et al. (2023). In planta expression of human polyQ-expanded huntingtin fragment reveals mechanisms to<br>prevent disease-related protein aggregation. Nat Aging. 2023 Nov;3(11):1345-1357.doi: 10.1038/s43587-023-00502-1.<br><u>Chong</u> et al. (2022) The tomato OST1-VOZ1 module regulates drought-mediated flowering. Plant Cell. 2022 Apr<br>26;34(5):2001-2018. doi: 10.1093/plcell/koac026. PMID: 35099557; PMCID: PMC9048945.<br><u>Bychkov</u> et al. (2022) The role of PAP4/FSD3 and PAP9/FSD2 in heat stress responses of chloroplast genes. Plant Sci.<br>2022 Sep;322:111359. doi: 10.1016/j.plantsci.2022.111359. Epub 2022 Jun 20. PMID: 35738478.<br><u>Cvetkovska</u> et al. (2022) A constitutive stress response is a result of low temperature growth in the Antarctic green alga<br>Chlamydomonas sp. UWO241. Plant, Cell & Environment, 45, 156– 177. https://doi.org/10.1111/pce.14203 |

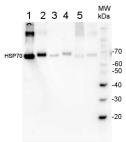


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Wang et al. (2022) 17-(Allylamino)-17-demethoxygeldanamycin treatment induces the accumulation of heat shock proteins and alleviates senescence in broccoli. Postharvest Biology and Technology,Volume 186, 2022, 111818, ISSN 0925-5214, https://doi.org/10.1016/j.postharvbio.2021.111818. Kumari et al. (2021) In-depth assembly of organ and development dissected Picrorhiza kurroa proteome map using mass spectrometry. BMC Plant Biol. 2021 Dec 22;21(1):604. doi: 10.1186/s12870-021-03394-8. PMID: 34937558;





200 fmoles of HSP70 protein standard product number AS08 371S (1), 1 µg of total protein from samples such as *Lycopersicum esculentum* leaf (2), *Nicotiana tabaccum* leaf, (3), *Zea mays* leaf (4), *Hordeum vulgare* leaf (5), *Arabidopsis thaliana* leaf (6) were extracted with Agrisera Protein Extraction Buffer PEB (AS08 300). Samples were diluted with 1X sample buffer (NuPAGE LDS sample buffer (Invitrogen) supplemented with 50 mM DTT and heat at 70 °C for 5 min and keept on ice before loading. Protein samples were separated on 4- 12% Bolt Plus gels, LDS-PAGE and blotted for 70 minutes to PVDF using tank transfer. Blots were blocked immediately following transfer in 2% blocking reagent or 5% non-fat milk

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dissolved in 20 mM Tris, 137 mM sodium chloride pH 7.6 with 0.1% (v/v) Tween-20 (TBS-T) for 1h at room temperature with agitation. Blots were incubated in the primary antibody at a dilution of 1: 10 000 (in blocking reagent) for 1h/RT with agitation. The antibody solution was decanted and the blot was rinsed briefly twice, and then washed 1x15 min and 3x5 min with TBS-T at room temperature with agitation. Blots were incubated in secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated, recommended secondary antibody <u>AS10 1489</u>, Agrisera) diluted to 1:25 000 in blocking reagent for 1h at room temperature with agitation. The blots were washed as above. The blot was developed for 5 min with chemiluminescence detection reagent in extreme femtogram range, according the manufacturers instructions. Images of the blots were obtained using a CCD imager (VersaDoc MP 4000) and Quantity One software (Bio-Rad). Exposure time was 30 seconds.