Involvement of S-type anion channels in disease resistance against an oomycete pathogen in Arabidopsis seedlings

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ABSTRACT
Pharmacological indications suggest that anion channel-mediated plasma membrane (PM) anion efflux is crucial in early defense signaling to induce immune responses and programmed cell death in plants. Arabidopsis SLAC1, an S-type anion channel required for stomatal closure, is involved in cryptogein-induced PM Cl− efflux to positively modulate the activation of other ion fluxes, production of reactive oxygen species and a wide range of defense responses including hypersensitive cell death in tobacco BY-2 cells. We here analyzed disease resistance against several pathogens in multiple mutants of the SLAC/SLAH channels of Arabidopsis. Resistance against a biotrophic oomycete Hyaloperonospora arabidopsidis Noco2 was significantly enhanced in the SLAC1-overexpressing plants than in the wild-type, while that against a bacteria Pseudomonas syringae was not affected significantly. Possible regulatory roles of S-type anion channels in plant immunity and disease resistance against bacterial and oomycete pathogens is discussed.
related genes, *PRI* and *AtRbohD*, triggered by flg22, a microbe-associated molecular pattern (MAMP) from bacteria, was not affected significantly in comparison with the control [4]. These data were consistent with a previous study showing that the membrane potential change triggered by flg22 as well as elf18, another typical bacterial MAMP, was not affected by anion channel inhibitors including DIDS, or by a T-DNA insertional mutation in *SLAC1* or *SLAH3* gene in mesophyll cells [11,12]. These results suggest that the SLAC/SLAH family may not play a major role in MAMP-triggered immunity (PTI) in Arabidopsis. Effects of PM anion efflux or SLAC/SLAH channels in plant disease resistance against pathogens have not been so far studied.

In order to investigate the possible involvement of anion efflux mediated by SLAC/SLAH channels in disease resistance by Arabidopsis to bacterial and oomycete pathogens, *SLAC1*-overexpressor (*SLAC1-OE*), *slac1/slah3* double-mutant, and the wild-type seedlings were infected with a virulent strain of an obligate biotrophic oomycete *Hyaloperonospora arabidopsidis* (*Hpa Noco2*). As shown in Figure 1(a,b), the spread of conidiophore formation (indicative of oomycete reproduction) was significantly lower in the *SLAC1-OE* than in wild-type Col-0 seedlings (compatible line). Moreover, the frequency of the spread of conidiophores was also lower in the *SLAC1-OE* than in the wild-type (Figure 1(c)), indicating the enhanced resistance against the oomycete at the seedling stages. The conidiophore

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**Figure 1.** Depletion of *SLAC1/SLAH3* and overexpression of *SLAC1* result in increased disease resistance against the obligate biotrophic oomycete *Hyaloperonospora arabidopsidis* (*Hpa Noco2*). (a) Plant cell death and hyphal growth in representative Arabidopsis plants 7 days after infection (5.0 × 10⁴ spores ml⁻¹) as visualized by trypan-blue staining; the resistant Lever line were used as a control to display host cell death at infection sites. Scale bar: 100 μm. Data are representative of three experiments. Similar results were obtained in three independent experiments. (b) The number of conidiophores shown in (a) were quantified. Data are mean ± SE; n = 3 independent samples. * P < 0.05; significantly different from the control (Col-0). N.d.: not detected. (c) Categorization of conidiophore formation in 2-week-old Arabidopsis plants. Data are representative of three experiments. Similar results were obtained in three independent experiments.
formation was also slightly restricted in the slac1/slah3 mutant phenotype (Figure 1(b,c)). Microscopic examinations revealed that fungal hyphal growth monitored by trypan blue staining was observed in all plant lines, except for the resistant line La-er (Figure 1(a-c)).

Resistance against biotrophic microbial pathogens is assumed to be predominantly due to the salicylic acid (SA)-dependent mechanisms of resistance [13–15]. Since the SLAC1-OE plants were found to be more resistant to the biotrophic oomycete Hpa Noco2, we investigated whether SA-dependent defense gene activation was constitutively enhanced in the SLAC1-OE plants by analyzing the expression of the SA-inducible marker genes, PR1, PR2, and PR5. As shown in Figure 2, the transcript levels of these SA marker genes in uninfected seedlings were comparable among the wild-type, SLAC1-OE and slac1/slah3 mutant, suggesting that SA-dependent pathway is not affected by the expression levels of SLAC/SLAH channels at least in uninfected seedlings. Either defense signaling responses upon infection of the oomycete or regulation of SA-independent immune responses against an oomycete pathogen may be up-regulated by the overexpression of SLAC1.

The present results suggest importance of SLAC/SLAH-mediated anion fluxes in the defense signaling pathway against the oomycete in Arabidopsis seedlings. Resistant phenotypes of the SLAC1-OE plants are more critical than that of the slac1/slah3 mutant (Figure 1(b,c); Supplemental Figure 1), suggesting that SLAC/SLAH-mediated anion fluxes positively modulate plant defense responses against the oomycete Hpa Noco2 at least in Arabidopsis seedlings. This hypothesis is consistent with the enhanced defense responses by the ectopic-overexpression of Arabidopsis SLAC1 in tobacco BY-2 cells triggered by cryptogein [4]. The potential roles of anion fluxes mediated by SLAC/SLAH family in the regulation of disease resistance against Hpa Noco2 is an important topic for future research.

It currently remains unclear why the resistance trend against oomycetes was observed in Arabidopsis slac1/slah3 mutant. SLAC/SLAH family plays important roles in the control of nitrate loading of the root xylem [16]. Limitations in several nutrient sources may be responsible for reduced conidiophore formation in these plants. The limited availability of major plant carbon or nitrogen sources may explain changes in conidiophore formation in the slac1/slah3 mutant.

To investigate whether SLAC/SLAH channels also affect disease resistance against bacteria pathogens, we infiltrated slac1/slah3 mutant and SLAC1-OE plants

**Figure 2. Effects of expression levels of SLAC/SLAH family genes on SA-dependent defense gene activation at the basal level in Arabidopsis.** The expression of SA-dependent defense genes (PR1, PR2, and PR5) in Arabidopsis leaves. The amount of each mRNA was calculated from the threshold point located in the log-linear range of RT-PCR. Total RNA was isolated from Arabidopsis grown under normal growth conditions. Data are the mean ± SE of three independent experiments.
with a bacterial pathogen *Pseudomonas syringae*. The overexpression of *SLAC1* and a defect in *SLAC1/SLAH3* genes did not have a significant effect on the growth of the virulent hemibiotrophic bacterial strain *P. syringae* pv. *tomato* DC3000 (*Pst DC3000*) (Figure 3(a)), suggesting limited roles of the SLAC/SLAH channels in basal resistance against a bacteria pathogen. Since PTI plays critical roles in basal resistance [17], this result is consistent with the previous studies showing limited roles of the SLAC/SLAH channels in PTI [4,11]. Growth of *Pst-avrRpt2* bacteria (the avirulent strain) was also comparable between wild-type, *slac1/slah3* mutant, and *SLAC1-OE* plants (Figure 3(b)), suggesting their limited roles in the effector-triggered immunity against the bacterial pathogen.

Collectively, the present results suggest possible involvement of S-type anion channels in the resistance against the oomycete pathogen, while we did not find significant effects on the resistance against a bacterial pathogen, *Pst DC3000*. In Arabidopsis cultured cells, pharmacological indications suggest that R-type anion channels play key roles in cell death and ROS production triggered by the non-specific plant pathogen *Xanthomonas campestris* [18]. Other anion channel members including the R-type may be functionally redundant in some pathways. Roles of various anion channels may vary among cell types and the triggering signals/types of plant-pathogen interactions.

**Materials and methods**

**Plant materials and growth conditions**

*slac1*-*3* (SALK_099139) and *slah3*-*4* (SALK_111623) seeds were obtained from the Arabidopsis Biological Resource Center. *slac1*-*3/slah3*-*4* double mutant was obtained through standard crossing and genotyping in the F2 generation.

Surface-sterilized seeds of *Arabidopsis thaliana* (Col-0, La-er) and a *SLAC1* overexpression plant [4] as well as a *slac1*-*3/slah3*-*4* mutant line were germinated on MS medium containing 0.8% agar and grown in a growth chamber under long day conditions (16 h light/8 h darkness, 22°C).

**Real-time RT-PCR quantification**

Total RNA was isolated using NucleoSpin RNA Plus kit (TaKaRa, Shiga, Japan) according to the manufacturer’s protocol and quantified with a spectrophotometer. First-strand cDNA was synthesized from 500 ng total RNA using ReverTra Ace qPCR RT Kit (TOYOBO, Osaka, Japan).

**Figure 3.** Effects of expression levels of *SLAC/SLAH* family genes on disease resistance against the bacterial strain *Pseudomonas syringae* in *Arabidopsis*. Wild-type Col-0, slac1/slah3, and *SLAC1* overexpressing plants grown under long-day conditions for 3 weeks were injected into plant leaves with a syringe-inoculated with 1.0 × 10^5 cfu ml⁻¹ of virulent *Pseudomonas syringae* pv. *tomato* DC3000 (*Pst DC3000*) as well as the avirulent strain *Pst-avrRpt2*. After 0, 1, 2, and 3 days, bacterial growth was quantified by counting colony-forming units. Results represent means ± SDs (n ≥ 6), and the entire experiment was repeated at least three times with similar results. dpi; days post-inoculation.

Real-time PCR was performed using an ABI PRISM 7300 sequence detection system (Life
technologies) with THUNDERBIRD SYBR qPCR Mix (TOYOBO, Osaka, Japan) and gene specific primers. The following PCR primers were used: PR1-RealF, 5’-GTGGGTTAGCGAGAAGGCTA-3’; PR1-RealR, 5’-ACTTTGGCACATCCGAGTCT-3’; PR2-RealF, 5’-AGTCGATAGTCGATGCTAGTG-3’; PR2-RealR, 5’-GTACGTAGCTAAGCATGATC-3’; PR5-RealF, 5’-GTCAGCATGAAACAATGGCA-3’; PR5-RealR, 5’-GCTAGCATAGCTAAAGGCC-3’; AtTUB2-RealF, 5’-ATTCCCCGTCCTTTCACTTCT-3’; AtTUB2-RealR, 5’-CACATTCAGCATCTGCTGG-3’.

Relative mRNA abundances were calculated using the standard curve method and normalized to corresponding AtTUB2 gene levels. Standard samples of known template amounts were used to quantify PCR products.

**RT-PCR analysis**

Total RNA isolation and cDNA synthesis was performed as described above. PCR amplification was performed with an initial denaturation at 95°C for 3 min followed by the indicated number of cycles of incubations at 94°C for 30 s, 55°C for 90 s, and 72°C for 1 min, and a final extension at 72°C for 10 min using Arabidopsis SLAC1-specific primers (SLAC1(regular)-F, 5’-GCCATTAGCGTACCTCCCAT-3’; SLAC1(regular)-R, 5’-GCAGATATTCTTTCGCACTG-3’). AtACT2 gene (ACT2-F, 5’-GTAAGAGACATCAAGGAGAAGCTCTC-3’; ACT2-R, 5’-GGAGATCCACATCTGCTGGAAATG-3’) was used as an internal control. Aliquots of individual PCR products were resolved by agarose gel electrophoresis and visualized by ethidium bromide staining using a UV light.

**Infection assays**

For *Hyaloperonospora arabidopsidis* Noco2 (Hpa Noco2) assay, 2-week-old Arabidopsis plants were sprayed with a suspension of 5 × 10^4 spores ml⁻¹ and monitored by lactophenol-trypan blue staining as described previously [19]. To evaluate conidiospore production, more than 42 cotyledons of Arabidopsis in each line were harvested. After trypan-blue staining, total number of conidiophores were counted in each examination under bright-field microscopy with a haemocytometer. Statistical analyses have been performed from three independent experiments.

Infection assays with *Pst* DC3000 or *Pst* DC3000 (avrRpt2) were performed as described previously [20] on leaves of the same leaf stage of 3-week-old plants grown under long day condition. The virulent bacterial leaf pathogen *Pst* DC3000, which causes bacterial speck disease, was grown overnight at 28°C in NYGB liquid medium. The bacterial cells were collected by centrifugation and resuspended in 10 mM MgCl₂ to a final density of 1.0 × 10⁷ colony-forming units (cfu) ml⁻¹. For inoculation, the bacterial solution was injected into plant leaves with a syringe. The leaf discs that were excised from the infiltrated area after the indicated periods after inoculation were ground in 10 mM MgCl₂ and serially diluted to determine the bacterial numbers. Bacterial numbers were scored at 0, 1, 2 and 3 d post inoculation. For each sample, eight leaf discs were pooled and analyzed, and the experiment was repeated six times per data point.

**Statistical analyses**

Statistical Significance was determined using an unpaired Student *t*-test at *P* < 0.05.

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**Disclosure statement**

No potential conflict of interest was reported by the authors.

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