

# RBOHF activates stomatal immunity by modulating both reactive oxygen species and apoplastic pH dynamics in *Arabidopsis*

Dominique Arnaud<sup>\*</sup> , Michael J. Deeks  and Nicholas Smirnov<sup>\*</sup> 

Biosciences, Faculty of Health and Life Sciences, University of Exeter, Exeter EX4 4QD, UK

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<sup>\*</sup>For correspondence (e-mail [n.smirnov@exeter.ac.uk](mailto:n.smirnov@exeter.ac.uk); [dominiquearnaud.fr@gmail.com](mailto:dominiquearnaud.fr@gmail.com)).

## SUMMARY

Stomatal defences are important for plants to prevent pathogen entry and further colonisation of leaves. Apoplastic reactive oxygen species (ROS) generated by NADPH oxidases and apoplastic peroxidases play an important role in activating stomatal closure upon perception of bacteria. However, downstream events, particularly the factors influencing cytosolic hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) signatures in guard cells are poorly understood. We used the H<sub>2</sub>O<sub>2</sub> sensor roGFP2-Orp1 and a ROS-specific fluorescein probe to study intracellular oxidative events during stomatal immune response using *Arabidopsis* mutants involved in the apoplastic ROS burst. Surprisingly, the NADPH oxidase mutant *rbohF* showed over-oxidation of roGFP2-Orp1 by a pathogen-associated molecular pattern (PAMP) in guard cells. However, stomatal closure was not tightly correlated with high roGFP2-Orp1 oxidation. In contrast, RBOHF was necessary for PAMP-mediated ROS production measured by a fluorescein-based probe in guard cells. Unlike previous reports, the *rbohF* mutant, but not *rbohD*, was impaired in PAMP-triggered stomatal closure resulting in defects in stomatal defences against bacteria. Interestingly, RBOHF also participated in PAMP-induced apoplastic alkalisation. The *rbohF* mutants were also partly impaired in H<sub>2</sub>O<sub>2</sub>-mediated stomatal closure at 100 μM while higher H<sub>2</sub>O<sub>2</sub> concentration up to 1 mM did not promote stomatal closure in wild-type plants. Our results provide novel insights on the interplay between apoplastic and cytosolic ROS dynamics and highlight the importance of RBOHF in plant immunity.

**Keywords:** *Arabidopsis thaliana*, stomatal immunity, reactive oxygen species, *Pseudomonas syringae*, redox biosensor.

## INTRODUCTION

Plants are continuously exposed to diverse microorganisms and must develop strategies to avoid infection by restricting the entry and multiplication of pathogens. The main first line of plant defences against foliar bacteria is the closure of stomatal pores at the leaf epidermis. Stomata are formed by a pair of guard cells that mediate gas exchange between the plant and the environment and are critical during the plant's innate immune response (Melotto et al., 2006; Melotto et al., 2008). Stomatal immunity is initiated by the recognition of pathogen-associated molecular patterns (PAMPs) by pattern recognition receptors at the plasma membrane. In *Arabidopsis thaliana*, the receptor FLAGELLIN-SENSITIVE2 (FLS2) that recognises the peptide flg22, the biologically active epitope of the bacterial PAMP flagellin, plays a prominent role in stomatal immunity (Melotto et al., 2006; Zeng & He, 2010). Downstream of PAMP perception, calcium, reactive oxygen

species (ROS), nitric oxide and apoplastic alkalisation function as secondary messengers to promote stomatal closure (Arnaud & Hwang, 2015). Stomatal immunity was also shown to be modulated by hormonal pathways, mainly the abscisic acid (ABA) and salicylic acid (SA) signalling pathways (Arnaud & Hwang, 2015; Melotto et al., 2008). To counteract PAMP-mediated stomatal closure, pathogens have evolved virulence factors, such as the phytotoxin coronatine (COR) secreted by the bacterial pathogen *Pseudomonas syringae* pv tomato (*Pst*) strain DC3000, to counteract host stomatal defences by promoting stomatal reopening (Melotto et al., 2006).

During the immune response, the apoplastic ROS burst is mainly generated by the NADPH oxidase isoform RBOHD and to a lesser extent RBOHF (Torres et al., 2002; Zhang et al., 2007). While RBOHD participates in stomatal (pre-invasive) defence responses, its contribution to plant resistance against *Pst* bacteria is not clear (Chaouch

et al., 2012; Kadota et al., 2014; Zhang et al., 2007). Moreover, the role of RBOHD in PAMP-triggered stomatal closure has been questioned (Guzel Deger et al., 2015; Khokon et al., 2010). On the contrary, RBOHF is required for full post-invasive (late apoplastic) resistance against virulent *Pst* bacteria (Chaouch et al., 2012) and no role in stomatal defence has been described until now. The type III cell wall peroxidases PRX33 and PRX34 also participate in PAMP-mediated ROS production and play an important role in both pre- and post-invasive defences against *Pst* bacteria (Arnaud et al., 2017; Daudi et al., 2012). Although it is unknown how PAMPs activate PRXs, apoplastic alkalinisation is a prerequisite for PRX-mediated apoplastic ROS production (Bolwell et al., 2002). Apoplastic ROS, mainly hydrogen peroxide ( $H_2O_2$ ), diffuse into the cytosol through plasma membrane aquaporins (Rodrigues et al., 2017; Tian et al., 2016). However, recent results indicate that PAMP- and bacteria-induced oxidation of the cytosolic/nuclear localised  $H_2O_2$  sensor roGFP2-Orp1 in leaf discs is largely independent of RBOHD, RBOHF, PRX33 and PRX34 (Arnaud et al., 2023). Moreover, it remains unclear how apoplastic ROS produced by NADPH oxidases or cell wall peroxidases upon PAMP perception influences intracellular  $H_2O_2$ /ROS accumulation specifically in guard cells.

In this contribution, we used three different probes to analyse oxidative events after flg22 treatment. The chemiluminescent luminol and fluorescein  $H_2DCFDA$  probes detect, respectively, extracellular and intracellular ROS and are not specific to  $H_2O_2$  (Murphy et al., 2022). In contrast, the ratiometric biosensor roGFP2-Orp1 localised in the cytosol and nucleus is mainly oxidised by  $H_2O_2$  *in vivo* and can be reduced by the glutaredoxin/glutathione system permitting dynamic and real-time measurements (Arnaud et al., 2023; Nietzel et al., 2019). We show that PAMP-triggered oxidation of roGFP2-Orp1 is surprisingly stronger in guard cells of the *rbohF*, *prx33-3* and *prx34-2* mutants compared to WT plants. Despite an increase in  $H_2O_2$  in guard cells upon PAMP perception, we found a weak negative correlation between stomatal aperture and roGFP2-Orp1 oxidation. Moreover, roGFP2-Orp1 oxidation state in guard cells was not correlated with a stomatal aperture in control conditions. More importantly, we found that RBOHF plays a crucial role in stomatal immunity and defence against *Pseudomonas syringae* bacteria by a complex control of both ROS production and apoplastic pH.

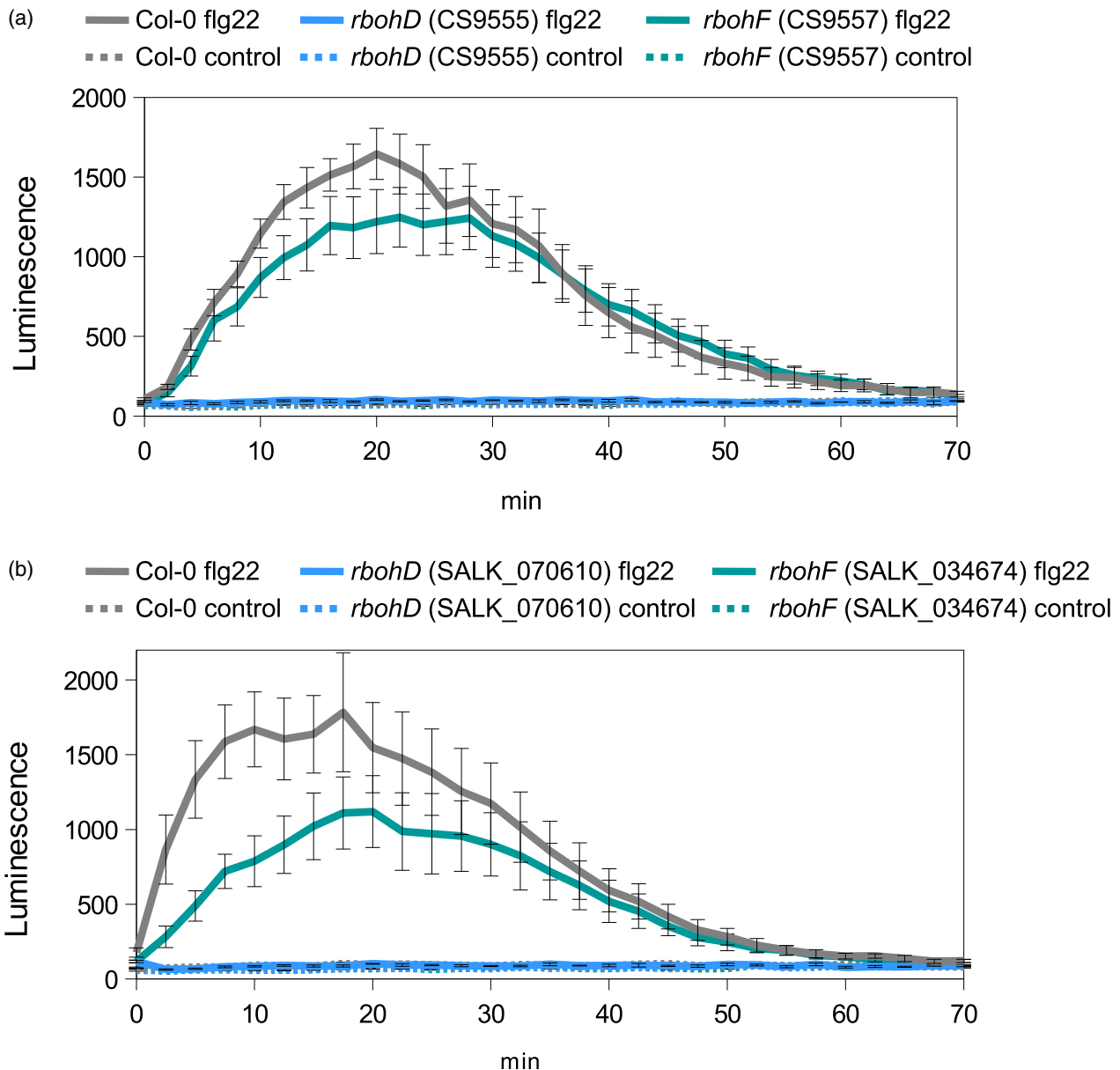
## RESULTS

### PAMP-induced oxidation of the $H_2O_2$ sensor roGFP2-Orp1 is higher in guard cells of the *rbohF* mutant

Using the chemiluminescent luminol probe to detect extracellular ROS production, RBOHD was shown to be the main NADPH oxidase implicated in PAMP-induced ROS

burst in the apoplast of Arabidopsis leaves (Mersmann et al., 2010; Zhang et al., 2007). Indeed, luminol assays performed on leaf discs showed that the *rbohD* mutants were defective in flg22-triggered apoplastic ROS burst while a non-significant decrease in ROS production was observed in the *rbohF* mutants compared to Col-0 WT (Figure 1). Surprisingly, while flg22 causes stomatal closure, ROS production in the cytosol of *rbohD* or *rbohF* guard cells upon flg22 activation has not been reported. Moreover, these two NADPH oxidases were not implicated in intracellular ROS increase in guard cells after treatment with yeast elicitors (Khokon et al., 2010). To analyse the contribution of apoplastic ROS burst to cytosolic  $H_2O_2$  accumulation during the immune response, we used Arabidopsis lines in which the  $H_2O_2$  sensor roGFP2-Orp1 has been introduced into the *rbohD*, *rbohF*, *prx33-3* and *prx34-2* mutants, respectively (Arnaud et al., 2023). We analysed PAMP-triggered roGFP2-Orp1 oxidation in pavement cells and guard cells of leaf discs after dual excitation at 405 nm and 488 nm (emission at 508–526 nm) by confocal laser scanning microscopy (Figure 2, Figure S1). No significant difference in roGFP2-Orp1 oxidation state was observed in *rbohD*, *rbohF*, *prx33-3* and *prx34-2* pavement cells and guard cells as compared to Col-0 in control conditions (Figure S1a,b). The *rbohD*, *prx33-3* and *prx34-2* mutants were not affected in roGFP2-Orp1 oxidation in pavement cells at 60 min after flg22 treatment. Surprisingly, the *rbohF* mutant exhibited a marked increase in flg22-mediated roGFP2-Orp1 oxidation compared to Col-0 in pavement cells (Figure S1a) suggesting lower antioxidant capacities in this mutant as observed before (Arnaud et al., 2023).

In Col-0 guard cells, roGFP2-Orp1 oxidation started to increase from 20 min after flg22 treatment (Figure S1d). The *rbohD* mutant was not impaired in flg22-mediated roGFP2-Orp1 oxidation in guard cells (Figure 2a). By contrast, the *rbohF* mutant showed higher roGFP2-Orp1 oxidation compared to Col-0 WT at 30 and 60 min after flg22 treatment. Similarly, the *prx33-3* and *prx34-2* mutants which are impaired in stomatal immunity (Arnaud et al., 2017) showed a significant increase in flg22-induced roGFP2-Orp1 oxidation in guard cells as compared to Col-0 (Figure S1b). These unexpected results suggest that a defect in PAMP-mediated stomatal closure is correlated with an enhanced  $H_2O_2$  or oxidation state of guard cell cytosol/nucleus. A more detailed comparison of roGFP2-Orp1 oxidation and stomatal aperture of individual guard cells in Col-0 WT after control or flg22 treatment for 30 and 60 min shows that while roGFP2-Orp1 is relatively reduced in control conditions (Figure 2b), there was no correlation between stomatal aperture and roGFP2-Orp1 oxidation ( $R = 0.07$ ,  $P = 0.48$ ). We found a negative correlation, albeit weak, between stomatal aperture and roGFP2-Orp1 oxidation across all flg22 treatments (flg22 30 min,



**Figure 1.** The *rbohD* mutants are defective in PAMP-triggered ROS burst in the apoplast.

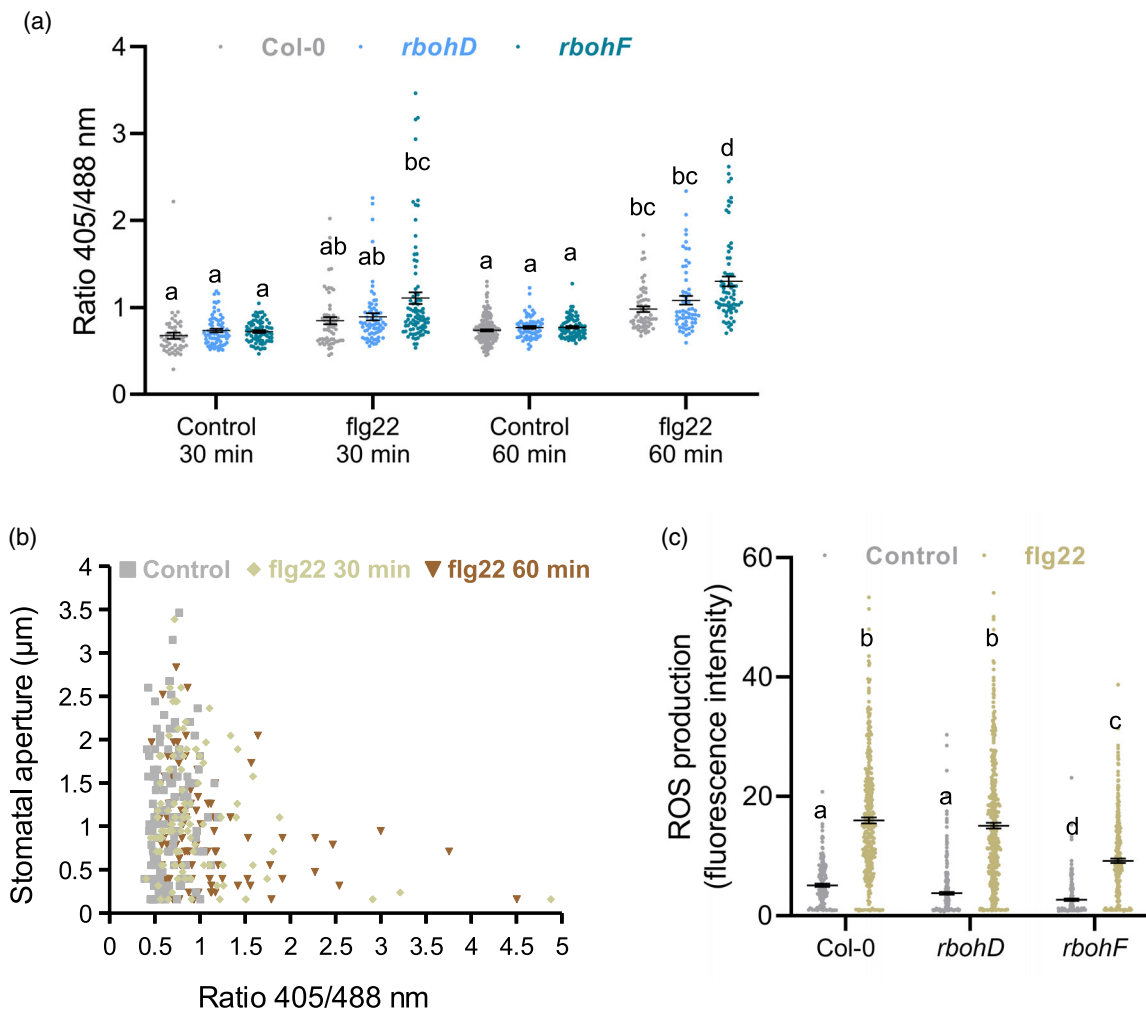
(a and b) PAMP-induced apoplastic ROS production detected by luminol assay in Col-0 WT, *rbohD* (CS9555) and *rbohF* (CS9557) mutants (a) and other allelic *rbohD* (SALK\_070610) and *rbohF* (SALK\_034674) mutants (b). The luminescence was measured over time after treatment with control solution or  $1 \mu\text{M}$  flg22 at  $t = 0$  min. Data are means  $\pm$  SE ( $n = 6$ ) from a representative experiment.

$R = -0.24$ ,  $P = 0.02$  and flg22 60 min,  $R = -0.24$ ,  $P = 0.01$ ) (Figure 2b). Guard cells with roGFP2-Orp1 in a reduced state (either control or flg22-treated) had a wide range of apertures, whereas flg22-treated guard cells with more oxidised roGFP2-Orp1 all exhibited small apertures.

#### **RBOHF is required for PAMP-triggered ROS production in guard cells**

As a complementary approach, we used the fluorescein probe H<sub>2</sub>DCFDA, which is not specific to H<sub>2</sub>O<sub>2</sub> (Murphy et al., 2022), to analyse PAMP-induced intracellular ROS

production in guard cells of NADPH oxidase mutants. H<sub>2</sub>DCFDA assays indicate that although the basal level of ROS was decreased in both *rbohD* and *rbohF* mutants, *rbohF* has a significantly smaller increase in ROS compared to Col-0 by 30 min after flg22 treatment (Figure 2c). These results suggest that only RBOHF contributes to flg22-mediated ROS production in guard cells but RBOHD does not. Importantly, based on H<sub>2</sub>DCFDA assays, PRX33 and PRX34 are also implicated in PAMP-mediated ROS accumulation in guard cells (Arnaud et al., 2017). Therefore, across these mutants, ROS measurement by



**Figure 2.** The  $\text{H}_2\text{O}_2$  sensor roGFP2-Orp1 is more oxidised in guard cells of the *rbohF* mutant.

(a) Oxidation state of roGFP2-Orp1 in Col-0 WT, *rbohD* and *rbohF* guard cells in response flg22. Leaf discs were exposed to control solution or  $1 \mu\text{M}$  flg22 and the ratio 405/488 nm of stomata was quantified at 30 min and 60 min after treatment from images of the fluorescence emission at  $517 \pm 9 \text{ nm}$  following excitation at 488 and 405 nm. Data are means  $\pm$  SE ( $n \geq 50$  guard cells) from a representative experiment. Different letters indicate significant differences at  $P < 0.05$  based on a Tukey's HSD test.

(b) Correlation between the stomatal aperture and the oxidation state of roGFP2-Orp1 in guard cells of leaf discs after treatment with control solution or  $1 \mu\text{M}$  flg22 for 30 min and 60 min. The scatter plots show the stomatal aperture as a function of the ratio 405/488 nm for each condition ( $n \geq 100$ ). The Pearson correlation coefficient ( $R$ ) is 0.0651 ( $P = 0.483$ ) for control treatment and  $-0.2428$  ( $P = 0.015$ ) and  $-0.2395$  ( $P = 0.011$ ) for flg22 treatment at 30 min and 60 min, respectively.

(c) ROS production detected by  $\text{H}_2\text{DCFDA}$  fluorescence in guard cells of Col-0 WT, *rbohD* and *rbohF* epidermal peels 30 min after treatment with control solution or  $5 \mu\text{M}$  flg22. Data are means  $\pm$  SE ( $n \geq 200$ ) from 3 independent experiments. Different letters indicate significant differences at  $P < 0.05$  based on a Tukey's HSD test.

$\text{H}_2\text{DCFDA}$  shows a correlation between defects in ROS production and defects in flg22-induced stomatal closure while, particularly in the case of *rbohF*, cytosolic roGFP2-Orp1 is more oxidised.

#### RBOHF is involved in stomatal defence responses

Stomatal closure is one of the first lines of defence against foliar bacteria (Melotto et al., 2006) and PAMP-mediated stomatal closure has been correlated with increased ROS production in guard cells (Desclos-Theveniau et al., 2012; Desikan et al., 2008). Both NADPH oxidase (RBOHD) and

apoplastic peroxidases (PRX33 and PRX34) participate in stomatal immunity (Arnaud et al., 2017; Macho et al., 2012; Mersmann et al., 2010). Indeed, pre-treatment with the non-specific inhibitor of NADPH oxidases (DPI) or with salicylhydroxamic acid (SHAM) or sodium azide (two non-specific inhibitors of apoplastic peroxidases) (Khokon et al., 2010) compromised flg22-mediated stomatal closure (Figure S2a). Stomatal aperture was analysed in *rbohD* and *rbohF* leaf discs after treatment with flg22 or inoculation with the coronatine-deficient *Pseudomonas syringae* pv. tomato DC3000 (*Pst* COR<sup>-</sup>) bacteria for 2 h (Figure 3a).

A comparison with epidermal peels was also made (Figures S2b,c). Because the toxin coronatine counteracts PAMP-mediated stomatal closure, the *Pst* COR<sup>-</sup> strain has been widely used to characterise mutants defective in stomatal immunity (Desclos-Theveniau et al., 2012; Kadota et al., 2014; Macho et al., 2012; Melotto et al., 2006). Contrary to previous observations showing that RBOHD is required for PAMP-induced stomatal closure (Kadota et al., 2014; Li et al., 2014; Macho et al., 2012; Mersmann et al., 2010), we found that in our experimental conditions, *rbohD* consistently exhibited WT stomatal closure upon PAMP or bacterial treatments on both leaf disc and epidermal peel assays (Figure 3a, Figure S2b,c). On the contrary, the *rbohF* mutant was clearly defective in PAMP- and bacteria-mediated stomatal closure (Figure 3a, Figure S2b,c). These results were confirmed for flg22 using different mutant alleles of these NADPH oxidases (Figure 3b). As a control, we confirmed that *rbohF*, but not *rbohD*, is partially impaired in abscisic acid-mediated stomatal closure (Figure S3a; Kwak et al., 2003; Macho et al., 2012). Moreover, as previously observed (Khokon et al., 2010), these NADPH-oxidases are not involved in salicylic acid-mediated stomatal closure (Figure S3b). Although RBOHD is reported to be more expressed in guard cells than RBOHF (Kwak et al., 2003; Morales et al., 2016), RT-qPCR measurements show that, contrary to RBOHD, RBOHF expression was induced in guard cells 2 h after flg22 treatment (Figure S2d). Compared with the PAMP-mediated induction of RBOHD expression in young seedlings (Morales et al., 2016), our results indicate tissue-specific induction of these two NADPH oxidases after PAMP activation and highlight the importance of RBOHF in stomatal immunity.

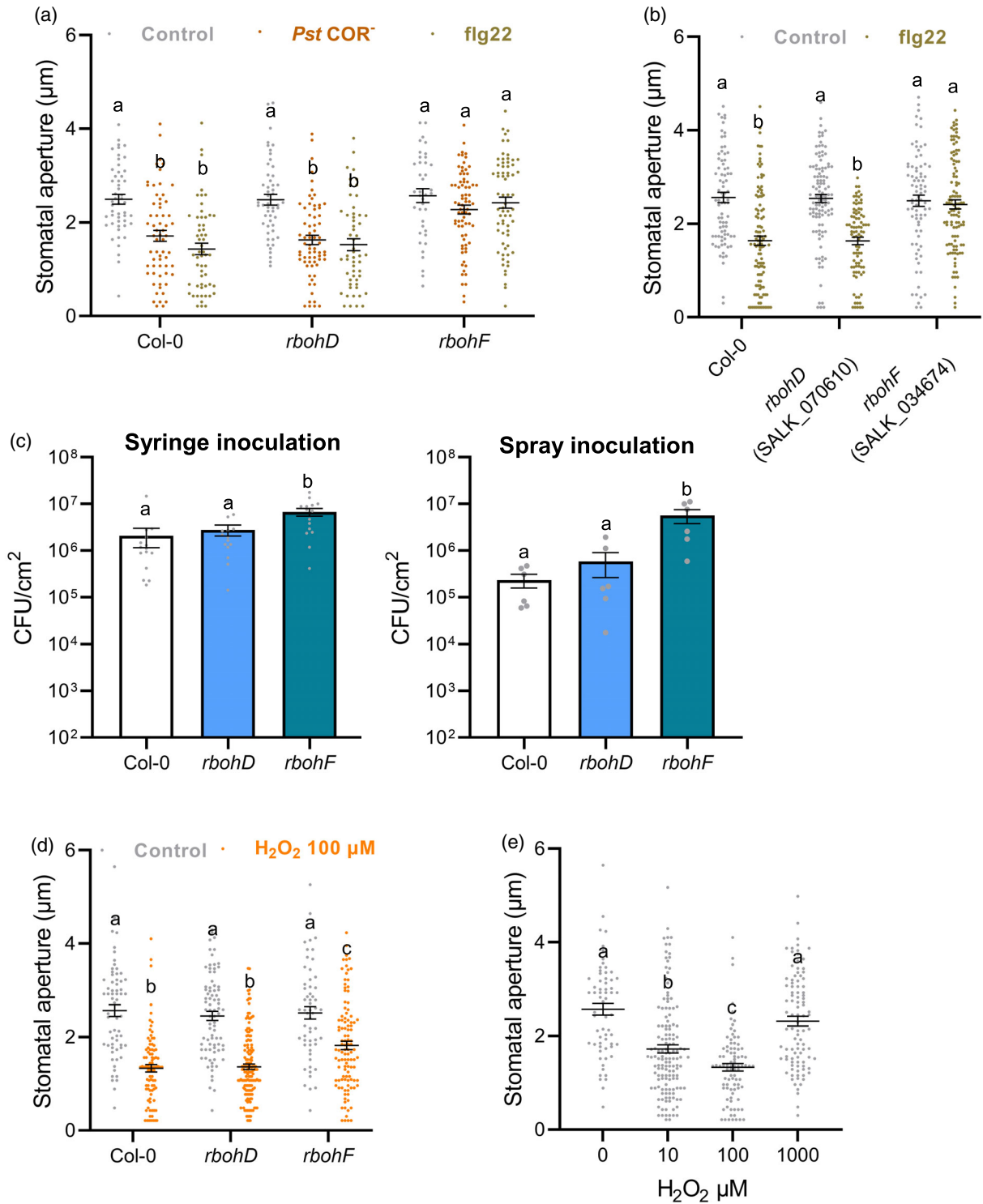
Because a defect in stomatal defence often results in enhanced susceptibility to bacteria (Melotto et al., 2006), leaves were spray-inoculated with *Pst* COR<sup>-</sup> bacteria and bacterial growth was evaluated 3 days later (Figure 3c). The *rbohF* mutant was more susceptible than the Col-0 control to surface-inoculated bacteria, with a 24-fold increase in the bacterial population. As observed before (Kadota et al., 2014; Macho et al., 2012) the *rbohD* mutant was as susceptible as Col-0 to surface-inoculated *Pst* COR<sup>-</sup> corroborating the absence of defects in bacteria-mediated stomatal closure. To verify any deficiency in post-invasive defences, plants were also inoculated with *Pst* COR<sup>-</sup> by syringe infiltration. Similar to previous results with the WT *Pst* DC3000 bacteria (Chaouch et al., 2012), the *rbohD* mutant exhibited WT susceptibility to bacteria while *rbohF* was slightly more susceptible to syringe-inoculated *Pst* COR<sup>-</sup> with a threefold increase in bacterial titre as compared to Col-0 (Figure 3c). These results indicate that RBOHF is important for resistance to *Pst* DC3000 bacteria and contributes more to pre-invasive than post-invasive defence responses.

### The *rbohF* mutant is partially defective in H<sub>2</sub>O<sub>2</sub>-mediated stomatal closure

We also tested the stomatal response of *rbohD* and *rbohF* mutants to H<sub>2</sub>O<sub>2</sub> which closes stomata through the activation of Ca<sup>2+</sup> channels and inhibition of the inward K<sup>+</sup> channels (Pei et al., 2000; Zhang et al., 2001). The results indicate that *rbohF* mutants were partially impaired in H<sub>2</sub>O<sub>2</sub>-mediated stomatal closure while WT stomatal closure was observed for the *rbohD* mutants (Figure 3d, Figure S3c). Therefore, RBOHF may also act downstream of H<sub>2</sub>O<sub>2</sub> during PAMP-mediated stomatal closure. While H<sub>2</sub>O<sub>2</sub> at 100 μM is widely used for inducing stomatal closure, this effect of H<sub>2</sub>O<sub>2</sub> at this concentration is dependent upon the experimental setup (Li et al., 2021; Pei et al., 2000). Moreover, the *rbohF* mutant showed increased oxidation of the H<sub>2</sub>O<sub>2</sub> sensor roGFP2-Orp1 in guard cells after PAMP treatment (Figure 2a) and in whole leaf discs after H<sub>2</sub>O<sub>2</sub> treatment (Arnaud et al., 2023). Therefore, we also tested the effect of different H<sub>2</sub>O<sub>2</sub> concentrations on stomatal aperture in our experimental conditions using Col-0 WT leaf discs in stomatal buffer. Surprisingly, while H<sub>2</sub>O<sub>2</sub> at 10 and 100 μM effectively induced significant stomatal closure, we found that stomatal apertures at a higher H<sub>2</sub>O<sub>2</sub> concentration (1 mM) were similar to the control treatment (Figure 3e). These results suggest that excess H<sub>2</sub>O<sub>2</sub> accumulation in *rbohF* guard cells above a certain threshold is unable to induce stomatal closure.

### The *rbohF* mutant is affected in flg22-mediated apoplastic alkalinisation

Apoplastic alkalinisation is a well-characterised early event induced upon PAMP perception (Felix et al., 1999; Felle et al., 2004) and constitutively active plasma membrane H<sup>+</sup>ATPase OST2/AHA1 impedes PAMP-mediated stomatal closure and apoplastic ROS burst (Keinath et al., 2010; Liu et al., 2009). Otte and collaborators showed that elicitor-induced alkalinisation of the apoplast was inhibited by DPI in chickpea (Otte et al., 2001) and NADPH oxidases could contribute to extracellular pH changes through their electrogenic activities and/or the consumption of protons during superoxide dismutation (Segal, 2016). To test this hypothesis, we used the ratiometric Oregon green fluorescent dye which has been successfully used in Arabidopsis (Geilfus & Muehling, 2011; McLachlan et al., 2016) and adapted the method for multi-well fluorimetry on leaf discs. Interestingly, assays with Oregon green indicate that the apoplastic pH was higher in the *rbohF* mutant in unstressed conditions while it remained unchanged in the *rbohD* background (Figure 4a). This result nicely correlates with the impaired growth of *rbohF* rosette leaves observed previously (Chaouch et al., 2012; Torres et al., 2002) and in our laboratory conditions suggesting that RBOHF is important for the acidic growth of plant cells. Although *rbohD*



and *rbohF* showed WT increase in absolute pH after flg22 treatment (Figure S3d), the flg22-triggered increase in pH relative to the initial pH in unstressed conditions was significantly lower in the *rbohF* mutant compared to Col-0 WT

(Figure 4b). These results suggest that the *rbohF* mutant is partly defective in PAMP-induced apoplastic alkalisation. Therefore, as observed for the constitutive active *ost2-1D* and *ost2-2D* mutants (Liu et al., 2009; Merlot et al., 2007),

**Figure 3.** Analysis of the function of RBOHD and RBOHF during stomatal defence responses.

(a) Stomatal apertures in WT Col-0, *rbohD* and *rbohF* leaf discs exposed to mock control (10 mM MgCl<sub>2</sub>), 10<sup>8</sup> cfu/ml COR-deficient *Pst* DC3000 (*Pst* COR<sup>-</sup>) bacteria and 5 μM flg22 for 2 h. Data are means ± SE ( $n \geq 100$ ) from a representative experiment.  
 (b) Stomatal apertures in Col-0 WT and other allelic mutants of *rbohD* (SALK\_070610) and *rbohF* (SALK\_034674). Epidermal peels were exposed to control solution or 5 μM flg22 for 2 h. Data are means ± SE ( $n \geq 80$ ) from a representative experiment.  
 (c) Bacterial growth in WT Col-0, *rbohD* and *rbohF* mutants assessed at 3 days after syringe-infiltration with *Pst* COR<sup>-</sup> at 10<sup>6</sup> cfu/ml (left panel) or after spray-inoculation with *Pst* COR<sup>-</sup> at 10<sup>8</sup> cfu/ml (right panel). Values are the means ± SE ( $n = 6$ ).  
 (d) The *rbohF* mutant is partially defective in H<sub>2</sub>O<sub>2</sub>-mediated stomatal closure. Stomatal apertures in WT Col-0, *rbohD* and *rbohF* leaf discs exposed to Control or 100 μM H<sub>2</sub>O<sub>2</sub> for 2 h. Data are means ± SE ( $n \geq 60$ ) from a representative experiment.  
 (e) Dose response of Col-0 WT stomata to H<sub>2</sub>O<sub>2</sub> indicate that high H<sub>2</sub>O<sub>2</sub> concentration does not close stomata. Stomatal apertures in WT Col-0 leaf discs exposed to different concentrations of H<sub>2</sub>O<sub>2</sub> (0, 10, 100 and 1000 μM) for 2 h. Data are means ± SE ( $n \geq 60$ ) from a representative experiment. Different letters indicate significant differences at  $P < 0.001$  (A, B, D and E) and  $P < 0.05$  (C) based on a Tukey's HSD test.

defects in apoplastic alkalinisation may also contribute to the impairment of *rbohF* in H<sub>2</sub>O<sub>2</sub>, ABA- and PAMP-mediated stomatal closure.

## DISCUSSION

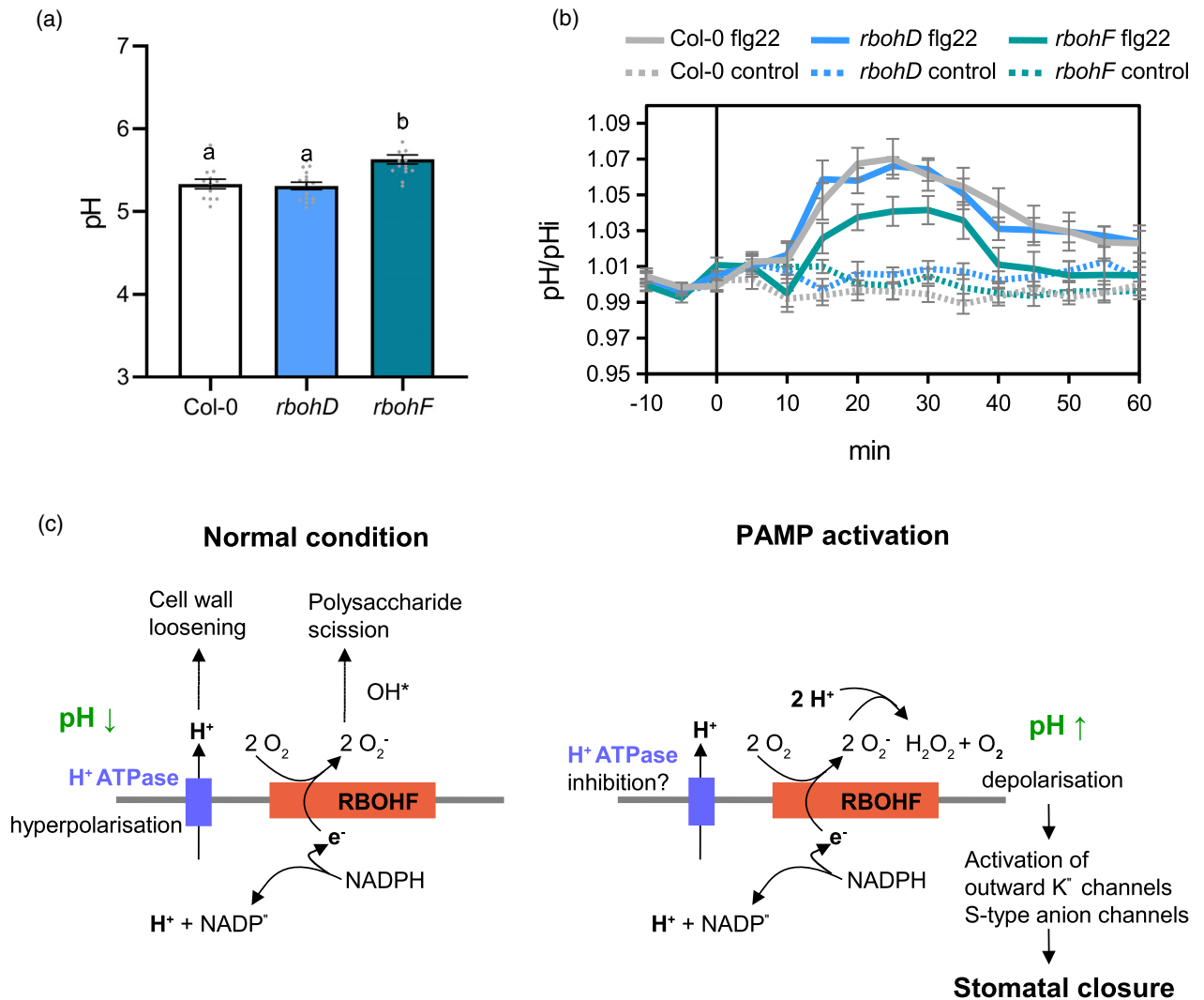
### The role of RBOHF in stomatal defences

PAMP-mediated stomatal closure is considered to involve ROS production (Arnaud & Hwang, 2015). Previous reports suggest that the RBOHD isoform is required for closure, presumably mediated by ROS production initially in the apoplast (Kadota et al., 2014; Li et al., 2014; Macho et al., 2012; Mersmann et al., 2010). However, the defect in PAMP-mediated stomatal closure observed in the *rbohD* mutant was not correlated with increased susceptibility to *Pst* bacteria (Kadota et al., 2014; Macho et al., 2012). A more recent study showed that RBOHD and RBOHF are not involved in flg22-triggered stomatal closure (Guzel Deger et al., 2015). Contrary to previous work, and using multiple mutant alleles, we found that stomatal closure in response to flg22 and to *Pst* COR<sup>-</sup> bacteria depends on RBOHF but not on RBOHD. Correspondingly, there was increased bacterial growth in the *rbohF* mutant but not *rbohD*. It should be noted that the *rbohD* and *rbohF* mutants showed the expected responses for the apoplastic burst in response to flg22. Our results and previous data suggest that the relative contribution of RBOHD and RBOHF in stomatal immunity likely depends on experimental conditions. While unquantified differences in growth conditions may be involved, it should also be noted that immune responses are strongly dependent on the circadian clock which could, for example, mean that different NADPH oxidase isoforms influence responses at different times of day (Zhang et al., 2013).

### Is cytosolic H<sub>2</sub>O<sub>2</sub> important for stomatal closure?

The relationship between ROS production and stomatal closure was complex. The widely used probe H<sub>2</sub>DCFDA, which it is not specific for any particular ROS and not oxidised directly by H<sub>2</sub>O<sub>2</sub> (Murphy et al., 2022) showed decreased ROS production in *rbohF* guard cells in response to flg22, which is commensurate with defects in stomatal closure. roGFP2-Orp1 and GRX1-roGFP2 have been used to demonstrate oxidation in guard cell cytosol

during NaHS-induced stomatal closure (Scuffi et al., 2018). However, during the flg22 response, roGFP2-Orp1 oxidation state did not show a strong relationship with the aperture and the probe was more oxidised in *rbohF* guard cells despite a lower H<sub>2</sub>DCFDA oxidation. As suggested previously for whole leaves (Arnaud et al., 2023; Chaouch et al., 2012), a lower antioxidant capacity in *rbohF* guard cells may explain the higher roGFP2-Orp1 oxidation upon PAMP activation. This could reduce the thiol-reducing capacity affecting roGFP2-Orp1 regeneration without influencing H<sub>2</sub>DCFDA oxidation. Added to this, stomatal aperture in Col-0 had a poor correlation with roGFP2-Orp1 oxidation state in control conditions. Using epidermal strips incubated in stomatal opening buffer, the stomatal aperture seems not to be affected by high roGFP2-Orp1 oxidation in guard cells (Scuffi et al., 2018). A possible explanation is that roGFP2-Orp1 is specifically reporting cytosolic H<sub>2</sub>O<sub>2</sub> which could be derived from multiple compartments. For example, mitochondrial ROS contribute to ABA-mediated stomatal closure (Postiglione & Muday, 2023). The use of catalase mutants has shown that, despite higher intracellular H<sub>2</sub>O<sub>2</sub>, stomatal aperture is unaffected although still responsive to exogenous H<sub>2</sub>O<sub>2</sub> (Jannat et al., 2011). This may explain why stomata remain open despite photosynthetic ROS production in the light. Our results, therefore, support the uncoupling of cytosolic H<sub>2</sub>O<sub>2</sub> from stomatal aperture and that the extracellular production of superoxide or H<sub>2</sub>O<sub>2</sub> by RBOHF is critical for flg22-induced closure. The LRR receptor kinase HPCA1 is a proposed extracellular H<sub>2</sub>O<sub>2</sub> sensor which, when oxidised on extracellular cysteines, activates calcium channels leading to increased intracellular Ca<sup>2+</sup> (Wu et al., 2020). This sensor provides an explanation for the requirement of apoplastic H<sub>2</sub>O<sub>2</sub> for stomatal closure. Intriguingly, our results suggest that the ROS probe H<sub>2</sub>DCFDA could indicate that ROS other than H<sub>2</sub>O<sub>2</sub> are involved in triggering closure while roGFP2-Orp1 indicates cytosolic conditions that are at least partly independent of stomatal aperture. The results with guard cells therefore partly mirror the recent findings with whole leaves (Arnaud et al., 2023) where the RBOHD-dependent apoplastic oxidative burst measured by luminol is independent of subsequent prolonged oxidation of cytosolic roGFP2-Orp1.



**Figure 4.** RBOHF influences apoplastic pH.

(a) Apoplastic pH in Col-0 WT, *rbohD* and *rbohF* mutant leaves. Apoplastic pH in untreated condition was determined by multiwell fluorimetry (sequential excitation at 440 ± 8 and 495 ± 8 nm; emission, 525 ± 20 nm) on Oregon green dextran-infiltrated leaf discs from 5 week-old plants (See Methods for details). Data are means ± SE (*n* ≥ 12) from a representative experiment. Different letters indicate significant differences at *P* < 0.001 based on a Tukey's HSD test.

(b) Kinetics of flg22-induced leaf apoplastic alkalisation in Col-0 WT, *rbohD* and *rbohF* mutants. Oregon green dextran-infiltrated leaf discs were exposed at *t* = 0 min to control solution or 1 μM flg22, the apoplastic pH was measured over time by multiwell fluorimetry and expressed relative to the mean initial pH (pHi) before treatment (pH/pHi). Data are means ± SE (*n* ≥ 18) from 3 independent experiments. Significant differences at *P* < 0.05 between flg22-treated Col-0 and *rbohF* were found between 10 and 60 min based on two-way ANOVA and uncorrected Fisher's LSD analyses for each time point.

(c) Hypothetical model of the regulation of apoplastic pH by RBOHF during PTI activation. In normal unstressed condition, the efflux of electron (e<sup>-</sup>) produced by the NADPH oxidase activity is compensated by an efflux of proton (H<sup>+</sup>) probably generated by plasma membrane H<sup>+</sup> ATPases. Both the acidification of the apoplast and the production of hydroxyl radicals (OH<sup>\*</sup>) from superoxide (O<sub>2</sub><sup>-</sup>) through the Haber–Weiss and Fenton reactions contribute to cell expansion. The plant immune response may induce the inhibition of H<sup>+</sup>ATPases and the activation of RBOHF together with superoxide dismutases (SOD) or germin-like proteins which dismutates O<sub>2</sub><sup>-</sup> to H<sub>2</sub>O<sub>2</sub>, a reaction that consumes H<sup>+</sup>. The resulting alkalisation of the apoplast induces a depolarisation of the plasma membrane ultimately leading to stomatal closure via the activation of ion channels.

### Putative role of RBOHF in PAMP-mediated apoplastic alkalisation

The *rbohF* mutant was affected in stomatal closure induced by exogenous H<sub>2</sub>O<sub>2</sub> suggesting that, besides ROS production, RBOHF also functions in other mechanisms important for stomatal immunity. Interestingly, despite a higher apoplastic pH under normal conditions, the *rbohF*

mutant was affected in flg22-induced apoplastic alkalisation. In the *rbohC* mutant, the amplitude of apoplastic pH fluctuations during root hair elongation is affected (Monshausen et al., 2007). It was shown recently that damage-induced membrane depolarisation in roots is impaired in *rbohD* and *rbohF* mutants (Marhavý et al., 2019) and RBOHD might be important in propagating the systemic



electric potential induced by wounding in leaves (Suzuki et al., 2013). These results suggest that besides superoxide production, plant NADPH oxidases like their animal counterparts (Segal, 2016) are electrogenic through electron transport across the plasma membrane which is coupled to proton efflux probably through H<sup>+</sup>-ATPases (Figure 4c). These extracellular H<sup>+</sup> ions could participate in cell wall loosening *via* the expansion activities in normal conditions (Cosgrove, 2005). During PAMP-triggered immunity, superoxide dismutation to H<sub>2</sub>O<sub>2</sub>, probably catalysed by unknown PAMP-activated superoxide dismutases or germin-like proteins (Smirnov & Arnaud, 2019), consumes protons and could increase apoplastic pH (Figure 4c) which in turn activates apoplastic peroxidases (Bolwell et al., 2002). Alternatively, *rbohF* could have an indirect effect on the activity of unknown proton transporters or H<sup>+</sup>-ATPases at the plasma membrane (Segal, 2016). It has been suggested that flg22 inhibits AHA1 and AHA2 H<sup>+</sup>-ATPase *via* dephosphorylation (Elmore & Coaker, 2011; Nuhse et al., 2007). More importantly, apoplastic alkalisation driven by RBOHF or AHA1/2 inhibition likely contributes to plasma membrane depolarisation which is well known to activate outward K<sup>+</sup> channels and S-type anion channels leading to stomatal closure (Kollist et al., 2014). The function of RBOHF, RBOHD and other NADPH oxidases in connecting extra- and intracellular ROS production, apoplastic and cytosolic pH changes and plasma membrane polarisation during plant response to environmental stresses needs to be further explored.

## EXPERIMENTAL PROCEDURES

### Statistical analysis

The experiments reported here were repeated at least three times with similar results unless otherwise mentioned. Experiments were analysed by Student's *t*-tests or ANOVAs followed by Tukey's honestly significant difference (HSD) post hoc test using R software (R Core Team, <https://www.R-project.org>). Raw data for stomatal aperture measurements are in Supplemental Table S1. The time course experiment was analysed by 2-way ANOVA. Significant differences between each treatment at each time were determined by uncorrected Fisher's LSD test using GraphPad Prism v8 (GraphPad, San Diego, California, USA).

### Plant materials and growth conditions

The roGFP2-Orp1 line (Nietzel et al., 2019) and all mutant lines are in *Arabidopsis* Col-0 background. *rbohD* (CS9555), *rbohF* (CS9557), *prx33-3* (GK-014E05) and *prx34-2* (GK-728F08), were previously described (Arnaud et al., 2017; Torres et al., 2002). The *rbohF* (SALK\_034674) and *rbohD* (SALK\_070610) lines were obtained from the European Arabidopsis Stock Centre. All T-DNA insertion mutants were confirmed by PCR genotyping prior to use (Supplemental Table S2). The genotyping of *rboh* mutants is shown in Figures S4 and S5. The obtention of roGFP2-Orp1 *rbohD*, roGFP2-Orp1 *rbohF*, roGFP2-Orp1 *prx33-3*, roGFP2-Orp1 *prx34-2* lines were previously described (Arnaud et al., 2023). Four- to 5-week-old plants grown on soil in a growth chamber under short-day conditions (10 h light at 22°C/14 h dark at 19°C), at 60% humidity and

illuminated with fluorescent tubes at 100 μmol m<sup>-2</sup> sec<sup>-1</sup> light intensity were used for all the experiments. For the experiments described below, treatments with chemicals or bacteria were performed 5–6 h after the light switch was on.

### Chemicals

Purified chemicals, except the flg22 peptide (Peptron, Korea), were purchased from Sigma. Control solutions were stomatal buffer containing 1% ethanol for 2 mM salicylhydroxamic acid (SHAM), 0.1% DMSO for 20 μM diphenyleneiodonium chloride (DPI) and water for 1 μM sodium azide, 10 μM to 1 mM hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>), 5 μM flg22.

### Bacterial infection assay

The bacterial strain *Pst* DC3000 COR<sup>-</sup> (DB4G3) (Brooks et al., 2004) was cultivated overnight at 28°C in King's B medium supplemented with Kanamycin and Rifampicin (each at 100 μg/ml). Bacteria were collected by centrifugation at 3000 *g* for 5 min at room temperature and washed twice in 10 mM MgCl<sub>2</sub>. Plants were surface-inoculated by spraying with a bacterial solution of 10<sup>8</sup> cfu/ml in 10 mM MgCl<sub>2</sub> containing 0.02% Silwet L-77, and plants were covered to maintain high humidity until disease symptoms developed. Alternatively, rosette leaves were syringe-infiltrated with a bacterial solution of 10<sup>6</sup> cfu/ml in 10 mM MgCl<sub>2</sub>. After 3 days, bacterial growth in the apoplast of three leaves per plant and six plants per genotype was determined as previously described (Katagiri et al., 2002).

### Luminol assay

Leaf discs of 6 mm diameter were cut into four equal pieces, immersed in distilled water and incubated for 3 h minimum at room temperature for recovery after wounding. Before starting the assay, water was exchanged by a solution containing 100 μM luminol and 10 μg/ml horseradish peroxidase (≥ 250 units/mg solid). After adding a control solution or 1 μM flg22, the luminescence was measured immediately using a CLARIOstar plate reader (BMG Labtech, Aylesbury, UK) with a reading time of 2 sec.

### Confocal imaging and image analysis

Leaf discs from rosette leaves were immersed in 10 mM MES-KOH pH 6.15, 30 mM KCl, incubated for 2 h at 21°C under laboratory lighting (PPFD ~ 10 μmol m<sup>-2</sup> sec<sup>-1</sup>) for recovery after wounding and subsequently treated with control solution or 1 μM flg22 for 30 and 60 min. Leaf discs were mounted under a Zeiss confocal microscope LSM 880. Images were collected with a 20X lens (Plan-Apochromat, 0.8 numerical aperture) and roGFP2-Orp1 was excited sequentially at 405 and 488 nm and emission was detected at 508–526 nm, with the pinhole set to 1 airy unit. Single-plane images were processed with ImageJ software. Background fluorescence was insignificant and not subtracted. Fluorescence intensity values of individual guard cells selected as region of interest were quantified on 32-bit converted images and the ratio 405/488 nm was calculated.

### Measurements of stomatal aperture

Stomatal experiments were conducted as previously described (Desclos-Theveniau et al., 2012). Epidermal peels collected from the abaxial side of young fully expanded leaves or leaf discs were floated in stomatal buffer (10 mM MES-KOH pH 6.15, 30 mM KCl) for 2.5 h under light (100 μmol m<sup>-2</sup> sec<sup>-1</sup>) to ensure that most stomata were opened before treatments. Solutions of chemicals or bacterial suspension at 10<sup>8</sup> cfu.ml<sup>-1</sup> in 10 mM MgCl<sub>2</sub> were directly added to the stomatal buffer. After treatments, epidermal

peels or leaf discs were further incubated under light for 2 h and observed under a light microscope (Carl Zeiss, Axioplan 2). Stomatal apertures of stomata in random areas were measured using ImageJ 1.42 software. Measurements of stomatal aperture were performed 7–8 h after starting illumination.

### Monitoring ROS in guard cells

2',7'-dichlorofluorescein diacetate (H<sub>2</sub>DCFDA, Sigma) was used to measure ROS in guard cells (Pei et al., 2000). After 2.5 h incubation in stomatal buffer, epidermal peels were incubated with 50 μM H<sub>2</sub>DCFDA in 0.1% DMSO for 15 min. Excess H<sub>2</sub>DCFDA was then removed by washing 3 times for 20 min with stomatal buffer. Then, 5 μM flg22 or control solution was added. After 30 min incubation, H<sub>2</sub>DCFDA fluorescence was observed with a fluorescence microscope (Carl Zeiss, Axioplan 2) and fluorescence intensity of guard cells was analysed using ImageJ software.

### Gene expression analysis in guard cell protoplasts

For each condition, about 50 young fully expanded leaves were immersed for 2 h in stomatal buffer containing 0.02% (v/v) Silwet-L77 with or without 1 μM flg22. Guard cell protoplasts were isolated as previously described (Obulareddy et al., 2013) in the presence of transcriptional inhibitors 0.01% (w/v) cordycepin and 0.0033% (w/v) actinomycin D. For each condition, about 10<sup>6</sup> guard cell protoplasts were obtained and their purity was above 98%. RNA was extracted using the RNeasy Plant Mini Kit with in-column Dnase 1 digestion (Qiagen, Hilden, Germany). Two hundred nanograms of total RNA were reverse transcribed using 500 ng of oligo(dT)15 and the ImProm-II Reverse Transcription System following the manufacturer's instructions (Promega, Madison, WI, USA). Quantitative real-time PCR reaction was performed on a Light Cycler 2 (Roche, Basel, Switzerland) using 10 μl SYBR Premix Ex Taq (Takara, Kusatsu, Japan), 2 μl of twofold diluted cDNA and 0.5 μM of primers in a total volume of 20 μl per reaction. The cycling conditions were composed of an initial 20 sec denaturation step at 95°C, followed by 45 cycles of 95°C for 7 sec, 60°C for 10 sec and 72°C for 13 sec. A melting curve was run from 65°C to 95°C to ensure the specificity of the products. Data were analysed with the delta Ct method. Ubiquitin 1 (*UBQ1*) was used as a reference gene for normalisation of gene expression levels. Control treatment was considered as expression level = 1. As controls, the expression of the reference gene *ACT2* did not change between control and flg22 conditions while the expression of the PTI marker gene *PRX4* was induced by flg22 (Figure S2d). qRT-PCR primer sequences are listed in Supplemental Table S2.

### Ratiometric pH quantification using Oregon green

Fully expanded leaves were syringe-infiltrated with 25 μM of Oregon Green 488 dextran (ThermoFisher, Waltham, MA, USA) (Geilfus & Muehling, 2011; McLachlan et al., 2016). For the autofluorescence background, milliQ water was infiltrated into the apoplast. Plants were kept in the growth chamber for 2 h until excess water has evaporated. Oregon Green-treated leaf discs were placed in a 96-well plate, immersed in milliQ water and further incubated for 2 h at 21°C under laboratory lighting (PPFD ~ 10 μmol m<sup>-2</sup> sec<sup>-1</sup>) for recovery after wounding. Oregon Green was excited sequentially at 440 nm ± 8 nm and 495 nm ± 8 nm in a CLARIOstar plate reader (BMG Labtech) and emission was recorded at 525 nm ± 20 nm with a gain set at 1250 and 1000 for the 440 and 495 nm excitations, respectively. Each leaf disc was scanned from the top with the fluorescence recorded and averaged from 76 flashes per well-organised as a spiral of 5 mm diameter. The initial 495/440 ratio of the resting state of leaf discs was estimated by reading the wells for 15 min

before treatment. The emission of water-infiltrated leaf discs was averaged and subtracted for all the data points to correct for background fluorescence. To convert the fluorescence 495/440 ratio into pH values, a calibration curve with pH-buffered Oregon Green solution infiltrated into the apoplast of leaves was performed according to (McLachlan et al., 2016). The Boltzmann fit was chosen for fitting sigmoidal curves to calibration data and pH was determined according to the equation:

$$x = V50 - Slope \ln \left( \frac{Top - y}{y - Bottom} \right)$$

where  $x = \text{pH}$ ,  $y = 495/440$  ratio,  $Bottom = 0.4833$ ,  $Top = 2.045$ ,  $V50 = 4.513$  and  $Slope = 0.6267$ . Changes in pH over time were normalised to the initial pH<sub>i</sub> as pH/pH<sub>i</sub>.

### ACCESSION NUMBERS

The *A. thaliana* genes included in this study are as follows: *PRX33* (At3g49110); *PRX34* (At3g49120); *RDOHD* (At5g47910); and *RBOHF* (At1g64060).

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### AUTHOR CONTRIBUTIONS

DA designed and performed the research, analysed the data and wrote the paper. MJD designed the research, analysed the data and wrote the paper. NS designed the research, analysed the data and wrote the paper.

### CONFLICT OF INTEREST

The authors declare no conflict of interest.

### SUPPORTING INFORMATION

Additional Supporting Information may be found in the online version of this article.

**Figure S1.** Flg22-triggered roGFP2-Orp1 oxidation in pavement cells and guard cells.

**Figure S2.** The NADPH oxidase RBOHF activates stomatal immunity.

**Figure S3.** The *rbohF* mutant is partly defective in ABA- and H<sub>2</sub>O<sub>2</sub>-mediated stomatal closure.

**Figure S4.** Genotyping by PCR of *rbohD* and *rbohF* dSpm transposon mutants.

**Figure S5.** Genotyping by PCR of *rbohD* and *rbohF* Salk T-DNA mutants.

**Table S1.** Raw data of stomatal aperture measurements (in μm) involving the *rbohD* and *rbohF* mutants.

**Table S2.** PCR primers used for genotyping and RT-qPCR.

### OPEN RESEARCH BADGES



This article has earned an Open Data badge for making publicly available the digitally-shareable data necessary to reproduce the reported results. The data is available.

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