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

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# Product no AS12 1856 Anti-SOC1 | Suppressor of constans overexpression 1

#### **Product information**

Immunogen	KLH-conjugated synthetic peptidederived from Arabidpsis thaliana SOC1. UniProt: O64645, TAIR: AT2G45660
Host	Rabbit
Clonality	Polyclonal
Purity	Immunogen affinity purified serum in PBS pH 7.4.
Format	Lyophilized
Quantity	50 µg
Reconstitution	For reconstitution add 50 $\mu$ l 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 : 5000 (WB)
Expected   apparent MW	24 kDa
Confirmed reactivity	Arabidopsis thaliana
Predicted reactivity	<i>Brassica sp., Cardamine sylvatica, Sinapsis juncea</i> Species of your interest not listed? <u>Contact us</u>
Not reactive in	No confirmed exceptions from predicted reactivity are currently known
Selected references	Cuerda-Gil et al. (2022) A plant tethering system for the functional study of protein-RNA interactions in vivo. Plant Methods. 2022 Jun 4;18(1):75. doi: 10.1186/s13007-022-00907-w. PMID: 35658900; PMCID: PMC9166424.

## application example

2 μg of *Arabidopsis thaliana* total protein from 15 days old seedlings **(1)** from 200ng/μl extracted with buffer containing Tris (pH7.5), NaCl, Triton-x100 and protease inhibitors and yeast protein extracts from SG335 cells transformed with pGADT7/SOC1 construct **(2)** were separated on 12 % SDS-PAGE and blotted 40 min. to PVDF. Blots were blocked with 5% milk for 1h at room temperature (RT) with agitation. Blot was incubated in the primary antibody at a dilution of 1:5000 for 1.5h at RT 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 secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1:10 000 in for 45min at RT with agitation. The blot was washed as above and developed for 5 min with ECL according to the manufacturer's instructions. Exposure time was 7 min.

1. 2. 3.

Double band in the yeast sample might be due to post-translational mofifications as experiment with yeast harbouring vector showed no cross-reactivity.

Courtesy of Dr Theoni Margaritopoulou, Agricultural University of Athens, Greece