Anti-NAI2 ΔSP (At) antibody, rabbit polyclonal

Product code	81-103
Size	200 µg
Storage	-20℃
Concentration	2.0 mg/ml
Buffer	PBS- with 50% glycerol
Purity	Purified IgG fraction with protein A from rabbit antiserum.
Immunogen	Purified recombinant His6-tagged NAI2 (amino acids 272-502) protein lacking signal peptide (*A. thaliana*).
Isotype	Rabbit IgG
Reactivity	NAI2 protein of Arabidopsis thaliana. Not tested in other species.
Special notes	Validation: Specific reactivity has been validated by western blot using *nai1* mutant extracts.
Application	1. Western blotting (1/2,000-1/4,000) 2. Immunofluorescence staining (1/1,000-1/3,000)
Background	Responsible for the ER body formation. Regulates the number and shape of the ER bodies and the accumulation of PYK10 in ER bodies, but is not involved in the expression of PYK10. Interacts directly or indirectly with MEB1 and MEB2. Expressed in roots. Detected in shoot apex. Induced by NAI1.Lenght:772: amino acids. Predicted molecular mass: 85,016 Subcellular location: Endoplasmic reticulum lumen. Modification: Elimination of 24-amino acid signal peptide from N-terminus.
Data Link	UniProtKB Q9LSB4(NAI2_ARATH)

Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES. NOT FOR MILITARY USE.

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Data Images: 81-103 Anti-NAI2 ΔSP (At) antibody, rabbit polyclonal

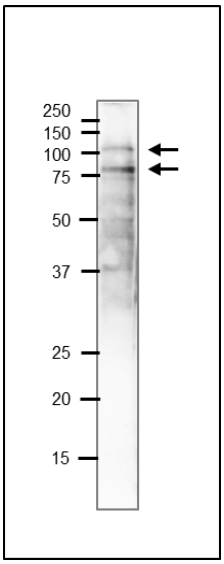

Fig.1 Western Blot of NAI2 in extract of arabidopsis.seedling

Crude extract of 7day old seedling of *Arabidopsis thaliana* was run on 12.5% SDS-PAGE and blotted overnight to PVDF membrane by wet system. Anti-MEB1 antibody was used at 1/4,000 dilution. Secondary antibody (goat anti-rabbit IgG antibody HRP-conjugated, ab97051) was used at 1/10,000 dilution.

Among the two bands the lower band was suggested to be degradation product of the upper band (Ref.1)

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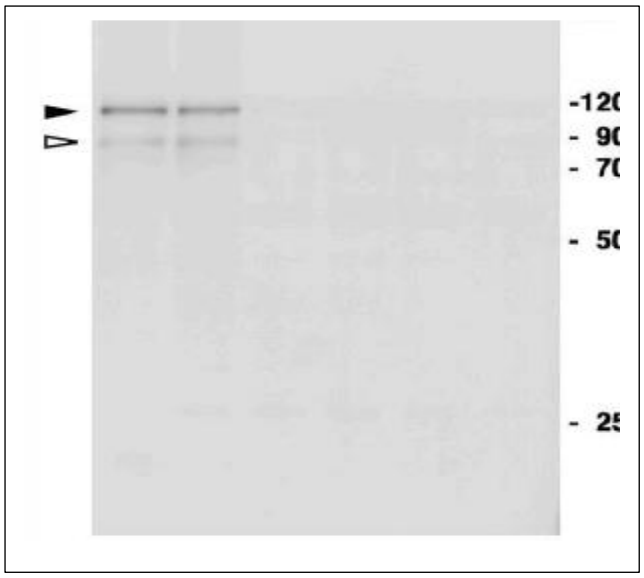

Fig.2　Validation of the anti-NAI2 (ΔSP)the antibody specificity by western blot using extracts of mutant seedling of Arabidopsis.

Total proteins were extracted from 20 plants of 7-d-old seedlings using 200 ml of 23 sample buffer (20 mM Tris-HCl buffer, pH 6.8, 40% glycerol, 2% SDS, and 2% 2-mercaptoethanol). The extracts (10 ml) were subjected to SDS-PAGE (12.5% acrylamide gel). The separated proteins were transferred to a nylon membrane and subjected to western blot analysis using anti-NAI2(ΔSP) antibody at 1/2,000 dilution.

As the second antibody, goat anti-rabbit IgG antibody HRP-conjugated was used at 1/10,000 dilution.

Samples. 1. Wild-typ. 2 Wild-type with GFP-h 3.*nai2-1.* 4. *nai2-2.* 5. *nai2-3*　6.*nai1-1*

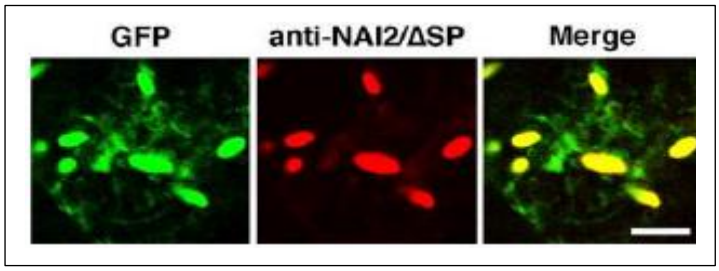

Fig.3 Immunofluorescence staining of NAI peotein in ER bodies.

Immunofluorescence analysis of NAI2 in 7-d-old GFP-h seedlings. Left panels, the ER-targeted GFP signals; middle panels, the NAI2 signals, which were detected by antibodies against NAI2/ΔSP, and Cy3-labeled second antibodies; right panels, the merged images. Bars = 10 mm.

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Reference: This antibodyhas been described I Ref.1 and used in the following publications.

1. Yamada K et al. NAI2 is an endoplasmic reticulum body component that enables ER body formation in Arabidopsis thaliana. Plant Cell. 2008 Sep;20(9):2529-40. PMID: 18780803 **WB, IF (Arabidopsis)**
2. Yamada K et al. Identification of two novel endoplasmic reticulum body-specific integral membrane proteins. Plant Physiol. 2013 Jan;161(1):108-20. PMID: 23166355 **WB (Arabidopsis)**
3. Ueda H et al. Endoplasmic Reticulum (ER) Membrane Proteins (LUNAPARKs) are Required for Proper Configuration of the Cortical ER Network in Plant Cells. Plant Cell Physiol. 2018 Oct 1;59(10):1931-1941. PMID: 30010972 **WB (Arabidopsis)**

Related Products

81-101 Anti-MEB1 (At) antibody, rabbit polyclonal

81-102 Anti-MEB2 (At) antibody, rabbit polyclonal

81-104 Anti-NAI2 C-terminal (At) antibody, rabbit polyclonal

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