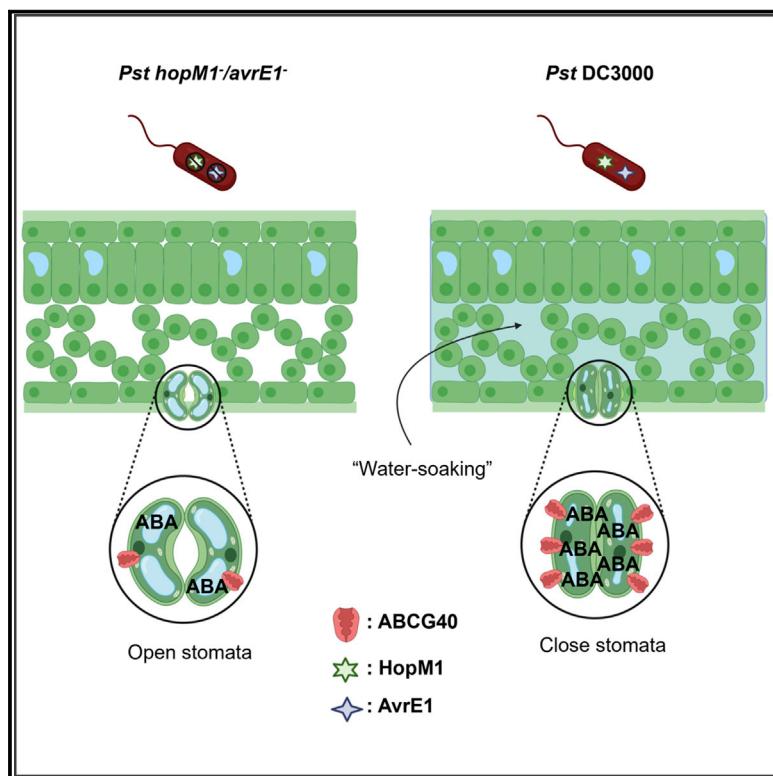


# Evolutionarily conserved bacterial effectors hijack abscisic acid signaling to induce an aqueous environment in the apoplast

## Graphical abstract



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## In brief

Roussin-Léveillée et al. demonstrate that two *Pseudomonas syringae* effector proteins induce abscisic acid (ABA) biosynthesis and signaling in *Arabidopsis*. Although stomatal opening early in infection allows access to the leaf apoplast, they show that ABA induces stomatal closure in later stages, leading to water-soaking lesions that benefit the bacteria.

## Highlights

- *Pseudomonas syringae* HopM1 and AvrE1 markedly affect the *Arabidopsis* transcriptome
- HopM1 and AvrE1 induce stomatal closure by inducing ABA biosynthesis and signaling
- Bacterial manipulation of ABA pathways is required to cause water-soaking lesions
- Induction of water soaking by HopM1 involves the *Arabidopsis* ABA transporter ABCG40



## Article

# Evolutionarily conserved bacterial effectors hijack abscisic acid signaling to induce an aqueous environment in the apoplast

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## SUMMARY

High atmospheric humidity levels profoundly impact host-pathogen interactions in plants by enabling the establishment of an aqueous living space that benefits pathogens. The effectors HopM1 and AvrE1 of the bacterial pathogen *Pseudomonas syringae* have been shown to induce an aqueous apoplast under such conditions. However, the mechanisms by which this happens remain unknown. Here, we show that HopM1 and AvrE1 work redundantly to establish an aqueous living space by inducing a major reprogramming of the *Arabidopsis thaliana* transcriptome landscape. These effectors induce a strong abscisic acid (ABA) signature, which promotes stomatal closure, resulting in reduced leaf transpiration and water-soaking lesions. Furthermore, these effectors preferentially increase ABA accumulation in guard cells, which control stomatal aperture. Notably, a guard-cell-specific ABA transporter, ABCG40, is necessary for HopM1 induction of water-soaking lesions. This study provides molecular insights into a chain of events of stomatal manipulation that create an ideal microenvironment to facilitate infection.

## INTRODUCTION

Many biotrophic pathogens secrete proteins, known as effectors, that manipulate their host to their benefit. These effectors are delivered, in many cases, to the cytoplasm of their host's cells via diverse mechanisms. Bacterial pathogens deliver such effectors via a needle-like structure, known as the type-3 secretion system (T3SS). In a compatible host-pathogen interaction, effector activity results in effector-triggered susceptibility (ETS) to infection. A conserved and essential strategy used by microbial invaders is to employ effectors to subvert the host immune system (Macho, 2016; Toruño et al., 2016). However, this aspect of ETS is not sufficient for optimal pathogen growth as full pathogenicity is not restored to effector-less pathogens in mutant plants that are near completely compromised in their immune system (Xin et al., 2016). This indicates that effectors are required to induce physiological microenvironments favorable to pathogen growth in addition to inhibiting plant defenses.

Recent reports indicate that an important aspect of ETS involves the establishment of an ideal microenvironment in the apoplast (Bezutyczny et al., 2018; Cohn et al., 2014; Cox et al., 2017; Xin et al., 2016). Elevated humidity levels in the environment are widely acknowledged as a key contributing factor to

disease development in the field. During infection, such conditions have been shown to allow microbial pathogens to induce an aqueous living space in the apoplast, referred to as "water-soaking lesions." In addition to suppressing host defenses (Hauck et al., 2003; Lozano-Durán et al., 2014; Xin et al., 2015), the widely conserved effectors HopM1 and AvrE1 were previously identified as the main inducers of water-soaking lesions in an *Arabidopsis-Pseudomonas* pathosystem (Xin et al., 2016). HopM1 has been shown to interact with the regulator of endocytic trafficking MIN7 (Nomura et al., 2006). It has also been reported that HopM1 triggers proteaphagy, whereas the selective autophagy receptor NBR1 antagonizes HopM1-induced water soaking (Üstün et al., 2018). A functional homolog of AvrE1 from *Pantoea stewartii*, WstE, interacts with the protein phosphatase 2A B' (PP2A) subunit in maize (Jin et al., 2016). However, the molecular and physiological basis for the ability of HopM1 and AvrE1 to induce water-soaking lesions remain elusive.

Stomata are formed by specialized cells that mediate gas exchange and water transpiration in plants. Many plant pathogens gain access to the leaf interior through these same structures. Upon sensing of microbe-associated molecular patterns (MAMPs), stomata close, thereby preventing pathogen entry (Melotto et al., 2006). This phenomenon, known as stomatal



immunity, is the first line of defense in MAMP-triggered immunity (MTI) and is often targeted by pathogens. *Pseudomonas syringae* pv. tomato DC3000 (*Pst* DC3000) produces a phytotoxin, known as coronatine (COR), that induces a reopening of closed stomata, allowing the pathogen access to the host apoplastic environment (Melotto et al., 2006). Although stomatal behavior during the early stages of infection has been the subject of numerous studies, their role in late stages is still unclear.

In this study, we have investigated the molecular and physiological basis for HopM1/AvrE1 induction of an aqueous microenvironment in the apoplast. Transcriptional profiling of *Arabidopsis thaliana* during an infection revealed that HopM1 and AvrE1 are functionally redundant in triggering abscisic acid (ABA) biosynthesis and signaling pathways. We show that ABA accumulation induced by HopM1 and AvrE1 triggers stomatal closure after an initial opening by COR and that manipulation of stomata has a dramatic effect on pathogenesis. Our study identifies a molecular and physiological explanation behind humidity-driven pathogenesis through the manipulation of ABA signaling.

## RESULTS

### Water-soaking effectors induce a major reprogramming of the *Arabidopsis* transcriptome and induce ABA pathways

To characterize the role of water-soaking effectors, we profiled the transcriptome of *A. thaliana* inoculated with *Pst* using RNA sequencing (RNA-seq). We included *hopM1*<sup>-</sup> and *avrE1*<sup>-</sup> single and double mutants of *Pst*, as these two effectors have been reported to be required for the induction of water-soaked lesions (Xin et al., 2016). Leaves were syringe infiltrated with a low dose of inoculum ( $1 \times 10^6$  CFU/mL) and tissues harvested during the *Pst* DC3000 exponential growth phase at 36 h post inoculation (hpi), as determined by bacterial growth curve dynamics (Figures S1A and S1B). This time point was selected to provide insights into the transcriptional profile of *Arabidopsis* during the active growth phase of the pathogen, when we expect effector proteins to be highly active. Transcriptional responses induced by *Pst* were highly influenced by the presence of HopM1 and AvrE1 (Figures 1A–1C). A principal component analysis of the RNA-seq data revealed a close transcriptional relationship between *Pst* DC3000-, *hopM1*<sup>-</sup>-, and *avrE1*<sup>-</sup>-infected plants, but a more distant relationship to the *hopM1*<sup>-</sup>/*avrE1*<sup>-</sup> (*h*<sup>-</sup>/*a*<sup>-</sup>)-infected plants (Figure 1B). The number of differentially expressed genes (DEGs) in *Arabidopsis* Col-0 plants infected by *Pst* DC3000 was strikingly higher than in their *h*<sup>-</sup>/*a*<sup>-</sup> infected counterparts (9,623 versus 4,906, respectively) (Figures 1C and S1C–S1G). We also compared the transcriptome of plants infected with *Pst* *h*<sup>-</sup>/*a*<sup>-</sup> with a previously described dataset from plants treated with the MTI-inducing peptide flg22 (Winkelmüller et al., 2021) to compare HopM1/AvrE1 transcriptional reprogramming in contrast to the response of an *Arabidopsis* immune response. Although these datasets were generated from different experimental conditions, we nonetheless find that they result in distinct transcriptional signatures. That is, less than half of the genes upregulated by the flg22 response were also induced by *Pst* *h*<sup>-</sup>/*a*<sup>-</sup> and there was only a very minor overlap in downregulated genes (Figure S1G). This suggests that *Pst* effectors other than HopM1 and AvrE1 contribute to countering

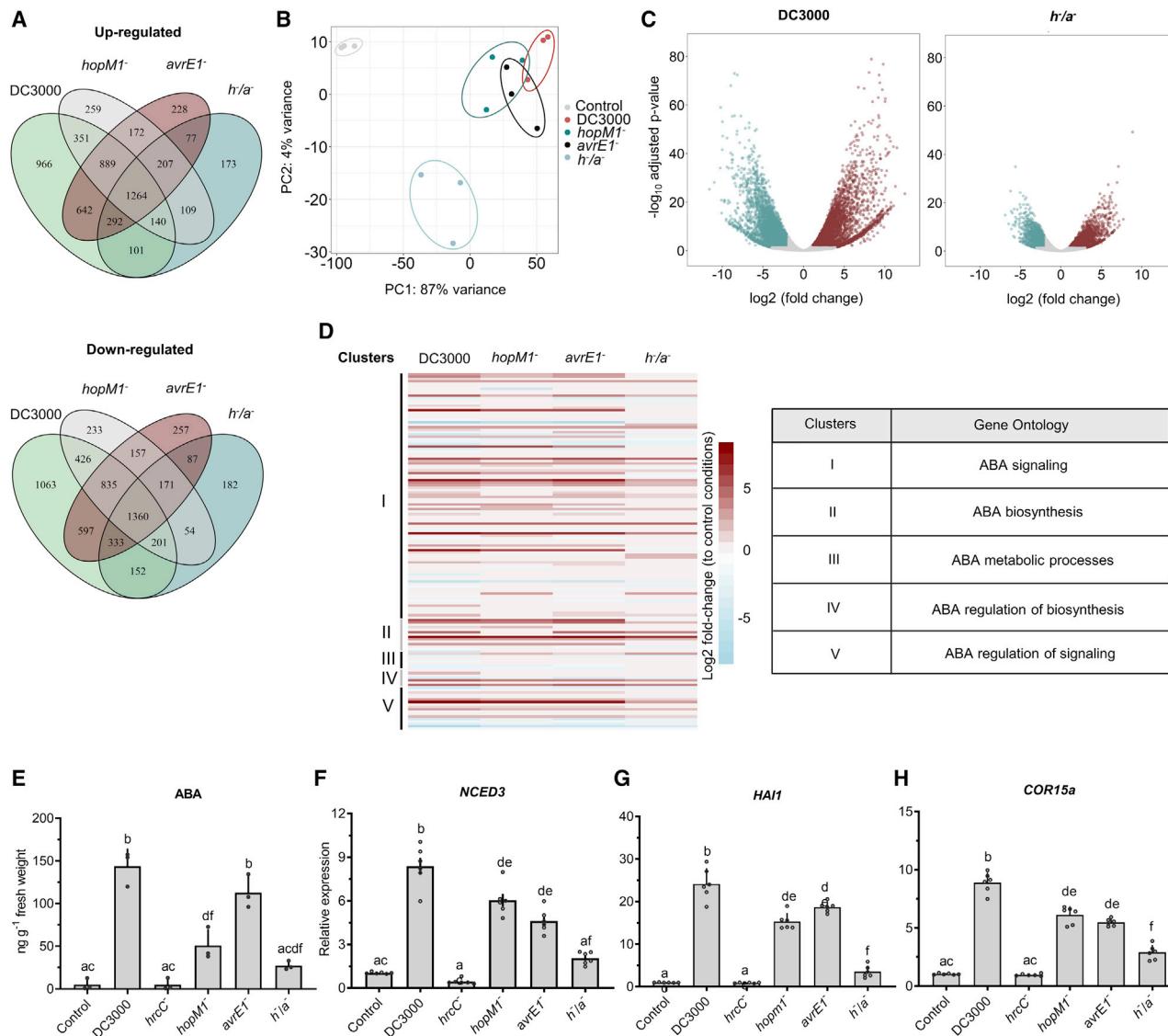
the transcriptional reprogramming associated with an MTI response.

Gene ontology (GO) analysis associated with the transcriptional response of *Arabidopsis* to infection by *Pst* DC3000, *hopM1*<sup>-</sup>, *avrE1*<sup>-</sup>, and *h*<sup>-</sup>/*a*<sup>-</sup> indicated that HopM1 and AvrE1 induce a gene expression pattern related to an ABA response (Figure 1D; Table S1). ABA signaling is a well-known response to water stress and can impact multiple regulatory pathways. Thus, we investigated changes in ABA responses mediated by HopM1 and AvrE1. Using ultraperformance liquid chromatography-mass spectrometry (UPLC-MS), we observed an increase in ABA levels in plants infected with *Pst* DC3000, compared with uninfected plants, but not with *h*<sup>-</sup>/*a*<sup>-</sup> (Figure 1E). In concordance with this, *A. thaliana* plants infected with *Pst* DC3000, *hopM1*<sup>-</sup>, and *avrE1*<sup>-</sup> single mutants, but not with *h*<sup>-</sup>/*a*<sup>-</sup> double mutant, showed significantly higher expression levels of the ABA biosynthesis marker gene *NCED3* than mock-infected plants (Figure 1F). Furthermore, inoculation with *Pst* DC3000 resulted in an increase in expression of known components of the ABA signaling pathway, *HAI1* (negative regulator of ABA signaling; ~24-fold increase) and *COR15a* (cold- and ABA-regulated gene; ~9-fold increase) (Baker et al., 1994; Yoshida et al., 2006). In contrast, the *h*<sup>-</sup>/*a*<sup>-</sup> double mutant showed only a moderate increase (~2.5-fold for both *HAI1* and *COR15a*) compared with control plants, while the *hopM1*<sup>-</sup> and *avrE1*<sup>-</sup> single mutants showed an intermediate level of expression (at least half the level of induction induced by DC3000) (Figures 1G and 1H). Importantly, this ABA signature was observed at 24 hpi, at a time point when bacterial densities do not significantly differ between wild-type (WT) and mutant strains lacking HopM1 and/or AvrE1 (Figure 2F). Together, these results indicate a role for water-soaking effectors in increasing ABA accumulation and subsequent responses.

### HopM1 and AvrE1 induce stomatal closure following leaf entry

Under high levels of humidity, *Pst* DC3000 induces clear water-soaking lesions, which are crucial for its virulence (Xin et al., 2016). Since stomata mediate gas exchange, including loss of water vapor, between the leaf apoplast and the extracellular environment, we hypothesized that *Pst* DC3000 might induce stomatal closure following leaf entry to maintain high internal levels of humidity. As stomata respond rapidly to changes in relative humidity settings, we optimized a stomatal fixation method to prevent stomatal movement during analyses (Eisele et al., 2016). Briefly, we found that treatment of leaves with a solution containing 4% formaldehyde allowed for the processing of large numbers of leaves in a context where stomatal aperture does not change upon transfer of leaves from experimental to analytical conditions (Figures S2A–S2E).

We investigated *A. thaliana* stomatal responses to *Pst* DC3000 under elevated relative humidity over the 3-day course of an infection. Plants dip-inoculated with *Pst* DC3000 displayed the well-documented phenomenon of stomatal closure within 1 h of inoculation, followed by reopening of stomata after 4 h. Interestingly, stomata were closed at the 24 hpi time point in *Pst* DC3000-inoculated plants, suggesting a dynamic manipulation of stomatal movement (Figure 2A). In agreement with this, leaves infected with *Pst* DC3000 underwent slower water loss than

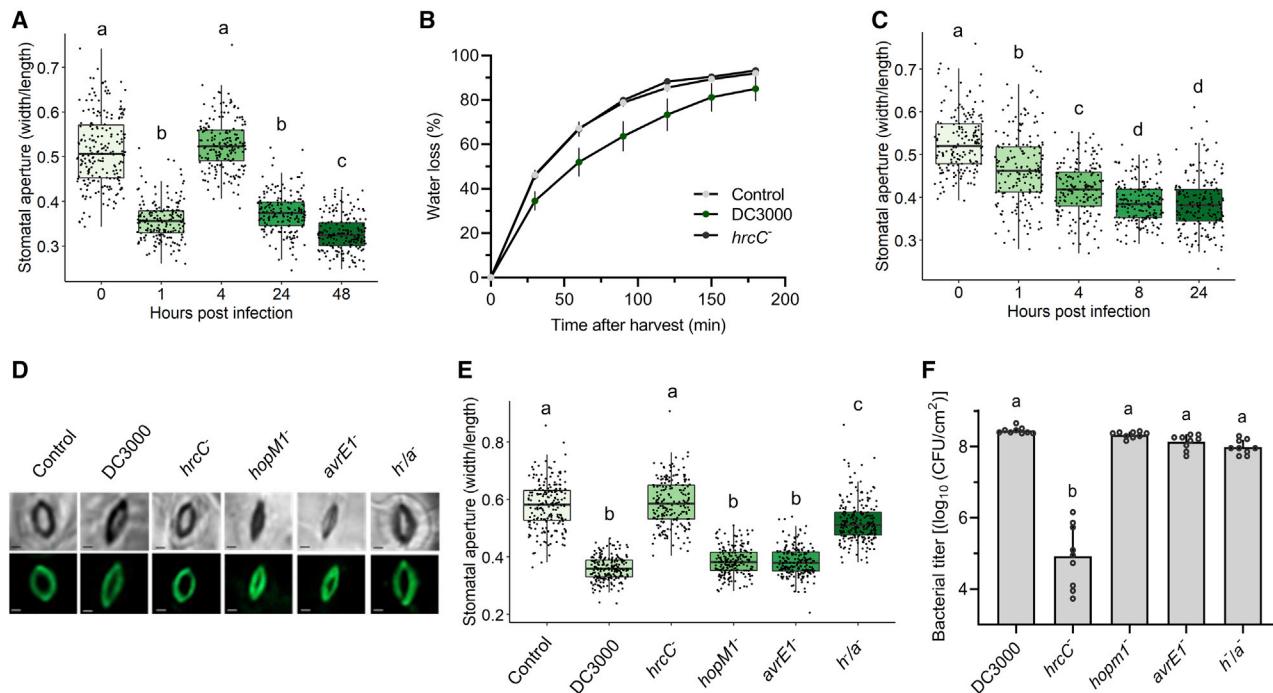


**Figure 1.** Water-soaking effectors induce a major transcriptional reprogramming and increase ABA biosynthesis

- (A) Venn diagrams displaying specific and shared differentially expressed genes (DEGs) in *A. thaliana* Col-0 challenged with *Pst* DC3000, *hopM1<sup>-</sup>*, *avrE1<sup>-</sup>*, and *hopM1<sup>-</sup>/avrE1<sup>-</sup>* (*h/a<sup>-</sup>*) ( $1 \times 10^6$  CFU/mL) at 36 h post inoculation (hpi).
- (B) Principal component (PC) analysis showing expression changes of transcriptomic data from the conditions described in (A), as indicated.
- (C) Volcano plot representing DEGs in *Arabidopsis* plants infected with *Pst* DC3000 or *h/a<sup>-</sup>* ( $1 \times 10^6$  CFU/mL at 36 hpi). Blue, red, and gray dots represent significantly downregulated, upregulated, and not significantly altered DEGs, respectively.
- (D) Heatmap representing expression differences of genes with gene ontologies involved in ABA-related processes, as indicated, upon infection of *Arabidopsis* plants with the indicated *Pst* strains.
- (E) ABA quantification in *Arabidopsis* Col-0 leaves infected with  $0.5 \times 10^8$  CFU/mL of the indicated *Pst* strains by UPLC-MS at 24 hpi.
- (F) Relative expression of the ABA biosynthesis marker gene *NCED3* in *Arabidopsis* Col-0 plants infiltrated with the indicated *Pst* strains ( $1 \times 10^8$  CFU/mL) at 24 hpi measured by real-time quantitative PCR (RT-qPCR).
- (G and H) Relative expression of the ABA signaling pathway marker genes *HAI1* (G) and *COR15a* (H) in *Arabidopsis* Col-0 plants inoculated as in (E). Data are represented as mean of total experimental replicates  $\pm$  SEM. Different letters indicate statistically significant differences (adjusted  $p < 0.01$ , one-way ANOVA).

leaves in control plants (Figure 2B). These phenomena required the bacterial T3SS as a *Pst* mutant lacking this system (*Pst* *hrcC<sup>-</sup>*) did not affect stomatal aperture or water loss (Figures 2B, 2D, and 2E). Stomatal closure was also observed in *Arabidopsis* plants that were syringe infiltrated with *Pst* DC3000, although no reopening was observed at 4 hpi (Figure 2C).

Given the importance of HopM1 and AvrE1 in inducing ABA responses, we tested the importance of these effectors in inducing stomatal closure. Interestingly, *Arabidopsis* plants infected with *Pst* *hopM1<sup>-</sup>* and *avrE1<sup>-</sup>* single mutants displayed closed stomata at 24 hpi, whereas those infected with the *h/a<sup>-</sup>* double mutant show stomatal aperture values similar to uninfected or



**Figure 2. *Pseudomonas syringae* water-soaking effectors induce stomatal closure in the late stages of infection**

(A) *Arabidopsis* Col-0 plants were dip-inoculated in a *Pst* DC3000 bacterial suspension ( $2 \times 10^8$  CFU/mL) containing 0.025% Silwet L-77. Epifluorescence microscopy images were used to measure stomatal apertures ( $n > 150$ ) at the indicated hpi.

(B) Water loss in *Arabidopsis* Col-0 plants inoculated as in A with mock (control), WT (DC300), or T3SS defective (*hrcC*<sup>-</sup>) strains at 24 hpi. Leaves were detached from plants and stomata were fixed in stomatal fixation solution at the beginning of water loss measurement.

(C) Stomatal aperture measurements of *Arabidopsis* Col-0 plants syringe infiltrated with *Pst* DC3000 ( $1 \times 10^8$  CFU/mL) measured over time ( $n > 150$  stomata).

(D) Microscopy images of abaxial sides of 4-week-old *A. thaliana* Col-0 leaves infiltrated with MgCl<sub>2</sub> 10 mM (control), *Pst* DC3000, *hrcC*<sup>-</sup>, *hopM1*<sup>-</sup>, *avrE1*<sup>-</sup> and *h*<sup>-</sup>/*a*<sup>-</sup> ( $1 \times 10^8$  CFU/mL), as indicated, at 24 hpi. Upper panel: bright field, lower panel: GFP field to visualize rhodamine 6G fluorescence. Bar scales represent 2  $\mu$ m.

(E) Stomatal aperture measurements of *Arabidopsis* Col-0 plants inoculated as in (D) at 24 hpi.

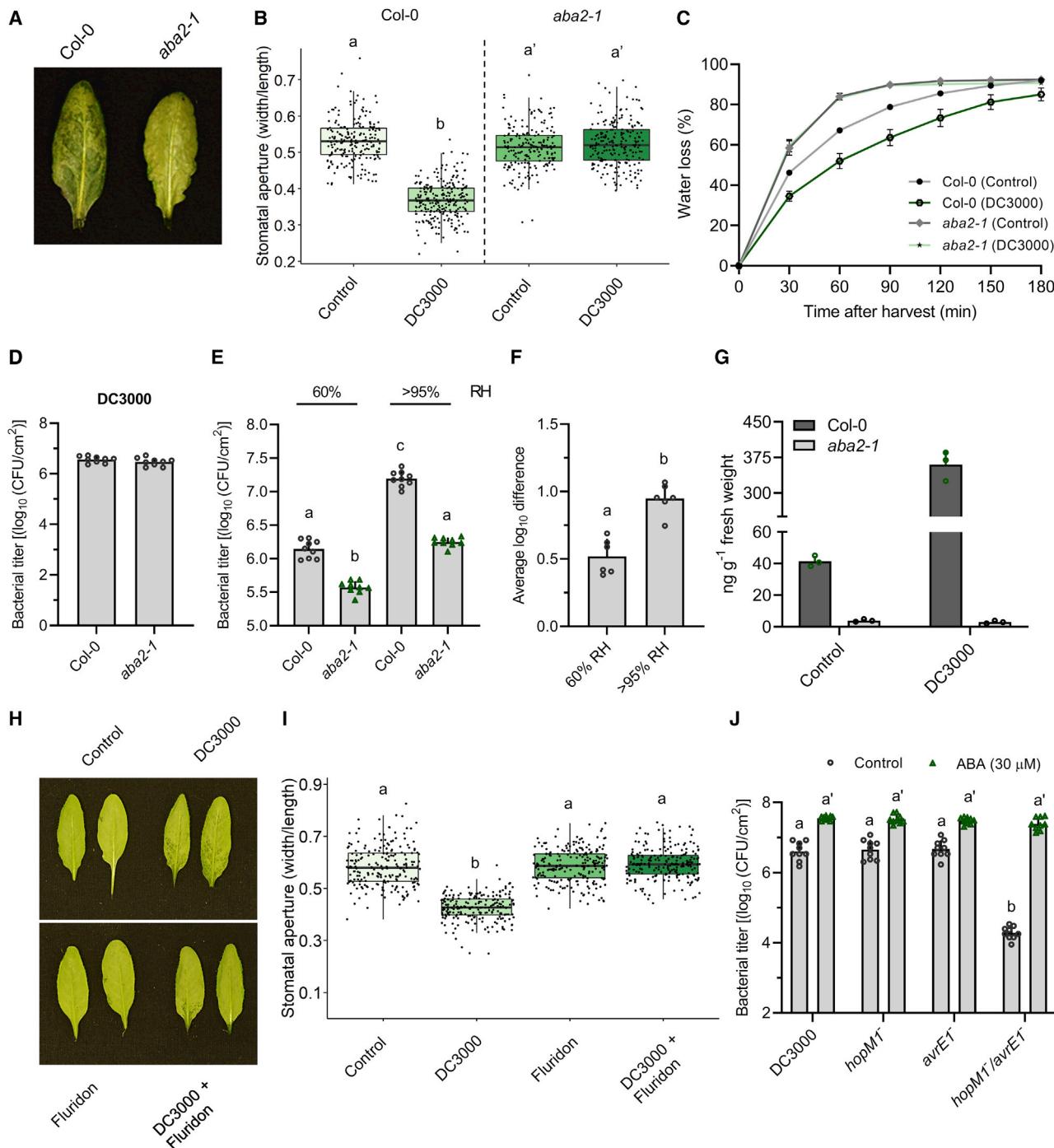
(F) Bacterial counts found in plants analyzed in (D) and (E). Data are represented as median values, with minimum and maximum values indicated (A, C, E) or as means of total experimental replicates,  $\pm$  SEM (B, F). Different letters indicate statistically significant differences (adjusted  $p < 0.01$ , one-way ANOVA).

*hrcC*<sup>-</sup> infected plants (Figures 2D and 2B). The differences in aperture opening cannot be due to differences in bacterial density as bacterial levels in the apoplast are not significantly different at this time point (Figure 2F). To further validate the role of these two effectors in inducing stomatal closure, we used transgenic *Arabidopsis* lines expressing HopM1 and AvrE1 under the control of a dexamethasone (DEX)-inducible promoter. Upon DEX treatment, HopM1- and AvrE1-expressing lines displayed significant stomatal closure compared with WT and non-DEX-treated plants, as well as induction of *NCED3* expression (Figures S3A and S3B). These results suggest that *Pst* has evolved a strategy to initially open stomata to facilitate entry into the apoplast, after which it induces stomatal closure to maintain high levels of humidity in its environment.

#### ABA-mediated stomatal closure as the primary mechanism behind water-soaking lesions

Control of gas exchange by stomata is primarily orchestrated by the phytohormone ABA (Kollist et al., 2019). Interestingly, ABA pathway manipulation by microbial invaders is also known to contribute to susceptibility in many plant pathosystems (Fan et al., 2009). Since ABA is known to be exploited by *Pst*

DC3000 to increase its virulence (de Torres-Zabala et al., 2007), we hypothesized that this enhanced susceptibility to infection might be caused by the pathogen's ability to induce water-soaking lesions in an ABA-dependent manner. To test this, we inoculated WT *Arabidopsis* and an *aba2-1* mutant, which is known to produce considerably less ABA (Rook et al., 2001). The induction of water-soaked lesions by *Pst* was completely abolished in the *aba2-1* mutant at 24 hpi (Figure 3A). Likewise, *Pst* inoculation did not result in stomatal closure or in the production of ABA in the *aba2-1* mutant (Figures 3B and 3G), and leaves from *aba2-1* plants showed the same degree of water loss in the presence or absence of bacteria (Figure 3C). Importantly, the levels of bacteria were the same at the time points tested, ruling out bacterial density as a factor in these observations (Figure 3D). Furthermore, it is reported that bacterial levels of *Pst* DC3000 in ABA biosynthesis or signaling mutants are similar to the levels of *Pst* *h*<sup>-</sup>/*a*<sup>-</sup> in WT *Arabidopsis* (Hu et al., 2022). To strengthen the link between ABA and water-soaking lesions, we co-infiltrated *Pst* with fluridone, which has been shown to interfere with carotenoid and ABA biosynthesis (Gamble and Mullet, 1986). We found that co-infiltrating the pathogen with fluridone prevented *Pst* from causing water-soaking lesions through stomatal



**Figure 3. ABA biosynthesis is required for humidity-driven pathogenesis in *Arabidopsis***

(A) Leaves of *Col-0* *Arabidopsis* and the *aba2-1* mutant were infiltrated with *Pst* DC3000 ( $1 \times 10^8$  CFU/mL) and plants grown at 95% humidity. Photos taken at 24 hpi.

(B) Stomatal aperture measurements of *Col-0* and the *aba2-1* mutant inoculated with *Pst* DC3000 under the same conditions as in (A) ( $n > 150$  stomata).

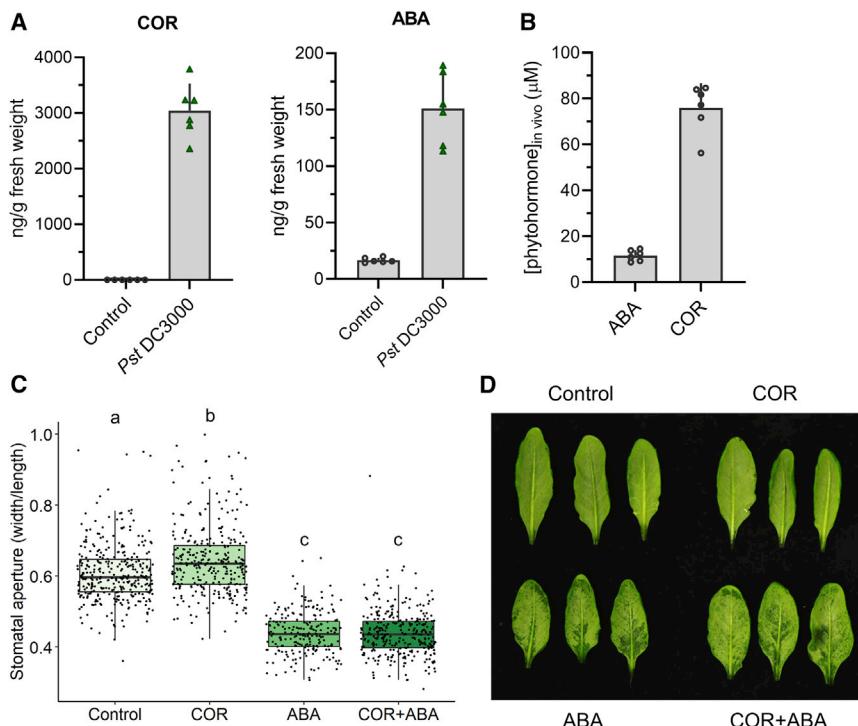
(C) Water loss was measured in leaves of *Arabidopsis* *Col-0* and *aba2-1* mutant plants inoculated with 10-mM MgCl $_2$  (control) or *Pst* DC3000 ( $1 \times 10^8$  CFU/mL). Leaves were detached at 24 hpi and stomata fixed in stomatal fixation solution at the beginning of water loss measurements.

(D) Bacterial titers measured in plants analyzed in (A) at 24 hpi.

(E) *Col-0* and *aba2-1* mutant plants were infected with *Pst* DC3000 ( $1 \times 10^5$  CFU/mL) and kept under either 60% or >95% RH levels. Bacterial titers were quantified 3 days post-infection.

(F) Average log difference in bacterial titers between WT and *aba2-1* plants assayed under the conditions as in (E).

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**Figure 4.** COR levels found *in vivo* do not prevent ABA from inducing stomatal closure and water-soaking lesions

(A) COR and ABA levels found *in planta* during a *Pst* DC3000 infection. COR and ABA levels were quantified by UPLC-MS in *Arabidopsis* Col-0 leaves mock inoculated (control) or infiltrated with *Pst* DC3000 ( $5 \times 10^7$  CFU/mL) at 24 hpi.

(B) Average calculated concentration of ABA and COR found in *Arabidopsis* leaves infiltrated with *Pst* DC3000 ( $5 \times 10^7$  CFU/mL) at 24 hpi ( $n = 6$  plants from three independent experimental replicates).

(C) Stomatal apertures of *Arabidopsis* Col-0 leaves infiltrated with ABA (10  $\mu$ M) and/or COR (80  $\mu$ M), as measured from epifluorescence images from stomata-fixed leaves at 8 h post-treatment.

(D) Appearance of water-soaking lesions in leaves described in (C). Data are represented as mean of total experimental replicates  $\pm$  SEM (A, B), or as median values, with minimum and maximum values indicated (C). Different letters indicate statistically significant differences,  $p < 0.05$ , one-way ANOVA.

closure, supporting a role for ABA in inducing stomatal closure during an infection (Figures 3H and 3I).

To better understand the role of ABA in humidity-driven pathogenesis, we infected WT and *aba2-1* mutant plants with *Pst* and maintained two different humidity levels over the course of the infection. Differences in bacterial levels between Col-0- and *aba2-1*-infected plants were greater under elevated humidity conditions as compared with ambient humidity levels, suggesting that ABA contributes to humidity-driven pathogenesis (Figures 3E and 3F). Next, we questioned whether treating plants with ABA would affect the growth rate of the *h<sup>-</sup>/a<sup>-</sup>* mutant. A significant difference in bacterial growth at three dpi was observed between the *h<sup>-</sup>/a<sup>-</sup>* double mutant in *Arabidopsis* as compared with DC3000, as well as *hopM1<sup>-</sup>* and *avrE1<sup>-</sup>* single mutants, as expected (Figure 3J). However, treatment of plants with ABA resulted in a complete rescue of the double mutant growth rate to the same levels as *Pst* DC3000 (Figure 3J). We suggest that ABA's ability to rescue the *h<sup>-</sup>/a<sup>-</sup>* mutant growth deficiency is due to its ability to induce stomatal closure, which in turn results in water-soaking lesions (Figures S4A–S4D) that are beneficial to the bacteria. As such, ABA appears to be both necessary and sufficient for the induction of stomatal closure.

#### Coronatine does not prevent ABA-mediated stomatal closure

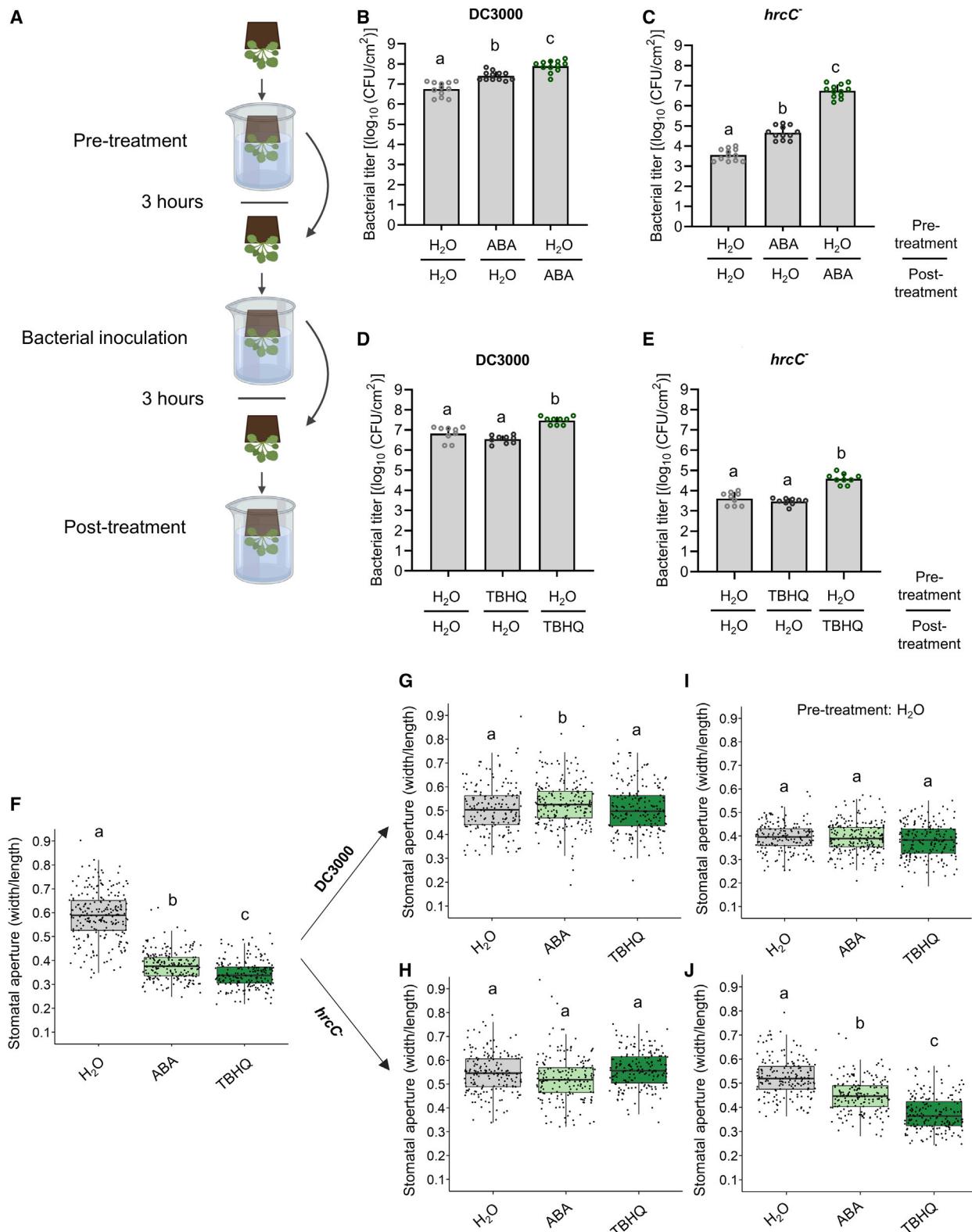
*Pst* produces COR in the early stages of an infection to reopen closed stomata to gain access to the apoplastic environment (Melotto et al., 2006). We questioned whether the amount of COR produced by *Pst* *in planta* was sufficient to antagonize ABA-induced stomatal closure once the pathogen has entered the apoplast. We measured *in planta* levels of COR and ABA, as well as several other phytohormones, in *Arabidopsis* plants infected by *Pst* DC3000 over a time course of 24 h under two different humidity settings (Figures S5A–S5F). Approximate concentrations of these molecules *in planta* were calculated based on the amount of water found in *Arabidopsis* leaves, as determined by measuring fresh and dry weight in leaves of plants grown at 95% humidity (Figures 4A and 4B). ABA and COR were then infiltrated into *Arabidopsis* Col-0 leaves at concentrations approximating those found *in vivo* during a *Pst*-*Arabidopsis* interaction. As expected, we found that at concentrations found in *Pst*-infected leaves, COR-induced stomatal opening and ABA was sufficient to induce stomatal closure and water soaking in the absence of bacteria. However, at physiologically relevant concentrations, COR did not prevent ABA from inducing stomatal closure or from inducing water-soaking lesions (Figures 4C–4D).

(G) Quantification of ABA in *Arabidopsis* Col-0 and *aba2-1* mutant plants mock inoculated (control) or inoculated with *Pst* DC3000 ( $1 \times 10^8$  CFU/mL), as measured by UPLC-MS at 24 hpi.

(H) Water-soaking lesion phenotypes in *Arabidopsis* leaves syringe-inoculated with MgCl<sub>2</sub> 10 mM (control), *Pst* DC3000 ( $1 \times 10^8$  CFU/mL), fluridone (100  $\mu$ M), or both *Pst* DC3000 and fluridone.

(I) Stomatal aperture measurements from leaves described in (H).

(J) ABA-mediated rescue of the *hopM1<sup>-</sup>/avrE1<sup>-</sup>* double mutant. *Arabidopsis* Col-0 plants were infected with the indicated *Pst* mutants ( $1 \times 10^5$  CFU/mL) followed by treatment, at 2, 24, and 48 hpi, with either ABA (30  $\mu$ M) or control solution (0.5% ethanol). Plants were treated at 2, 24, and 48 hpi. Treatments were carried out by spraying each solution on the abaxial and adaxial part of each plant. Bacterial titers were evaluated at 3 dpi. Data are represented as median values, with minimum and maximum values indicated (B, I) or as mean of total experimental replicates,  $\pm$  SEM (C–G and J). Different letters indicate statistically significant differences,  $p < 0.0005$ , two-tailed Student's t test (B and F) or ANOVA (E, I, and J).



**Figure 5. A role for late stomatal closure in bacterial pathogenesis**

(A) Schematic diagram depicting the experimental design to evaluate the importance of pre- versus post-infection stomatal closure. Created with BioRender.  
(B–E) All plants were subjected to three dip treatments. In the pre-treatment, plants were first dipped in a solution containing either 0.05% ethanol plus 0.025% Silwet L-77 in water (H<sub>2</sub>O) or the same solution also containing either ABA (30  $\mu$ M) or TBHQ (100  $\mu$ M). 3 h after pre-treatment, plants were dip-inoculated with a

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and 4D). Furthermore, a *Pst* mutant deficient in the biosynthesis of COR, DB29 (Brooks et al., 2004), displayed no difference in terms of water-soaking symptoms or in its ability to induce stomatal closure, compared with WT (Figures S5G–S5I). Thus, although COR and ABA have been shown to have opposite effects on stomatal behavior in early stages of infection, the concentration levels found in the later stage of an infection result in stomatal closure.

### Importance of late stomatal closure in bacterial pathogenesis

The non-virulent *Pst* strain *hrcC*<sup>−</sup> is unable to grow well in the apoplastic environment of *Arabidopsis*. We therefore questioned whether forcing stomatal closure could re-establish virulence to *Pst hrcC*<sup>−</sup>. To test this, we established a protocol whereby *Arabidopsis* plants were subjected to a series of treatments to affect stomatal movement either before or after inoculation with *Pst DC3000* or *Pst hrcC*<sup>−</sup> (Figure 5A). Treatments included H<sub>2</sub>O alone, ABA or *tert*-butylhydroquinone (TBHQ). TBHQ is an aromatic compound derived from hydroquinone and has been shown to induce stomatal closure in an ABA-independent manner (Toh et al., 2018). We find that, similar to ABA, TBHQ treatment induces water-soaking lesions at high humidity, as well as stomatal closure, suggesting that the latter is sufficient for the former (Figures S4E and S4G).

We found that pre-treatment of *Arabidopsis* plants with ABA 3 h prior to bacterial dip-inoculation resulted in enhanced growth of both *Pst DC3000* and *Pst hrcC*<sup>−</sup>, compared with plants pre-treated with H<sub>2</sub>O (Figures 5B and 5C). However, when *Arabidopsis* plants were instead treated with ABA 3 h after bacterial inoculation, this enhanced growth was more dramatic, with *Pst hrcC*<sup>−</sup> growing at levels nearing those of *Pst DC3000* in untreated plants (Figures 5B and 5C).

Since ABA has been reported to dampen immune signaling (de Torres Zabala et al., 2009; Mine et al., 2017), we sought to test the effect of inducing stomatal closure without affecting immunity. *Arabidopsis* plants treated with TBHQ prior to pathogen challenge did not show any difference in proliferation of *Pst DC3000* or *Pst hrcC*<sup>−</sup> 3 days post inoculation (Figures 5D and 5E). However, plants that were treated with TBHQ 3 h post pathogen inoculation displayed enhanced growth of both *Pst DC3000* and *Pst hrcC*<sup>−</sup> (Figures 5D and 5E). As TBHQ did not alter the protection provided by a flg22 pre-treatment (Figure S4F), we suggest that TBHQ does not affect immune signaling, but rather, augments bacterial growth by inducing stomatal closure and causing an aqueous apoplastic environment.

The interpretation of the above results depends on the actual behavior of stomates in this set of experiments. As such, we measured the stomatal apertures of plants treated as above. As

expected, pre-treatment with ABA or TBHQ resulted in stomatal closure (Figure 5F). Interestingly, both *Pst DC3000* and *Pst hrcC*<sup>−</sup> induced a reopening of stomata 3 h after dip-inoculation, regardless of ABA or TBHQ pre-treatment (Figures 5G and 5H). This suggests that the stomatal reopening capacity of the pathogen overcomes the action of ABA and TBHQ in this situation. In plants pre-treated with H<sub>2</sub>O alone, post-treatments with H<sub>2</sub>O, ABA, or TBHQ 3 h after DC3000 inoculation all resulted in similar stomatal closure activity (Figure 5I). This is presumably because the action of DC3000 effectors is sufficient to induce stomatal closure. In contrast, post-treatment of plants inoculated with *Pst hrcC*<sup>−</sup> with H<sub>2</sub>O alone did not result in stomatal closure, whereas ABA and TBHQ treatments did (Figure 5J). Thus, stomatal closure following leaf entry appears to be an important element in bacterial pathogenesis in addition to immune suppression.

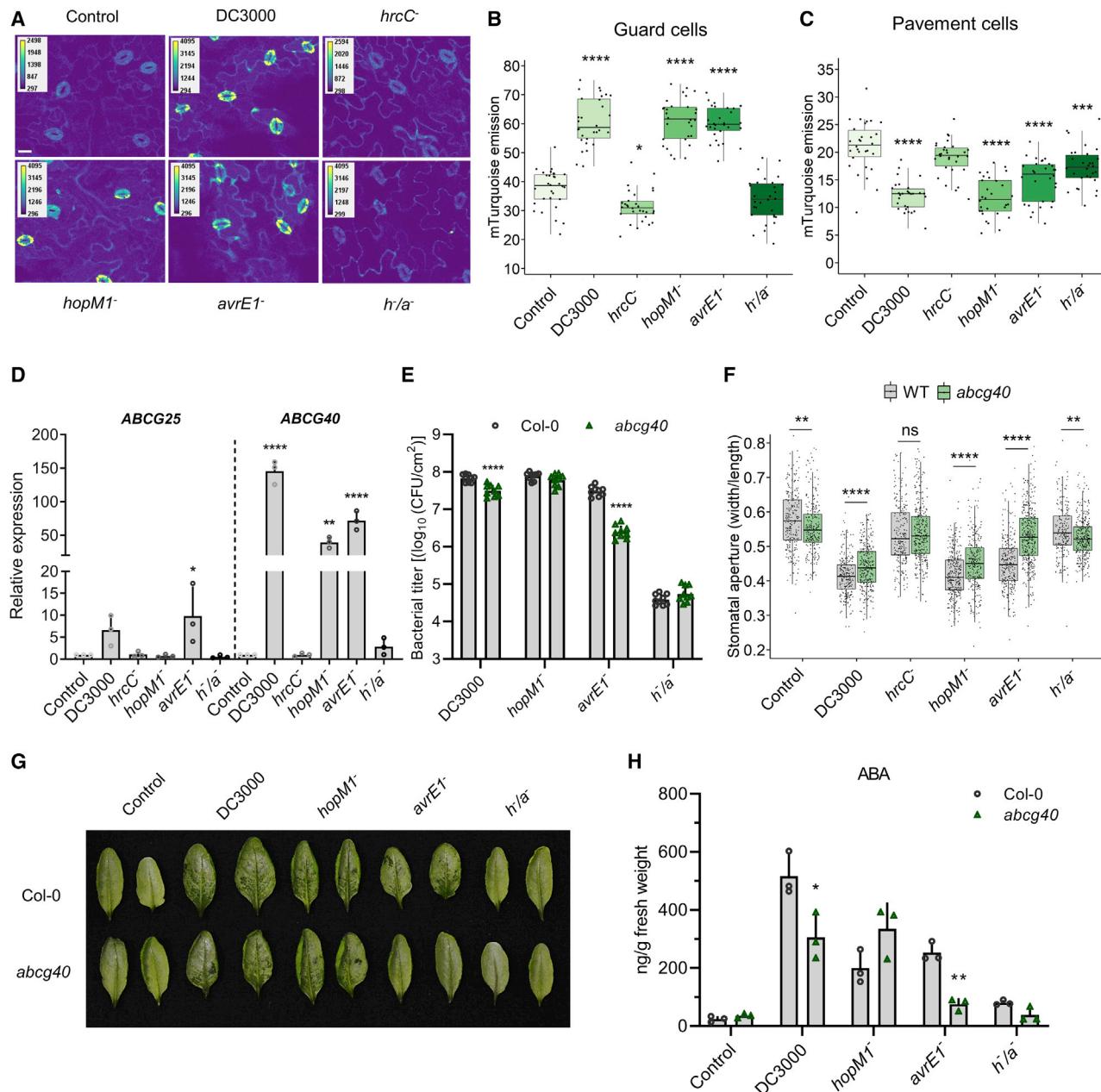
### HopM1 induction of water-soaking lesions requires the guard-cell-specific ABA transporter ABCG40

As guard cells respond rapidly to environmental and hormonal stimuli, we assessed whether HopM1 and AvrE1 increase ABA accumulation ubiquitously or in a more localized manner. To assess this question, we used *Arabidopsis* transgenic lines expressing ABALeon2.1, a recombinant protein reporter that undergoes FRET between mTurquoise and cpV173 fluorophores in the absence of ABA (Waadt et al., 2014). We took advantage of the molecular quenching that occurs between the mTurquoise and cpV173 fluorophores in the presence of ABA to directly visualize ABA concentration changes *in planta*. We found that mTurquoise emission rates were significantly higher in plants infiltrated with *Pst DC3000*, *hopM1*<sup>−</sup>, and *avrE1*<sup>−</sup> as compared with those infiltrated with *hrcC*<sup>−</sup>, *h*<sup>−</sup>/*a*<sup>−</sup>, and mock inoculated controls (Figure 6A). These observations suggest that cpV173 is less able to quench mTurquoise emission signal due to a higher quantity of ABA present in these cells. Interestingly, the mTurquoise emission appeared to be much greater in guard cells compared with pavement cells (Figures 6B and 6C).

Plant cells are equipped with a variety of ABA transporters involved in either ABA export or import, often expressed in a cell-type- and/or tissue-specific manner (Kuromori et al., 2018). We explored the possibility that *Pst DC3000* might increase ABA accumulation in guard cells by increasing the expression of a guard-cell-specific ABA transporter. We found that inoculation of plants with *Pst DC3000*, *Pst hopM1*<sup>−</sup>, and *Pst avrE1*<sup>−</sup> induced a dramatic increase in the expression levels of ABCG40, which is involved in stomatal ABA import (Kang et al., 2010) (Figure 6D). In contrast, transcripts encoding the non-tissue-specific ABA exporter ABCG25 were not upregulated by *Pst* infection to the same extent as ABCG40 (Figure 6D). Bacterial growth assays showed that WT *Pst DC3000* grew somewhat less in the *abcg40* mutant compared

solution of  $2 \times 10^8$  CFU/mL of *Pst DC3000* or *hrcC*<sup>−</sup>, as indicated. All plants were subsequently dip-inoculated in H<sub>2</sub>O, ABA, or TBHQ solutions, as in the pre-treatment (post-treatment). (B–E) Bacterial titers were evaluated 3 days post-infection (dpi). Bacterial titers are shown from plants subjected to different combinations of pre- and post-treatments, as indicated, for DC3000 (B and D) or *hrcC*<sup>−</sup> (C and E). Data are represented as mean of total experimental replicates,  $\pm$  SEM.

(F–J) Stomatal apertures measured in WT *Arabidopsis* plants treated as in (B–E). Apertures were measured 3 h after pre-treatment with H<sub>2</sub>O, ABA, or TBHQ (F). (G and H) Plants were pre-treated with H<sub>2</sub>O, ABA, or TBHQ, as indicated, followed by bacterial inoculation. Stomatal apertures were measured 3 h after inoculation with *Pst DC3000* (G) or with *hrcC*<sup>−</sup> (H). (I and J) Plants were pre-treated with H<sub>2</sub>O only, followed by inoculation with *Pst DC3000* (I) or with *hrcC*<sup>−</sup> (J) and post-treatment with H<sub>2</sub>O, ABA, or TBHQ, as indicated. Stomatal apertures were measured 3 h after post-treatment. Data are represented as median values, with minimum and maximum values indicated (F–J). Different letters indicate statistically significant differences,  $p < 0.05$ , one-way ANOVA.



**Figure 6. *Pseudomonas syringae* enhances ABA accumulation in guard cells and requires the ABA transporter ABCG40 to induce water-soaking lesions**

(A) Confocal microscopy images of *Arabidopsis* Col-0 leaves expressing the ABAeon2.1 reporter, mock inoculated (control) or inoculated by syringe with different *Pst* strains, as indicated. Images represent the emission of mTurquoise following excitation to evaluate the quenching of cpv173 fluorescence in the presence of ABA. Leaves were infected with  $1 \times 10^8$  CFU/mL of bacteria and harvested at 24 hpi before being mounted on microscopy slides. Bar scale represents 20  $\mu$ m.

(B and C) mTurquoise emission in guard cells (B) and pavement cells (C) of plants infected and harvested as in (A). Each data point represents mean of mTurquoise emission in specific cells of one image ( $n = 30$  images/condition).

(D) Relative expression at 24 hpi of the guard cell ABA transporters involved in export (ABCG25) or import (ABCG40) in leaves inoculated with the indicated *Pst* strains ( $1 \times 10^8$  CFU/mL).

(E) Bacterial titers in Col-0 and *abcg40* mutant plants inoculated with  $1 \times 10^5$  CFU/mL of the indicated *Pst* strains at 3 dpi.

(F) Stomatal aperture measurements in WT and *abcg40* mutant plants infected with the indicated *Pst* strains at 24 hpi ( $1 \times 10^8$  CFU/mL).

(G) Water-soaking phenotypes in Col-0 and *abcg40* mutants evaluated in (F).

(H) ABA quantification in Col-0 and *abcg40* mutant plants at 24 hpi with the indicated *Pst* strains ( $1 \times 10^8$  CFU/mL). Data are represented as median values, with minimum and maximum values indicated (B, C) or as mean  $\pm$  SEM (D, E, H). \* $p < 0.05$ , \*\* $p < 0.01$ , \*\*\* $p < 0.005$ , \*\*\*\* $p < 0.0005$ , one-way ANOVA.

with WT, whereas *Pst hopM1<sup>-</sup>* growth was unaffected (Figure 6E). In contrast, *Pst avrE1<sup>-</sup>* displayed a more dramatic growth deficiency in *abcg40* plants, compared with Col-0 (Figure 6E). Consistent with this, *Pst avrE1<sup>-</sup>* can no longer induce stomatal closure or water-soaking lesions in the *abcg40* mutant, whereas *Pst DC3000* and *Pst hopM1<sup>-</sup>* show only minor difference between WT and *abcg40* plants in these assays (Figures 6F and 6G). In combination with the fact that only the *h<sup>-</sup>/a<sup>-</sup>* double mutant is affected in guard cell ABA accumulation (Figures 6A and 6B), this suggests a certain redundancy between HopM1 and AvrE1 in ABA induction and its concentration in guard cells. Interestingly, in the absence of ABCG40, *Pst avrE1<sup>-</sup>* induces almost no accumulation of ABA in contrast to *Pst hopM1<sup>-</sup>* (Figure 6H). Thus, since HopM1 cannot compensate for a lack of AvrE1 in the *abcg40* mutant, this suggests that the effects of HopM1 on ABA-associated virulence are mediated at least in part through ABCG40.

## DISCUSSION

High humidity following consecutive rains is a factor well known to contribute to disease development in the field (Aung et al., 2018). Such conditions are often associated with the induction of water-soaking lesions, which are thought to be crucial for virulence (Xin et al., 2016). Homologs of HopM1 and AvrE1 are present in the genomes of a variety of bacterial pathogens, and functional homologs of AvrE1 have been identified in oomycete pathogens (Deb et al., 2018; Degrave et al., 2015). Here, we provide evidence that HopM1 and AvrE1 induce stomatal closure upon pathogen entry, which prevents transpiration, resulting in an accumulation of water in the apoplast. Through transcriptome analysis, we show that HopM1 and AvrE1 act redundantly in affecting the *Arabidopsis* transcriptome (Figure 1A). Among the most prominent differences in the transcriptional signatures induced by WT *Pst* versus *Pst* lacking HopM1 and AvrE1, we found an important induction of genes associated with ABA-related pathways (Figure 1D). Some transcriptional changes between WT and mutant strains could be due to the differences in levels of bacteria present in the leaf at the time of sampling (36 h). However, ABA-related gene expression, as well as ABA levels, are clearly increased at earlier time points when bacterial levels are similar (Figures 1E–1H and 2F). It has been previously reported that transgenic expression of the *Pst* effector AvrPtoB leads to increased ABA biosynthesis in *Arabidopsis* (de Torres-Zabala et al., 2007). However, we found that expression levels of the ABA biosynthesis marker gene *NCED3* was unaltered in *Arabidopsis* plants infected with an *avrPtoB* mutant or with an effector-less strain in which AvrPto and AvrPtoB have been reintegrated (Figures S3C and S3D). Interestingly, ABA has also been implicated in the induction of water-soaking lesions in rice, and rice mutants with reduced ABA synthesis show decreased growth of *Xanthomonas* bacteria, likely due to increased transpiration rates (Peng et al., 2019; Zhang et al., 2019). Thus, manipulating the ABA pathway to mediate stomatal closure may be a widely used strategy employed by pathogens to induce water-soaking lesions in plants. At the same time, although stomatal closure induced by ABA or TBHQ is sufficient to induce water soaking and increase pathogenicity (Figures 5 and S4), additional mechanisms undoubtedly contribute to creating aqueous apoplastic environments, such as promoting

water intake in the apoplast (Schwartz et al., 2017). Whether such mechanisms are dependent on ABA or not remains to be elucidated.

Considering that many bacterial pathogens have evolved to overcome stomatal immunity by producing diverse toxins, such as COR and SyrA, manipulating ABA to close stomata while producing COR might seem counterintuitive (Melotto et al., 2006; Schellenberg et al., 2010). The actions of such toxins are presumably required to allow pathogens access to the leaf interior only in the early stages of infection and would be detrimental to water-soaking induction at later stages. Indeed, *Pst DC3000* can reopen stomata previously closed by ABA or TBHQ (Figures 5G and 5H). Thus, at 3 h post-treatment, stomatal opening activity appears to dominate. The interplay between COR and ABA at this early phase remains to be elucidated, however, as COR is barely detectable *in planta* at this time point (de Torres Zabala et al., 2009) (Figure S5E). Nonetheless, by measuring the levels of COR and ABA present in late-stage infected leaves, we find that physiologically relevant levels of ABA could override COR-induced stomatal closure. This would be consistent with a chain of events wherein COR allows the pathogen to gain access to the apoplast during the early stage of an infection but whose stomatal opening activity is subsequently countered by ABA after an eventual accumulation of T3SS effectors in infected tissues. At the same time, COR appears to have stomate-independent roles in virulence by interfering with immune signaling in the post-invasive stages (Brooks et al., 2005; Mine et al., 2017; Zheng et al., 2012). However, with respect to stomata, in later stages ABA signaling would appear to predominate over COR signaling, effectively closing the doors previously opened by COR.

Growth levels of *Pst hrcC<sup>-</sup>* in *Arabidopsis* plants treated post-infection with ABA reached those of *Pst DC3000* in untreated plants (Figures 5B and 5C). When stomatal closure and water soaking was induced with TBHQ, a similar effect was seen on *Pst hrcC<sup>-</sup>* growth, but to a lesser extent (Figures 5D and 5E). We interpret the greater effect of ABA on bacterial growth as being consistent with the ability of ABA to interfere with immune signaling in addition to inducing water-soaking lesions (de Torres Zabala et al., 2009).

We found that ABA appears to accumulate to a greater degree in guard cells than pavement cells in infected *Arabidopsis* leaves (Figures 6A–6E). One possible explanation for this would be that HopM1 and AvrE1 might be delivered in a cell-type-specific manner by the pathogen to induce ABA specifically in stomata. Alternatively, this could be due to a concentration of ABA in guard cells through the activity of ABA transporters. ABCG40, also known as PDR12, has previously been identified as an ABA transporter specifically expressed in guard cells (Kang et al., 2010). We found that *Pst DC3000* induces an increase of ABCG40 expression levels in a HopM1- and AvrE1-dependent manner. Moreover, *Pst avrE1<sup>-</sup>*, but not *Pst hopM1<sup>-</sup>*, was unable to cause stomatal closure or water-soaking lesions in an *abcg40* mutant plant. This suggests that HopM1 requires the presence of ABCG40 to induce stomatal closure. Interestingly, ABCG40 has been identified as an important contributor to plant immunity alongside PEN3 against *Botrytis cinerea* via camalexin secretion (He et al., 2019). Thus, ABCG40 might have different functionalities in host-pathogen interactions depending on the type of interaction. Another explanation for the observation of specific

ABA accumulation in guard cells may be related to global control of ABA movement and homeostasis. It has been proposed that during a drought response, less ABA is imported into mesophyll cells, whereas its accumulation reaches high levels in guard cells, possibly due to the activity of ABCG40 (Kang et al., 2010; Zhang et al., 2021). Therefore, HopM1 and AvrE1 modulation of ABA accumulation in guard cells could be attributable to the fact that they may prompt a drought-like transcriptional response during an infection. Indeed, it has been reported that the *Pst* transcriptional signature resembles what is observed during a drought stress (Gupta and Senthil-Kumar, 2017). Nonetheless, regardless of the redundancy of both effectors in inducing ABCG40 expression, the requirement for ABCG40 for HopM1-induced disease development appears to represent a divergence in the functions of these two effectors.

Previous studies have identified MIN7 and PP2A subunits as HopM1 and AvrE1 interactors, respectively (Jin et al., 2016; Nomura et al., 2006). *Arabidopsis min7* mutants have previously been reported to display partial water-soaking lesions following infiltration with *Pst* *h<sup>-</sup>/a<sup>-</sup>* (Xin et al., 2016). The *Arabidopsis min7* mutant is highly sensitive to changes in growth conditions (Nomura et al., 2011), and under the growth conditions used in this study for stomatal analyses, infection with *Pst* *h<sup>-</sup>/a<sup>-</sup>* did not lead to water-soaking lesions in the *min7* mutant. In these conditions, *min7* mutants showed no significant differences in either stomatal closure or in ABA accumulation following infection with *Pst* DC3000, *h<sup>-</sup>/a<sup>-</sup>*, or *hrcC<sup>-</sup>* when compared with WT plants (Figures S6A and S6B). Interestingly, however, *min7* mutant plants display an ABA hypersensitive phenotype (Figure S6C). Furthermore, in growth conditions where *min7* mutants display partial water-soaking lesions upon infection with *Pst* *h<sup>-</sup>/a<sup>-</sup>*, although there was no significant difference in the accumulation of *NCED3* transcripts between plants infected with *Pst* DC3000 or *h<sup>-</sup>/a<sup>-</sup>* at 24 h post-infection, intermediate *NCED3* transcript levels were observed at 7 h post-infection in *min7* plants (Figures S6D and S6E). Because HopM1 targets multiple host proteins (Nomura et al., 2006), the partial water-soaking phenotype of the *min7* mutant indicates that MIN7 is involved in a water-soaking step that does not lead to full-scale ABA induction. Water soaking induced by *Pst* *h<sup>-</sup>/a<sup>-</sup>* may be due in part to the fact that *min7* mutants are hypersensitive to ABA (Figure S6C), and future research is needed to determine which other HopM1 host targets contribute, additively or synergistically, to full-scale water soaking.

The interactions between several PP2A subunits and AvrE1 could also be linked to increased ABA signaling. Indeed, PP2A catalytic subunits have been associated with negative regulation of ABA signaling (Hu et al., 2014; Luo et al., 2006; Pernas et al., 2007). At the same time, it has been reported that AvrE1 targets type one protein phosphatases (TOPPs), which are also negative regulators of ABA signaling (Hu et al., 2022). Ablation of TOPPs leads to enhanced water soaking and stomatal closure, even in the absence of the pathogen, and recues the virulence phenotype of *Pst* *h<sup>-</sup>/a<sup>-</sup>*. These results further highlight the important role for ABA production and signaling in the establishment of an aqueous living space for the pathogen.

In conclusion, our data provide a comprehensive portrait of stomatal behavior during an infection and highlight the role of stomatal movement manipulation in bacterial pathogenesis. How HopM1 and AvrE1 induce the expression of ABA biosyn-

thesis and transporter genes remains to be investigated. By understanding how pathogens induce water-soaking lesions, we anticipate that broad disease resistance in plants may be achieved by interfering with these mechanisms.

## STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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## SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.chom.2022.02.006>.

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## AUTHOR CONTRIBUTIONS

C.R.-L. and P.M. conceived the project. C.R.-L., S.Y.H., and P.M. designed the experiments. C.R.-L. performed RNA-seq analysis. C.R.-L. and M.St.-A. performed bioinformatic analysis. C.R.-L., G.L., V.P.V., M.St.-A., G.S.-M., K.N., S.B., and A.B. performed the experiments. All authors provided feedback on the manuscript. S.Y.H. and P.M. supervised the project. C.R.-L. and P.M. wrote the manuscript.

## DECLARATION OF INTERESTS

The authors declare no competing interests.

## INCLUSION AND DIVERSITY

One or more of the authors of this paper self-identifies as an underrepresented ethnic minority in science.

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## STAR★METHODS

## KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
<b>Bacterial and virus strains</b>		
<i>Pst</i> DC3000 wild-type	Cuppels, 1986	N/A
<i>Pst hopM1</i> <sup>-</sup>	Badel et al., 2003	CUCPB5368
<i>Pst avrE1</i> <sup>-</sup>	Badel et al., 2006	CUCPB5374
<i>Pst hopM1</i> <sup>-</sup> / <i>avrE1</i> <sup>-</sup>	Badel et al., 2006	CUCPB5377
<i>Pst hrcC</i> <sup>-</sup>	Wei et al., 2000	N/A
<i>Pst</i> DB29	Brooks et al., 2004	N/A
<i>Pst</i> <i>avrPtoB</i>	Lin and Martin, 2005	N/A
<i>Pst</i> D28E + AvrPtoB	Cunnac et al., 2011	CUCPB6012
<b>Biological samples</b>		
Promix Premium Potting Mix	PremierTech Inc.	N/A
SUREMIX	Michigan Grower Products Inc.	N/A
<b>Chemicals, peptides, and recombinant proteins</b>		
Abscisic acid	Millipore-Sigma	CAT#C8115
Coronatine	Millipore-Sigma	CAT#45511
Jasmonic acid	Millipore-Sigma	CAT#14631
Fluridone	Millipore-Sigma	CAT#45511
Water suitable for HPLC	Millipore-Sigma	CAT#270753
Methanol	Millipore-Sigma	CAT#34860
Rhodamine 6G	Millipore-Sigma	CAT#R4127
Formaldehyde	Millipore-Sigma	CAT#252549
Dexamethasone	Millipore-Sigma	CAT#D4902
tert-Butylhydroquinone	Millipore-Sigma	CAT#112941
QIAzol Reagent	QIAGEN	CAT#79306
Advanced Q-PCR Mastermix With Supergreen LO-ROX	Wisent Bioproducts	CAT#800-431-UL
SuperScript™ II Reverse Transcriptase	ThermoFisher Sci.	CAT#18064022
Ambion™ DNase I (RNase-free)	ThermoFisher Sci.	CAT#AM2222
Jasmonic acid – isoleucine	Cayman Chemical	CAT#10740
Abscisic acid-d6	Cayman Chemical	CAT#29093
<b>Critical commercial assays</b>		
NEBNext® Multiplex Oligos for Illumina	New England BioLabs	E7335L
<b>Deposited data</b>		
GEO Submission (RNA-Seq)		GSE186836
<b>Experimental models: Organisms/strains</b>		
<i>A. thaliana</i> : Col-0 wild-type	TAIR	CS70000
<i>A. thaliana</i> : <i>aba2-1</i> ethylmethane sulfonate mutant	TAIR	CS156
<i>A. thaliana</i> : <i>abcg40</i> TDNA insertion mutant	TAIR	SALK_148565C
<i>A. thaliana</i> : <i>min7</i> TDNA insertion mutant	Nomura et al., 2006	N/A
<b>Oligonucleotides</b>		
All primers are listed in Table S3		N/A
<b>Software and algorithms</b>		
R statistical environment	<a href="https://www.r-project.org/">https://www.r-project.org/</a>	N/A
CutAdapt	Martin, 2011	N/A
featureCounts	Liao et al., 2014	N/A
DESEQ2	Love et al., 2014	N/A

(Continued on next page)

**Continued**

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Galaxy	Blankenberg et al., 2010	N/A
PANTHER	Mi et al., 2013	N/A
REVIGO	Supek et al., 2011	N/A

**RESOURCE AVAILABILITY**

**Lead contact**

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Peter Moffett ([peter.moffett@usherbrooke.ca](mailto:peter.moffett@usherbrooke.ca)).

**Materials availability**

This study did not generate new unique reagents.

**Data code availability**

- RNA-seq data have been deposited at GEO and are publicly available as of the date of publication. Accession numbers are listed in the [key resources table](#). Microscopy data reported in this paper will be shared by the lead contact upon request.
- This paper analyzes existing, publicly available data (GSE115991) from ([Winkelmüller et al., 2021](#)).
- This paper does not report original code.
- Any additional information required to reanalyze the data reported in this work paper is available from the Lead Contact upon request.

**EXPERIMENTAL MODEL AND SUBJECT DETAILS**

**Plant material and growth conditions**

The *Arabidopsis thaliana* (L.) Heynh. ecotype Columbia (Col-0) was used in this study. The T-DNA insertional mutants ([Alonso et al., 2003](#)) CS156 (*AtABA2*) and SALK\_148565C (*AtABCG40*) were obtained from the Arabidopsis Biological Resource Center (ABRC, <https://abrc.osu.edu/>).

*Arabidopsis thaliana* plants were grown in Promix™ Premium potting mix (PremierTech, Rivière-du-Loup, QC) in Sherbrooke or in “*Arabidopsis Mix*” soil (equal parts of SUREMIX [Michigan Grower Products Inc., Galesburg, MI], medium vermiculite and perlite) at Michigan State University and Duke University in growth chambers with 12 hours light/dark photoperiod, with relative humidity of approximately 60% at 21°C. *Nicotiana benthamiana* and *Solanum lycopersicum* plants were grown in BM6 soil (Berger) in growth chambers under with 16 hours light/8 hours dark photoperiod, with relative humidity of approximately 60% at 23°C.

Four to five-week-old *Arabidopsis* plants were used for bacterial disease assay and stomatal aperture measurement assay. Six to seven-week-old *Nicotiana* and *Solanum* plants were used for stomatal aperture measurement assay.

**Bacterial strains**

*Pseudomonas syringae* pv. *tomato* DC3000 strains were from laboratory stocks and originated from studies described in the [key resources table](#).

**METHOD DETAILS**

**Stomatal aperture assay**

Leaves were cut at the base of the petiole and immediately immersed in stomatal fixation solution (formaldehyde 4%, rhodamine 6G 0.5 µM) for 1 min to stop stomatal movement and excess of solution was removed by blotting. A quarter of each leaf was cut with a razor blade and stomata observed by epifluorescence microscopy (see [Figure S2A](#)). Stomatal aperture was measured by using the software OMERO. Between 150 to 400 stomata were measured for each data point. Data collection and analysis was performed by using double-blinded standards to avoid bias.

For analysis of stomata in infected leaves, plants were kept at room humidity levels for two hours to allow infiltrated apoplastic liquid to dissipate following bacterial infiltration. Once apoplastic water evaporated, plants were domed and kept at 95% relative humidity (RH) levels to allow development of disease symptoms, unless stated otherwise.

**Bacterial disease assay**

*Pst* DC3000 and mutant strains were cultured overnight at 28°C in Luria-Bertani (LB) media containing 50 mg/L of rifampicin (and/or other appropriate antibiotics if necessary). On the day of the infection, fresh LB media was inoculated with 0.5 mL of the overnight

culture and bacteria collected when OD<sub>600</sub> reaches between 0.8-1. Bacteria were centrifuged at 4000 g for 10 minutes and the pellet resuspended in MgCl<sub>2</sub> 10 mM. Bacterial density was adjusted to 0.2 (~1 × 10<sup>8</sup> CFU/ml) prior to further dilutions.

For syringe infiltration, bacterial suspensions were infiltrated directly into the apoplast. Infiltrated plants were kept under ambient humidity levels for 1-2 h to allow water to evaporate and let plant leaves return to a pre-infiltration appearance, unless stated otherwise. Thereafter, plants were domed with a plastic unit to maintain high humidity (>95% RH), unless stated otherwise. For dip-inoculation method, plants were dipped in the bacterial suspension of OD<sub>600</sub> = 0.2, with 0.025% Silwet L-77 added. Plants were domed as mentioned previously once surface liquid had evaporated to allow disease to develop.

Bacterial populations were monitored by harvesting infected *Arabidopsis* leaves, surface sterilizing in ethanol 80% for one minute and rinsing in sterile water twice. Leaf disks were taken from three leaves from the same plant (one per leaf; total of three leaf disks) using a cork borer (0.58 mm in diameter) and ground in sterile MgCl<sub>2</sub> 10 mM. This served as biological replicates. Three biological replicates were performed for each biological experiment. Colony-forming units were determined by making serial dilutions (10<sup>0</sup>-10<sup>-6</sup>) and plating on LB plates containing 50 mg/L of rifampicin. Each dilution was plated in three technical replicates. Experiments were at least three times for biological repeatability.

### Water loss assay

*Arabidopsis* leaves were detached at the base of the petiole and weighed over time to quantify water loss as described previously (Takahashi et al., 2018).

### Chemical treatments

To assess the potential of ABA and TBHQ in inducing water-soaking lesions, plants were either infiltrated, sprayed or dipped with/in an ABA (30 µM) or TBHQ (100 µM) solution. For spraying and dipping experiments, Silwet L-77 (0.025%) was added to each solution.

Inhibition of the ABA biosynthesis pathway was made possible by syringe infiltrating fluridone (10 µM) in *Arabidopsis* leaves.

### RNA extraction and qPCR

RNA was extracted from flash-frozen, ground leaf tissue followed with QIAZOL (QIAGEN) reagents followed by on-column DNase treatment (QIAGEN), according to the manufacturer's protocol. RNA purity was assessed with a spectrophotometer and quality by gel electrophoresis. cDNA was generated by using 5X All-In-One RT Master Mix (ABM).

Quantitative real-time PCR was performed with a Bio-Rad CFX96 machine. Each reaction was composed to 1X Wisent Advanced SYBR Master Mix, specific primers and a 1:20 dilution of 500 ng of cDNA stock. Amplification cycle setups will be as follow: 2 min at 95°C; 40 cycles of 6 seconds at 95°C and 30 seconds at 60°C. Melting curves were verified at the end of the 40 cycles for the confirmation of primer specificities. All reactions were performed in three technical and biological replicates. Three biological replicates were performed to confirm significance and repeatability. Average Cq values were normalized by ΔΔCT formula against the indicated reference gene ACT2. Primers used in this study are listed in Table S3.

### RNA-sequencing and data analysis

RNA integrity was evaluated by an Agilent Bioanalyzer 2100 with the Eukaryote Total RNA Nano Series II. cDNA libraries were generated with NEBNext® Multiplex Oligos for Illumina®, according to the manufacturer's protocol. cDNA libraries were sequenced by RNA-seq at the Université de Sherbrooke RNomics Platform using an Illumina NextSeq 500 system with 43-bp strand-specific paired-end read. Approximately 20 million reads were generated per sample.

Reads quality was assessed using FastQC and low-quality sequences removed by using cutadapt with a quality cutoff of 30. The resulting reads were mapped onto the *Arabidopsis thaliana* genome (TAIR10) using RNA STAR. Mapped reads were counted using featureCounts. Differential gene expression analysis was performed by using the DESeq2 package. A cutoff of q-value <0.01 and absolute log<sub>2</sub> fold change > 1 was applied to identify DEGs. The R function plotPCA was used for principal component analysis. GO analysis was performed with the R package for PANTHER/REVIGO. Differentially expressed genes (DEG) are listed in Table S2.

### Phytohormone extraction and quantification

Fully expanded four weeks-old *Arabidopsis* leaves were harvested and weighed for fresh weight calculation and immediately flash-freeze in liquid nitrogen. Tissues were ground with a plastic pestle and phytohormones extracted overnight using 0.5-1 ml of ice-cold extraction buffer (methanol: water (80:20 v/v), 0.1% formic acid, 0.1 g/L butylated hydroxytoluene and 100 nM ABA-d<sub>6</sub> as an internal standard). Extracted phytohormones were filtered using centrifugal filter units.

Filtered extracts were quantified using an Acquity Ultra Performance Liquid Chromatography system (Waters Corporation, Milford, MA) as described previously (Huot et al., 2017) with some modifications. Briefly, capillary, cone and extractor voltage were set at 3.5 kV, 25 V and 5 V, respectively. Desolvation and cone gas will be set to a flow rate of 600 L/hrs and 50 L/hrs, respectively. Selected ion monitoring was conducted in the negative ES mode for all processed analytes (SA, SAG, ABA, JA, JA-Ile, COR). Analyte responses based on peak area integrations relative to internal standard was determined by using QuanLynx 4.1 software (Waters, Milford, MA). All analytes were quantified based on their standard curve to calculate sample concentration (nM), which was converted to ng using the molecular weight of each specific compound and the extraction volume used. All data was normalized to initial fresh weight in grams.

**ABA visualization by molecular quenching**

Four-week-old *Arabidopsis* leaves expressing the ABAlon2.1 construct were harvested with forceps and a quarter of the leaf cut with a razor blade before being mounted on a microscopy slide to avoid any major veins. mTurquoise was excited at a wavelength of 440 nm and emission was monitored at 476 nm. Images were acquired on a FV-3000 Olympus confocal microscope. Images were analyzed and quantified using ImageJ software [National Institutes of Health (NIH)].

**QUANTIFICATION AND STATISTICAL ANALYSIS**

All experiments were carried out at least three times with at least three biological replicates for each experiment. Statistical significances correspond to Tukey's HSD test, Student's T-test, One-way or Two-way ANOVA as indicated in the respective figure legends. Statistical analyses were performed using the GraphPad Prism 8.4.3 software.

For bacterial titer quantification, dots in figures represent nine plants from three independent experimental replicates. For stomatal aperture assays, each dot in figures represent one stomate. At least 50 stomates were quantified per biological replicate and carried out three times (> 150 stomates/experiment). For phytohormone and COR quantification, data points represent three biological replicates from a representative experimental replicate.