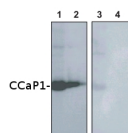


Product no **AS09 483****Anti-AtCCaP1 | vacuolar calcium-binding protein-related****Product information**

<b>Immunogen</b>	KLH-conjugated synthetic peptide derived from <i>Arabidopsis thaliana</i> CCaP1m UniProt: <a href="#">Q9SXE9</a> , TAIR: <a href="#">At1g62480</a>
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Purity</b>	Serum
<b>Format</b>	Lyophilized
<b>Quantity</b>	100 µl
<b>Storage</b>	Store lyophilized/reconstituted at -20°C; 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.
<b>Additional information</b>	0.1 % sodium azide is added as preservative. For antibody re-suspending information check the tube label.  Antibodies will detect target protein in a few µg of a crude preparation loaded per well. If purified preparations of vacuolar and plasma membranes are used, one µg load per well should be sufficient.

**Application information**

<b>Recommended dilution</b>	1 : 8000 (ELISA), 1 : 1000 (WB)
<b>Expected   apparent MW</b>	16.6 kDa
<b>Confirmed reactivity</b>	<i>Arabidopsis thaliana</i>
<b>Predicted reactivity</b>	<i>Arabidopsis thaliana</i>
<b>Not reactive in</b>	No confirmed exceptions from predicted reactivity are currently known
<b>Additional information</b>	Protein or membrane sample should be treated at 70°C for 10 min before loading on the gel.  Diluted antibody solution can be used 2 to 3 times within one month if it contains 0.1 % sodium azide as preservative and is stored at -20°C to -80°C.
<b>Selected references</b>	<a href="#">Ide et al. (2007)</a> . Transcriptional Induction of Two Genes for CCaPs, Novel Cytosolic Proteins, in <i>Arabidopsis thaliana</i> in the Dark. <i>Plant Cell Physiol.</i> 1:54-65.

**Application example**

5 µg (**1,3**) and 2.5 µg (**2,5**) of lysate from E.coli cells expressing CCaP1 protein/lane were separated on 12 % **SDS-PAGE** and blotted 1h to **PVDF membrane** (40 min. at 10 V using BioRad semidry transfer). Filters were blocked 1h with 5 % low-fat **milk powder** in TBS-T (0.05% Triton X.100). Membranes were washed 5 times with TBS-T, each time in a fresh polystyrene box and probed with anti-CCaP1 antibodies (AS09 483, **1:1000**, 1h) and secondary anti-rabbit (**1:2000**, 1 h). All steps were performed in RT with agitation.