

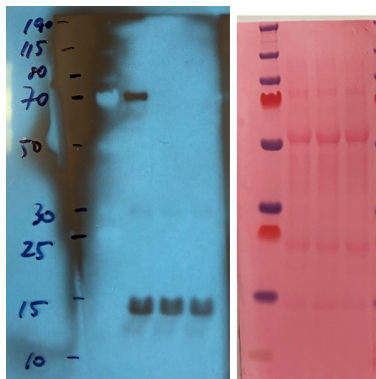
Product no **AS18 4221**
CRY2 | Cryptochrome 2

Product information

| | |
|-----------------------|---|
| Immunogen | Part of <i>Arabidopsis thaliana</i> CRY2 protein sequence, UniProt:Q96524, TAIR: At1g04400 Antigen used to elicit CR2 antibodies is not conserved in CRY1 protein. |
| Host | Rabbit |
| Clonality | Polyclonal |
| Purity | Immunogen 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 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. |

Application information

| | |
|-------------------------------|---|
| Recommended dilution | 1 : 1000 (WB) |
| Expected apparent MW | 69.5 70 kDa |
| Confirmed reactivity | <i>Arabidopsis thaliana</i> |
| Predicted reactivity | <i>Camelina sativa</i> , <i>Capsella rubella</i> Species of your interest not listed? Contact us |
| Not reactive in | <i>Poplar</i> sp. |
| Selected references | To be added when available, antibody available in January 2022. |



Samples:

- 50 µg of *Arabidopsis thaliana* whole seedlings (7 days old)
 1 - *cry1-304* (overexpressing line)
 2 - *cry2-1* (deletion mutant of both isoforms)
 3 - *cry1-304 cry2-1* (substitution mutant)

Mark: MW markers: PageRuler™ Plus Prestained Protein Ladder from Thermo Scientific™ (26619)

8 µl/well of total protein extracted freshly from 50 µg of *Arabidopsis thaliana* seedlings (7-day WL grown) with 8M Urea, 0.35M Tris-Cl (pH 7.5), 1× protease inhibitor cocktail, and denatured with 6x SDS buffer at 95°C for 10 min. Samples were separated on 4-12 % gradient SDS-PAGE (Invitrogen) and transferred 7 min to nitrocellulose (pore size of 0.2 µm), using transfer device (iBlot2, Thermofischer). Blot was blocked with 2 % milk 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1 : 1,000 for ON/4°C without agitation. The antibody solution was decanted and the blot was washed 3 times for 10 min in TBS-T at RT with agitation. Blot was incubated in Agrisera matching secondary antibody (goat anti-rabbit HRP conjugated [AS09 602](#)) diluted to 1:25 000 in for 3h/RT with agitation. The blot was washed as above and developed with AgriseraECLSuperBright [AS16 ECL-S-100](#), using Summit QCP. Exposure time was 5 seconds.



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

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Courtesy of Dr. Sanghwa Lee, Salk Institute for Biological Studies, USA