

Product no **AS16 4069****UVR3 | Protein UV repair defective 3****Product information**

Immunogen | KLH-conjugated peptide derived from *Arabidopsis thaliana* UVR3 protein sequence, UniProt: [Q48652-1](#), TAIR: [AT3G15620](#)

Host | Rabbit

Clonality | Polyclonal

Purity | Affinity purified serum, in PBS pH 7.4

Format | Lyophilized

Quantity | 50 µg

Reconstitution | For reconstitution add 50 µl, 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.

Additional information | Reactivity of this antibody on endogenous material remains to be determined. UVR3 is a low abundance protein, therefore use of specific cellular fraction or concentration through [TCA/acetone precipitation](#) is recommended.

Application information

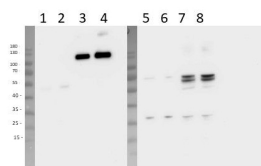
Recommended dilution | 1 : 1000 (WB)

Expected | apparent MW | 63.7 kDa

Confirmed reactivity | *Arabidopsis thaliana* UVR3-GFP

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

Selected references | To be added when available, antibody available in September 2022.



Samples: 1,2 - Cell lysate from *Arabidopsis thaliana* leaves, overexpressing UVR3 fused to GFP.

3,4 - UVR3-GFP immunoprecipitated using GFP-Trap Agarose beads (Chromotec) from *Arabidopsis thaliana* leaves overexpressing UVR3 fused to GFP Cell lysate from E.coli expressing recombinant protein UVR3 with his-tag. 5,6 – before induction
7,8 – after induction, and production of protein

Samples were separated on 12% SDS-PAGE, blotted 0,5h using semi-dry transfer. Nitrocellulose membrane were blocked with 0,5% milk for 1h. Blot was incubated in the primary antibody diluted 1:1 000 ON/4°C. After rising blot was incubated with secondary antibody goat anti rabbit IgG HRP conjugated (Agrisera. [AS09 602](#)) diluted 1:10 000. Detection: chemiluminescence. Mass Marker: Page Ruler Protein Ladder.

Courtesy of Dr. Justyna Łabuz Laboratory of Photobiology Malopolska Centre of Biotechnology Jagiellonian University, Kraków, Poland