

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

contact: support@agrisera.com

Agrisera AB | Box 57 | SE-91121 Vännäs | Sweden | +46 (0)935 33 000 | www.agrisera.com

Product no AS23 4886

Anti-UBC7/13/14 | Ubiquitin-conjugating enzyme E2 7/13/14

Product information

Immunogen KLH-conjugated peptide derived from Arabidopsis thaliana UBC7 UniProt: Q42540, TAIR: AT5G59300, UBC13:

UniProt: **Q42541**, TAIR: <u>AT3G46460</u> UBC14: UniProt: <u>F4IWU7</u> TAIR: <u>AT3G55380</u>

Host Rabbit

Clonality Polyclonal

Purity Antigern affinity purified serum, in PBS pH 7.4

Format Lyophilized

Quantity 50 μg

Reconstitution For reconstitution add 50 μl, of sterile or deionized 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.

Application information

Recommended dilution 1:1000 - 1:5000 (WB)

Expected | apparent MW 18.8 kDa

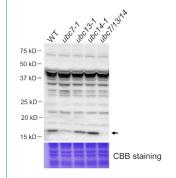
Confirmed reactivity | Arabidopsis thaliana

Predicted reactivity Brassica napus, Nicotiana tabacum, Solanum lycopresicum, Solanum tuberosum, Pisum sativum

Species of your interest not listed? Contact us

Not reactive in No confirmed exceptions from predicted reactivity are currently known

Selected references To be added when available, antibody available in November 2024.



Samples:

- 1 30 μg total proteins of Arabidopsis thaliana 10-day-old seedlings; wild type
- 2 $30~\mu g$ total proteins of $\textit{Arabidopsis thaliana}\ 10\mbox{-day-old seedlings};\ ubc7\mbox{-}1$
- 3 30 µg total proteins of Arabidopsis thaliana 10-day-old seedlings; ubc13-1
- 4 30 μg total proteins of Arabidopsis thaliana 10-day-old seedlings; ubc14-1
- 5 30 μg total proteins of Arabidopsis thaliana 10-day-old seedlings; ubc7/13/14

MW markers: BioRad Precision Plus Protein standards (bands indicated at the left)

 $30 \mu g/lane)$ of total proteins extracted freshly from 10-day-old seedlings of *Arabidopsis thaliana* wt and mutants, with the exaction buffer (50 mM Tris-HCl pH 8.0, 200 mM NaCl, 10 mM DTT, 1% (v/v) Triton X-100, Sigma protease inhibitor cocktail) and denatured with 4X SDS sample buffer at 95°C for 5 min. Samples were separated using 10% SDS-PAGE and transferred onto PVDF membrane (0.2 μ m pore size), using wet transfer. After blocking with 5% milk in PBS-T for 0.5h at RT (room temperature) with agitation, the blot was incubated in the primary antibody at a dilution of 1: 5000 (from the initial antibody solution at 1 μ g IgG/ μ l) in PBS-T at 4°C with agitation. The antibody solution was decanted, and the blot was washed 4 times, each for 10 min, in PBS-T at RT with agitation. The blot was incubated in the secondary antibody goat anti-rabbit IgG horse radish peroxidase conjugated, 1:10 000 for 1h at RT with agitation. It was washed and developed with Agrisera ECL SuperBright.

Note: Background signal can be decreased with further protocol adjustment.



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Courtesy of Dr. Hong Wang,

University of Saskatchewan, Canada