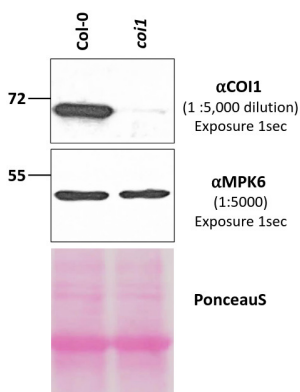


Product no **AS12 2637****COI1 | Coronate insensitive 1 (rabbit antibody)****Product information**

Immunogen	KLH-conjugated synthetic peptide derived from <i>Arabidopsis thaliana</i> COI1 protein, UniProt: O0419Z , TAIR: At2g39940
Host	Rabbit
Clonality	Polyclonal
Purity	Immunogen affinity purified serum in PBS pH 7.4.
Format	Lyophilized
Quantity	50 µg
Reconstitution	For reconstitution add 25 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	70 kDa
Confirmed reactivity	<i>Arabidopsis thaliana</i>
Predicted reactivity	<i>Brassica rapa</i> , <i>Glycine max</i> , <i>Lupinus luteus</i> , <i>Medicago tribuloides</i> , <i>Nicotiana tabacum</i> , <i>Oryza sativa</i> , <i>Pisum sativum</i> , <i>Populus trichocarpa</i> , <i>Ricinus communis</i> , <i>Solanum lycopersicum</i> , <i>Vitis vinifera</i> Species of your interest not listed? Contact us
Not reactive in	No confirmed exceptions from predicted reactivity are currently known
Additional information	Antibody detects 100 ng of recombinant COI1 protein, expressed in Hi5 suspension insect cells. The peptide used to elicit this antibody is partially conserved in <i>Zea mays</i> : ZmCOI1a, ZMCOI1b, ZMCO1c and ZmCOI2.
Selected references	Agrawal et al. (2022) MEDIATOR SUBUNIT17 integrates jasmonate and auxin signaling pathways to regulate thermomorphogenesis. Plant Physiol. 2022 Aug 1;189(4):2259-2280. doi: 10.1093/plphys/kiac220. PMID: 35567489.



Total proteins were isolated from 7-day old *Arabidopsis thaliana* Col-0 (wild-type) and *coi1* (null mutant) seedlings according to [ref 1]. Proteins were denatured at 65°C for 5 min. 30 µg of proteins were separated on a 10% SDS-PAGE and transferred for 70 min at 55V using a tank transfer system to nitrocellulose membrane. Blots were blocked with 1x PBS + 0.1% Tween 20 (PBS-T) + 5% milk for 1hr at room temperature (RT) with agitation. Blots were incubated in primary α -COI1 antibody (AS12 2637) diluted to 1:5,000 in PBS-T+5% milk at 4°C overnight with agitation. The primary antibody solution was decanted, and the blots were washed four times (6-8 minutes each) in 1x PBS-T at RT with agitation prior to incubation with secondary antibody Goat anti Rabbit IgG (H&L) – HRP conjugated ([AS09 602-trial](#)) diluted to 1:20,000 in 1x PBS-T + 5% milk for 2hr at RT with agitation. The blots were washed as above and developed for 4 min with chemiluminescent detection reagent ECL Bright ([AS16 ECL-N](#)). Exposure times (in seconds) to X-ray films were done as indicated. PonceauS stained membrane and membrane portion probed with α -MPK6 served as loading controls [ref 1].

Courtesy of Alani Antoine-Mitchell and Antje Heese, University of Missouri, Div. Biochemistry, IPG (USA)

References: [1] LaMontagne, ED., Collins, CA., Peck, S.C., and Heese, A. 2016. Isolation of microsomal membrane proteins from *Arabidopsis thaliana*. *Curr. Protoc. Plant Biol.* 1:217-234. doi: 10.1002/cppb.20020