

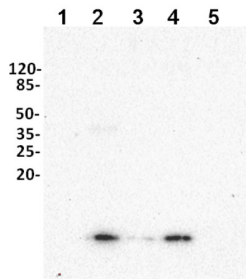
Product no **AS08 307A****Anti-UBQ11 | Ubiquitin (affinity purified)****Product information**

Immunogen	Recombinant <i>Arabidopsis thaliana</i> <u>ubiquitin</u> , ubq11, with N-terminal His-tag; TAIR: At4g05050
Host	Rabbit
Clonality	Polyclonal
Purity	Immunogen affinity purified serum in PBS pH 7.4.
Format	Lyophilized
Quantity	50 µg
Reconstitution	For reconstitution add 20 µl of sterile water to each tube before use.
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	Ubiquitin mutant is lethal, therefore could not be used in validation of this antibody.

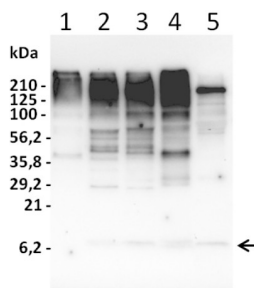
Application information

Recommended dilution	1 : 10 000 (WB)
Expected apparent MW	8.5 kDa
Confirmed reactivity	<i>Arabidopsis thaliana</i> , <i>Hordeum vulgare</i> , <i>Lupinus angustifolius</i> , <i>Malus domestica</i> , <i>Nicotiana benthamina</i> , <i>Oryza sativa</i> , <i>Pisum sativum</i> , <i>Solanum tuberosum</i> , <i>Zea mays</i> , <i>Vitis vinifera</i>
Predicted reactivity	<i>Brachypodium distachyon</i> , <i>Brassica napus</i> , <i>Capsella rubella</i> , <i>Cochliobolus heterostrophus</i> , <i>Citrus clementina</i> , <i>Citrus sinensis</i> , <i>Coffea canephora</i> , <i>Cymbidium faberi</i> , <i>Eucalyptus grandis</i> , <i>Erythraea guttata</i> , <i>Glycine max</i> , <i>Glycine soja</i> , <i>Ipomoea batatas</i> , <i>Medicago truncatula</i> , <i>Magnaporthe oryzae</i> , <i>Nicotiana tabacum</i> , <i>Phaseolus vulgaris</i> , <i>Populus trichocarpa</i> , <i>Prunus persica</i> , <i>Pyrus communis</i> , <i>Ricinus communis</i> , <i>Solanum nigrum</i> , <i>Sorghum bicolor</i> , <i>Triticum aestivum</i> , <i>Theobroma cacao</i> , <i>Vitis vinifera</i>
	Species of your interest not listed? Contact us
Not reactive in	Algae
Additional information	Antibody can be used to check ubiquitination status in the whole plant extracts.
	Technical note: It is very difficult to detect ubiquitin monomers in total cell extracts due to a great abundance of poly and multi-ubiquitinated proteins. Recommended is size separation of protein extracts before gel electrophoresis focused on good resolution of region between 6-10 kDa.
	Suggested extraction buffer: 100 mM Tris-HCl, pH 8.0, 0.1% [w/v] SDS, 0.5% [w/v] sodium deoxycholate, 1% [v/v] glycerol, 50 mM sodium metabisulfite, 20 mM N-ethylmaleimide (NEM) and protease inhibitor cocktail (Roche) (Orosa et al. 2018). This buffer will help to stabilize the conjugates and will help to detect any increase or decrease in conjugate accumulation using the antibodies.
Selected references	Lj et al. (2024) . Stress granules sequester autophagy proteins to facilitate plant recovery from heat stress. <i>Nat Commun</i> . 2024 Dec 30;15(1):10910. doi: 10.1038/s41467-024-55292-w. Xie et al. (2024) . Proteasome resides in and dismantles plant heat stress granules constitutively. <i>Mol Cell</i> . 2024 Sep 5;84(17):3320-3335.e7. doi: 10.1016/j.molcel.2024.07.033. Jeran et al. (2021) The PUB4 E3 Ubiquitin Ligase Is Responsible for the Variegated Phenotype Observed upon Alteration of Chloroplast Protein Homeostasis in Arabidopsis Cotyledons. <i>Genes (Basel)</i> . 2021 Sep 6;12(9):1387. doi: 10.3390/genes12091387. PMID: 34573369; PMCID: PMC8464772. Filippi et al. (2019) . Caspase-3-like activity and proteasome degradation in grapevine suspension cell cultures undergoing silver-induced programmed cell death. <i>J Plant Physiol</i> . 2019 Feb;233:42-51. doi: 10.1016/j.jplph.2018.12.003. Krasuska et al. (2016) . Nitric oxide-polyamines cross-talk during dormancy release and germination of apple embryos. <i>Nitric Oxide</i> . 2016 Nov 23. pii: S1089-8603(16)30178-1. doi: 10.1016/j.niox.2016.11.003. [Epub ahead of print]

Application example



The 10 µg of soluble protein extracts from 3 -week old *Solanum tuberosum* (1), one -week old seedlings of *Lupinus angustifolius* (2), *Pisum sativum* (3), *Zea mays* L. (4) and *Hordeum vulgare* (5) leaves were extracted with buffer containing 50 mM HEPES- KOH, pH 7.5, 330 mM sorbitol, 2 mM EDTA, 1 mM MgCl₂, 5 mM ascorbate, 0.05% BSA. All samples were mixed with sample buffer, denatured for 5 min at 70°C and separated on 10% Tricine -SDS -PAGE. Proteins were blotted 30 min to Hybond™ -PVDF membrane (Amersham Life Science) using semi -transfer (Bio -Rad) in standard transfer buffer in presence of 20% methanol. Transfer of proteins to the membrane was checked using 1% Ponceau S staining before the blocking step. Blots were blocked in buffer (5% low -fat milk in 1xPBS, 0.1% Tween) with for 60 min at room temperature (RT) with agitation. Blots were incubated in the primary antibody at a dilution of 1: 5 000 at RT with agitation. The antibody solution was decanted and the blot was rinsed briefly twice, then washed once for 15 min and 3 times for 5 min in P BS -T at RT with agitation. Blot was incubated in secondary antibody (goat anti -rabbit IgG, [AS09 602](#), Agrisera) diluted to 1 :50 000 for 1h at RT with agitation. The blot was washed as above and developed for 5 min with Clarity Western ECL Substrate and ChemiDoc detection system (Bio-Rad).



Total cell extracts of *Solanum tuberosum* (1), *Lupinus angustifolius* (2), *Pisum sativum* (3), *Zea mays* L. (4), *Hordeum vulgare* (5), were separated in conditions as described above. Marker - P restrained SDS -PAGE (Bio Rad, kat # 1610318). Monomers are marked by arrow. Exposure 30 seconds. For visualization of monomers, longer exposure time was applied.

Courtesy of Dr. Elena Poznidava Laboratory of Plant Gene Expression, Timiryazev Institute of Plant Physiology RAS, Moscow, Russia