

# Agrisera

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Product no **AS20 4420**

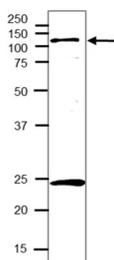
## NAI2 TSA1-like protein, C-terminal

### Product information

<b>Immunogen</b>	Purified recombinant C-terminal part of NAI2 of <i>Arabidopsis thaliana</i> , residues 636-772 with a His tag, UniProt: <a href="#">Q9LSB4</a> , TAIR: <a href="#">At3g15950</a>
<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Purity</b>	Total IgG, purified on Protein A
<b>Quantity</b>	200 µg
<b>Storage</b>	Store at -20°C; once 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 material adhering to the cap or sides of the tubes.
<b>Additional information</b>	Signal peptide of 24 amino acids is removed from the mature protein.

### Application information

<b>Recommended dilution</b>	1: 2000 - 1: 4000 (WB)
<b>Expected   apparent MW</b>	85   120 kDa
<b>Confirmed reactivity</b>	<i>Arabidopsis thaliana</i>
<b>Predicted reactivity</b>	Species of your interest not listed? <a href="#">Contact us</a>
<b>Not reactive in</b>	No confirmed exceptions from predicted reactivity are currently known.
<b>Selected references</b>	<a href="#">Yamada et al. (2008)</a> . NAI2 is an endoplasmic reticulum body component that enables ER body formation in <i>Arabidopsis thaliana</i> . <i>Plant Cell</i> . 2008 Sep;20(9):2529-40. doi: 10.1105/tpc.108.059345.



*Arabidopsis thaliana* 7 day seedling extract was freshly extracted with 2x SDS-sample buffer (+ 2ME) for SDS-PAGE and denatured with 4X SDS buffer at 95°C for 5 min. Sample was separated on 12.5 % SDS-PAGE and blotted at 15V overnight using wet transfer to PVDF membrane. Blot was blocked with 3 % skim milk/TBS-T, 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1: 2000 in TBS-T for 1h/RT. The antibody solution was decanted and the blot was washed 4 times for 10 min in TBS-T at RT with agitation. Blot was incubated in matching secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated) diluted to 1:10 000 in for 1h/RT with agitation. The blot was washed as above and developed with a chemiluminescent detection reagent, following manufacture's recommendation.

The nature of LMW band was not investigated.