Agrisera
Part of Olink® Group

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 AS16 4106

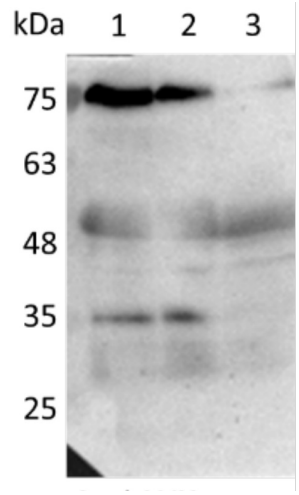
LUX | Transcription factor LUX

Product information

Immunogen	KLH-conjugated synthetic peptide derived from *Arabidopsis thaliana* LUX protein Q9SNB4, At3g46640
Host	Rabbit
Clonality	Polyclonal
Purity	Antigen 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.

Application information

Recommended dilution	1: 2000 (WB)
Expected | apparent MW	34.7 kDa
Confirmed reactivity	*Arabidopsis thaliana*
Predicted reactivity	*Brassica napus, Camelina sativa, Capsella rubella, Eutrema salsugineum, Raphanus sativus* 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 released in May 2023.



Samples:
1- 50 µg pEG202-LUX/Col-0 (FLAG-tag)
2- 50 µg Col-0
3- 50 µg lux mutant (SALK_119768) Expected product size: LUX (35kDa) and FLAG-LUX (35+1=36kDa)
All samples were 7-day-old seedlings grown under 22°C 16 light/8 dark.

50 µg/well of total protein extracted freshly from *Arabidopsis thaliana* seedlings. Exact buffer components were: and denatured with 100 mM Tris-HCl, pH7.8, 4 M Urea, 5% SDS, 15% glycerol, protease inhibitor cocktail, -Mercaptoethanol, and bromophenol blue dye at 100°C/10 min. Samples were separated in the cold on 10% SDS-PAGE and blotted to Immobilon®-P PVDF Membrane PVDF (pore size of 0.45 µm), using: wet transfer in the cold. Blot was blocked with 5 % milk 4°C/ON with agitation. Blot was incubated in the primary antibody at a dilution of 1:2000 for 1h with agitation in TBS-T 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 TBS-T at RT with agitation. Blot was incubated in matching secondary antibody (anti-rabbit HRP conjugated) diluted to 1:25

000 in TBS-T for 1h/RT with agitation. The blot was washed as above and developed with a chemiluminescent detection reagent: Agrisera Super Bright (AS16 ECL-S-10). Exposure time was 5 sec.

Courtesy of Dr. Chin-Mei Lee, Institute of Plant Biology Global Agriculture Technology and Genomic Science Master Program, National Taiwan University, Taiwan