

Anti-BGLU18 (At) antibody, rabbit polyclonal

Product code	81-105
Size	200 µg
Storage	-20°C
Concentration	2.0 mg/ml
Buffer	PBS- with 50% glycerol
Purity	Purified IgG fraction with protein A from rabbit antiserum.
Immunogen	Purified recombinant His6-Thioredoxin tagged BGLU18 protein (amino acids 27-528) of <i>A. thaliana</i> .
Isotype	Rabbit IgG
Reactivity	Arabidopsis thaliana. Not tested in other species.
Special notes	Validation: Specific reactivity has been validated by western blot using <i>bglu18</i> mutants.
Application	1. Western blotting (1/2,000-1/4,000) 2. Immunoelectron Microscopy (1/1,000)
Background	Hydrolyzes abscisic acid glucose ester (ABA-GE) which represents the predominant form of conjugated ABA (biologically inactive). No activity with beta-D-glucopyranosyl zeatin. The hydrolysis of ABA-GE in the endoplasmic reticulum (ER) forms free ABA and contributes to increase its cellular levels under dehydration conditions. ABA-GE hydrolyzing activity is enhanced by dehydration stress-induced polymerization into higher molecular weight forms. The ABA produced by BGLU18 contributes to the initiation of intracellular signaling as well as the increase in the extracellular ABA level. Length:528 amino acids. Predicted molecular mass:60,459 Subcellular location: Endoplasmic reticulum lumen. Modification: Elimination of 26-amino acid signal peptide from N-terminus.
Data Link	UniProtKB- Q9SE50 (BGL18_ARATH)
Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES. NOT FOR MILITARY USE.	

Data Images: 81-105 Anti-BGLU18 (At) antibody, rabbit polyclonal

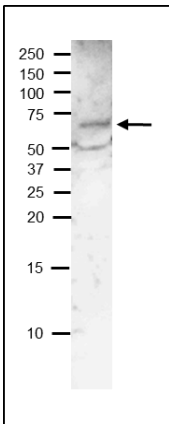


Fig.1 Western blot of BGLU18 in extract of arabidopsis seedling

Crude extract of 7day old seedling of *Arabidopsis thaliana* was run on 15-20% gradient SDS-PAGE and blotted overnight to PVDF membrane by wet system. Blocking was done with 3% skim milk. Anti-NAI2 antibody (C-terminal) was used at 1/1,000 dilution. Secondary antibody (goat anti-rabbit IgG antibody HRP-conjugated, ab97051) was used at 1/10,000 dilution.



Fig.2 Western blot analysis of BGLU18 protein accumulation.

BGLU18 accumulates in locally wounded cotyledons of both GFP plants (wild-type with GFP-fused with ER-retention signal) and *nai1* mutant but not in *bglu18* mutant.

Samples: Extracts of 12-day-old cotyledons from U: unwounded, L: locally wounded, S: systemically wounded. The anti-BGLU18 antibody was used at 1/2,000 dilution. As the second antibody, HRP-conjugated goat anti-rabbit IgG was used at 1/5,000 dilution.

Reference: This antibody has been described and used in the following publication.

- Ogasawara K et al. Constitutive and inducible ER bodies of *Arabidopsis thaliana* accumulate distinct beta-glucosidases. [Plant Cell Physiol.](#) 2009 Mar;50(3):480-8. PMID: [19147648](#) **WB, Immunoelectron Microscopy (Arabidopsis)**