

Product code	81-031
Size	200 μg
Storage	-20°C
Concentration	4.0 mg/ml
Buffer	PBS- with 50% glycerol
Purity	Purified IgG fraction with protein A from rabbit antiserum.
Immunogen	Purified recombinant Arabidopsis asparagine synthetase isoprotein 2 (ASN2),
	full-size, no-tag attached.
Isotype	Rabbit IgG
Reactivity	Plant ASN2 and ASN1 isoproteins
Special notes	Validation: The specificity of the antibody has been validated by western
	blotting with mutant plants.
Application	1. Western blotting (1/1,000-1/2,000 dilution)
	2. Immunohistochemistry, paraffin section (1/100-1/500)
	3. ELISA (assay dependent)
	Other applications have not been tested.
Background	Asparagine synthetase 2 (ASN2) is essential for nitrogen assimilation,
	distribution and remobilization within the plant via the phloem. ASN2 is
	expressed in leaf and ASN1 is expressed in floral organs.
	The amino acid sequences of Arabidopsis ASN1 and ASN2 are 76% identical.
	The amino acid sequences of Arabidopsis and Maize $\mathrm{ASN2}$ are 73.6% identical.
Data Link	UniProtKB <u>Q9LV77</u> (A. thaliana), <u>B5U8J7</u> (Z. mays)
Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC	
PROCEDURES. NOT FOR MILITARY USE.	

Anti-ASN (asparagine synthetase)(At) antibody, rabbit polyclonal

BioAcademia,Inc. Tel. 81-6-6877-2335 Fax. 81-6-6877-2336 info@bioacademia.co.jp https://www.bioacademia.co.jp/en/



Data Images: 81-031 Anti-ASN (asparagine synthetase)(At) antibody, rabbit polyclonal



Fig.1 Western Blot of ASN2 in arabidopsis leaf extract.

Anti-ASN2 antibody was used at 1/1,000 dilution. Secondary antibody (goat anti-rabbit IgG antibody HRP-conjugated, ab97051) was used at 1/10,000 dilution.

1. Arabidopsis leaf extract, 10 μg

Molecular mass of Arabidopsis ASN2 is 65 kDa.



Fig.2 Absence of ASN2 protein in leaf extracts of asn-2-1 and asn2-2 mutants.

AS is the position of ASN2 protein migrated at 65 kDa in SDS-PAGE and 70 kda is the position of pre-stained protein size marker. Col0 is wild-type Arabidopsis plant. *asn2-1* and *asn2-2 are* T1 insertion mutants.





Fig.3 Absence of ASN1 protein in extracts of floral organ of asn1 insertion mutant.

Wild type (Col-0) samples are analyzed in the upper window and *asn1* mutant samples are in the lower window. The upper numerals are the protein levels measured by densitometric tracing of western blot. Developmental stages are indicated below. Marker proteins are 72 and 55 kDa.



Fig.4 Immunofluorescence analysis of ASN2 in plant leaf section.

Arabidopsis thin leaf section was subjected to indirect immunofluorescence analysis using the Anti-ASN antibody as the primary antibody.

Goat anti-rabbit IgG labelled with Alexa 405 (Molecular Probes) was used as a secondary antibody.

Reference: This product has been used in the following publications.

- Gaufichon L, Masclaux-Daubresse C, Tcherkez G, Reisdorf-Cren M, Sakakibara Y, Hase T, Clément G, Avice JC, Grandjean O, Marmagne A, Boutet-Mercey S, Azzopardi M, Soulay F, Suzuki A. "Arabidopsis thaliana ASN2 encoding asparagine synthetase is involved in the control of nitrogen assimilation and export during vegetative growth." Plant Cell Environ. 2013 Feb;36(2):328-42. PMID: <u>22789031</u> WB, IHC ;arabidopsis
- 2. Gaufichon L, et al. ASN1-encoded asparagine synthetase in floral organs contributes to nitrogen filling in Arabidopsis seeds. <u>Plant J.</u> 2017 Aug;91(3):371-393. PMID:<u>28390103</u> WB, IHC; arabidopsis