

## RESEARCH ARTICLE

# Ethylene signaling increases reactive oxygen species accumulation to drive root hair initiation in *Arabidopsis*

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## ABSTRACT

Root hair initiation is a highly regulated aspect of root development. The plant hormone ethylene and its precursor, 1-amino-cyclopropane-1-carboxylic acid, induce formation and elongation of root hairs. Using confocal microscopy paired with redox biosensors and dyes, we demonstrated that treatments that elevate ethylene levels lead to increased hydrogen peroxide accumulation in hair cells prior to root hair formation. In the ethylene-insensitive receptor mutant, *etr1-3*, and the signaling double mutant, *ein3eil1*, the increase in root hair number or reactive oxygen species (ROS) accumulation after ACC and ethylene treatment was lost. Conversely, *etr1-7*, a constitutive ethylene signaling receptor mutant, has increased root hair formation and ROS accumulation, similar to ethylene-treated Col-0 seedlings. The *caprice* and *wereewolf* transcription factor mutants have decreased and elevated ROS levels, respectively, which are correlated with levels of root hair initiation. The *rhd2-6* mutant, with a defect in the gene encoding the ROS-synthesizing RESPIRATORY BURST OXIDASE HOMOLOG C (RBOHC), and the *prx44-2* mutant, which is defective in a class III peroxidase, showed impaired ethylene-dependent ROS synthesis and root hair formation via EIN3EIL1-dependent transcriptional regulation. Together, these results indicate that ethylene increases ROS accumulation through RBOHC and PRX44 to drive root hair formation.

**KEY WORDS:** Ethylene, Reactive oxygen species, Root hairs, Peroxidase, RBOH

## INTRODUCTION

The initiation of root hairs is genetically programmed and environmentally sensitive, making root hairs an ideal model for studying single-cell differentiation in plants. Root hairs are single-cell extensions that differentiate from longitudinal epidermal cell files, known as trichoblasts (Leavitt, 1904; Salazar-Henao et al., 2016). The formation of root hairs is modulated by environmental changes to increase root surface area to allow for efficient water and nutrient uptake (Bruex et al., 2012), while also anchoring plants in soil to reduce erosion (De Baets et al., 2020). In *Arabidopsis*, the root epidermis consists of an alternating pattern of trichoblasts,

which form root hairs, and atrichoblasts, which are non-hair-forming cells (Pemberton et al., 2001; Salazar-Henao et al., 2016). Root hair formation is dictated by cell positioning; epidermal cells overlying two cortical cells can become hair-forming cells, whereas those overlying one cortical cell will become non-hair cells (Berger et al., 1998; Salazar-Henao et al., 2016). Root hair development is separated into two processes: root hair initiation and root hair elongation (Duckett et al., 1994). A recent report divided this process into ten precise stages and examined the molecular mechanisms that drive this process. During stages -7 to -1, RHO OF PLANTS (ROP) proteins (ROP2, ROP4 and ROP6) and ROP guanine nucleotide exchange factors (ROP GEFs), which regulate ROP activity, accumulate at the future site of initiation (Denninger et al., 2019; Molendijk et al., 2001). This is followed by polymerization of actin filaments along which vesicles move to deposit the membrane needed for polarized tip growth (Salazar-Henao et al., 2016). During tip growth, tip-focused reactive oxygen species (ROS) (Foreman et al., 2003; Gayomba and Muday, 2020; Monshausen et al., 2007) and Ca<sup>2+</sup> gradients (Carroll et al., 1998) have been shown to promote exocytosis of the cell wall and membrane materials, driving subsequent root hair elongation. These processes are separable as mutants with either impaired root hair initiation or elongation have been identified (Masucci and Schiefelbein, 1994; Schiefelbein and Somerville, 1990).


Genetic screens in *Arabidopsis thaliana* have provided a wealth of insights into the proteins that drive root hair development (Lee and Schiefelbein, 1999; Masucci and Schiefelbein, 1996; Rerie et al., 1994). For example, the mutants *transparent testa glabra* (*tig*), *glabra2* (*gl2*) and *wereewolf* (*wer*) have root hairs that form from both trichoblast and atrichoblast cells (Di Cristina et al., 1996; Galway et al., 1994; Lee and Schiefelbein, 1999). Many of the protein products of these mutants have been mapped to transcriptional cascades that drive root hair differentiation (Shibata and Sugimoto, 2019) and have been shown to localize to root hair cells prior to root hair initiation (Denninger et al., 2019). In non-hair cells, a transcriptional complex containing three transcription factors (TFs), WER, GLABRA 3 (GL3) or its functionally redundant ENHANCER OF GLABRA (EGL3), and TTG1, functions as a transcriptional activator of the GL2 protein, which leads to repression of root hair initiation (Grebe, 2012; Salazar-Henao et al., 2016). In trichoblasts, WER expression is repressed, which allows for the formation of an alternate transcriptional complex comprising CAPRICE (CPC) or the functionally redundant proteins ENHANCER OF TRY AND CPC1 (ETC1), ETC3 or TRYPTICHON (TRY) (Schiefelbein et al., 2014). When this pathway is active, GL2 is not expressed and root hair initiation proceeds (Salazar-Henao et al., 2016).

Another genetic screen identified the *root hair defective* (*rhd*) mutants, which have impaired root hair initiation, elongation or structure (Masucci and Schiefelbein, 1994; Schiefelbein and Somerville, 1990). The mutation in *rhd2* was mapped to the RBOHC (RESPIRATORY BURST OXIDASE HOMOLOG

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*C/NADPH OXIDASE*) gene (Foreman et al., 2003). The *rhod2* mutant was identified in a screen for altered root hair elongation (Schieffelin and Somerville, 1990), but was recently reported to also have impaired root hair initiation (Gayomba and Muday, 2020). RBOHs are integral plasma membrane proteins that produce superoxide, which can be dismutated to hydrogen peroxide via superoxide dismutase (SOD) or other non-enzymatic mechanisms (Chapman et al., 2019).  $H_2O_2$  can then enter into the cell through aquaporins (Bienert et al., 2007), where it can act as a signaling molecule to drive cellular processes. Signaling-induced ROS regulates protein function by reversibly oxidizing cysteine residues to sulfenic acids (Cys-SOH) (Poole and Schoneich, 2015).

There are ten RBOH family members (RBOHA-RBOHJ) in *Arabidopsis* and each play distinct roles in organ development and stress response (Chapman et al., 2019). RBOHs can be regulated transcriptionally or enzymatically by a number of mechanisms, including calcium binding, phosphorylation and phosphatidic acid binding (Kobayashi et al., 2007; Postiglione and Muday, 2020; Suzuki et al., 2011). RBOH-induced ROS production is also regulated by hormone signaling, as many plant hormones generate ROS as a mechanism to drive growth and developmental processes (Chapman et al., 2019; Kwak et al., 2003; Mittler et al., 2011; Postiglione and Muday, 2020). For example, abscisic acid (ABA), a hormone involved in the abiotic stress response, has been shown to induce RBOH-derived ROS production to prevent water loss in leaves (Kwak et al., 2003). Auxin induces root hair initiation (Gayomba and Muday, 2020) and elongation (Mangano et al., 2017) through a localized increase in ROS, suggesting that RBOH may drive hormone-induced root hair elongation.

The plant hormone ethylene enhances root hair initiation and elongation. Treatment with ethylene, or its precursor 1-aminocyclopropane-1-carboxylic acid (ACC), leads to proliferation of root hairs, with substantial increases in their length (Tanimoto et al., 1995). The ethylene induction of root hairs occurs through the canonical ethylene signaling pathway, which is initiated when ethylene binds to one of the five receptors ETR1, ERS1, ETR2, ERS2 or EIN4 (Binder, 2020; Bleecker, 1999). When ethylene is absent, the receptors are in the 'on' state, leading to activation of the CTR1 Raf-like kinase (Kieber et al., 1993); this turns the pathway off via phosphorylation and subsequent degradation of the EIN2 transmembrane protein (Alonso et al., 1999; Ju et al., 2012). When ethylene binds to its receptors they shift to the 'off' state, which prevents activation of CTR1. This allows for cleavage and translocation of the EIN2 C terminus into the nucleus (Ju et al., 2012; Wen et al., 2012) where it stabilizes the EIN3/EIL1 TFs leading to ethylene-responsive gene expression. Previous work has shown that ethylene regulates root hair elongation through the EIN3 transcription factor (Feng et al., 2017) and that EIN3 also controls the formation of ectopic root hairs when ethylene reaches very high levels (Qiu et al., 2021). EIN3 physically interacts with RHD6, a positive regulator of root hair development, to form a transcriptional complex that binds to and induces expression of *RSL4*, resulting in increased root hair elongation (Feng et al., 2017). However, the mechanistic events that drive ethylene-induced root hair initiation have not been fully described and the role of ROS as a downstream molecule in ethylene-induced root hair development has not been reported.

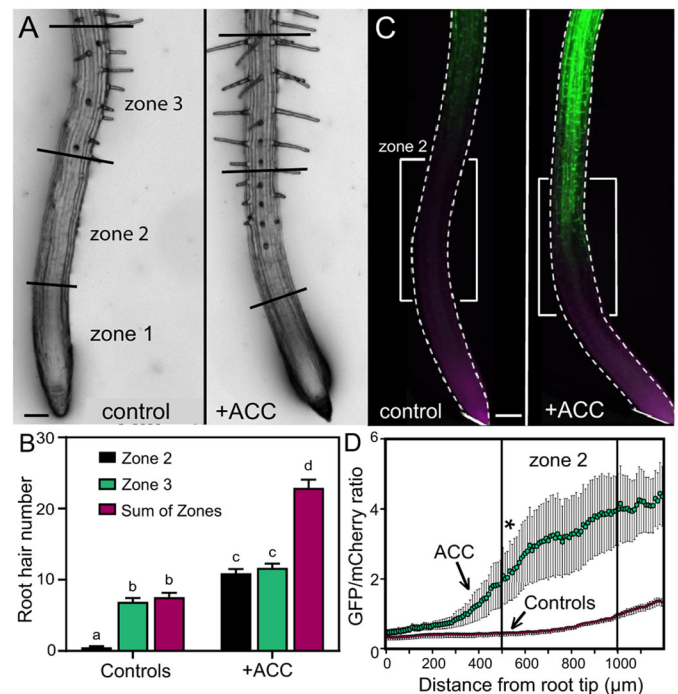
Here, we investigated whether ethylene acts to increase ROS levels to drive root hair initiation. Fluorescent dyes and biosensors that report ROS levels were used to examine ROS accumulation after ethylene or ACC treatment in trichoblast cells in the differentiation zone. This response was shown to be dependent on ETR1 receptor activity and EIN3 and/or EIL1 TF activity. The role of RBOHC and

class III peroxidase enzymes in ethylene-dependent ROS synthesis was demonstrated using mutants in genes encoding these ROS-generating enzymes. Together, these experiments demonstrated that ROS is a signaling molecule in ethylene-induced root hair formation and identified several enzymes that participate in producing ethylene-induced ROS.

## RESULTS

### Root hair number increases in ACC-treated roots

The role of ethylene signaling in root hair initiation was examined by testing the effects of short-term treatments with the ethylene precursor ACC on the number and position of root hairs in 5-day-old seedlings. Root hairs were visualized in wild-type (Col-0) seedlings grown in the presence of  $0.7 \mu\text{M}$  ACC for 4 h (Fig. 1A). The root tip was divided into three 500- $\mu\text{m}$  zones starting from the root tip (Fig. 1A) and the number of initiated root hairs formed in each zone was quantified (Fig. 1B). We defined root hair initiation as root hairs that were at stage +2 and above as described previously (Denninger et al., 2019). In zone 1, root hairs did not form in either untreated or ACC-treated seedlings. In zone 2, there were



**Fig. 1. ACC treatment increases root hair number and ROS-dependent gene expression along the root after 4 h.** (A) Representative images of root tips of Col-0 with and without ACC treatment for 4 h. Zones 1, 2 and 3 represent 500  $\mu\text{m}$  root sections. (B) Root hair quantification of untreated and 4 h ACC-treated seedlings. No root hairs formed in zone 1. Data are mean $\pm$ s.e.m. of total root hairs in zones 2 and 3, and all zones from three experiments ( $n=18-24$  seedlings per experiment). Columns with different letters indicate statistically significant differences ( $P<0.0001$ ) determined by two-way ANOVA followed by Tukey's multiple comparisons test. (C) Representative images of root tips of Col-0 containing the ZAT12p-ROS reporter with or without ACC treatment for 4 h with GFP signal in green and mCherry signal in magenta. White brackets indicate zone 2 of the root. (D) Quantification of the ratio of GFP/mCherry fluorescence intensity using a line profile along the roots of an average of 6-12 seedlings of three independent experiments of untreated or ACC-treated seedlings. Data are mean $\pm$ s.e.m. The asterisk at  $\sim 510 \mu\text{m}$  from the root tip represents the shortest distance from the root tip at which the values for GFP fluorescence in the presence of ACC became significantly different ( $P<0.0001$ ) from the controls as determined by unpaired two-tailed Student's *t*-test. Scale bars: 100  $\mu\text{m}$ .

very few root hairs in untreated roots; however, ACC treatment increased the number of root hairs by tenfold. There was also a twofold increase in root hair number in zone 3 of ACC-treated seedlings compared with untreated controls. This dose of ACC did not cause ectopic root hair formation in non-hair cells, but rather increased the number of hair cells forming root hairs in these two zones. It is also evident from Fig. 1 that this treatment increased the length of root hairs in both zones 2 and 3, consistent with prior reports (Feng et al., 2017; Harkey et al., 2018). These data suggest that ACC-induced root hair initiation begins between 500 and 1000  $\mu\text{m}$  from the root tip. Therefore, to understand the mechanisms driving the process of root hair initiation, our experiments focused on this region.

Long-term ACC and ethylene treatment also result in a shorter primary root due to reduced elongation of root cells (Harkey et al., 2018), so we examined whether the effect on root hair initiation was the result of altered length. The effect of 4 h of treatment with low doses of ACC (0.7  $\mu\text{M}$ ) on root elongation were examined. Primary roots were stained with the cell wall-specific dye propidium iodide (PI). We then measured the length of five epidermal cells from six to eight roots per treatment condition, measuring the length of cells at either end and in the middle of zone 2 (designated zone 2A, 2B and 2C; Fig. S1). There was no difference in cell length in any of these three regions after 4 h of ACC treatment compared with untreated controls. These results are consistent with short-term and low-dose ACC treatments increasing root hair number by inducing root hair formation from trichoblasts in zone 2, rather than as an indirect effect of a shorter primary root.

#### A ROS-dependent transcriptional reporter increases in ACC-treated roots

We investigated whether ethylene leads to elevated ROS to drive ethylene-induced root hair initiation. ROS-dependent gene expression along the root was examined using the ZAT12p-ROS ratiometric biosensor (Lim et al., 2019). This reporter construct contains the promoter of the ROS-sensitive transcription factor ZAT12, driving GFP, and the constitutive ubiquitin10 promoter, driving mCherry. We compared the fluorescent signal of ZAT12p-GFP in the presence and absence of ACC as visualized by laser-scanning confocal microscopy (LSCM), with GFP reported as green and mCherry reported as magenta (Fig. 1C). The ratio of signal of GFP (green) to mCherry was quantified across the entire root as a distance from the root tip. In ACC-treated roots, the GFP/mCherry ratio increased beginning at approximately 500  $\mu\text{m}$  from the root tip, which corresponded to zone 2, where root hair induction was maximal upon treatment with ACC (Fig. 1D). To examine the possibility that the ZAT12 promoter is ethylene regulated (rather than ROS regulated), we examined transcriptomic datasets in which roots were treated with ACC or ethylene (Harkey et al., 2018). These three datasets did not show ACC- or ethylene-driven transcriptional changes in ZAT12 transcript abundance.

#### Ethylene increases the signal of the H<sub>2</sub>O<sub>2</sub>-selective dye Peroxy Orange1 in trichoblast cells

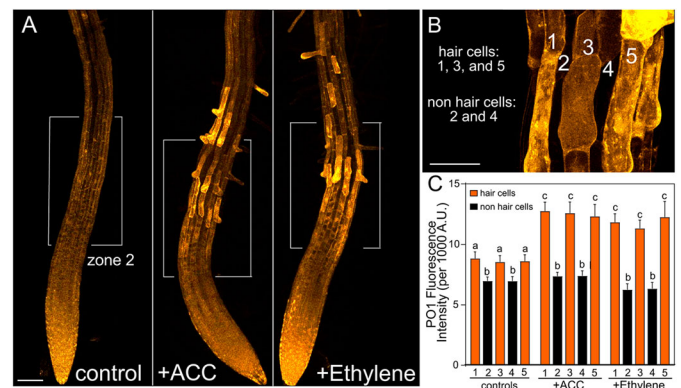
The hydrogen peroxide-selective dye Peroxy-Orange 1 (PO1) was used to determine whether there were cell type-specific increases in ROS in response to treatment with ACC or ethylene gas. PO1 is a permeable, boronate-based dye that is non-fluorescent in its reduced form, but becomes fluorescent when irreversibly oxidized by H<sub>2</sub>O<sub>2</sub> (Dickinson et al., 2010). Col-0 seedlings were treated with either 0.7  $\mu\text{M}$  ACC or 0.05 ppm of ethylene gas for 4 h followed by PO1

staining. In Col-0 roots, PO1 fluorescence was visualized using LSCM in zone 2 of roots treated with control, ACC or ethylene gas for 4 h (Fig. 2). In *Arabidopsis*, root hairs form in alternating patterns, so that every root hair-forming cell (trichoblast) is adjacent to a non-hair cell (atrichoblast). We quantified ROS accumulation by analyzing PO1 signal after confocal imaging, by drawing a line across the width of the root that spanned five epidermal cell files so that the PO1 signal in three trichoblasts (numbered 1, 3 and 5) and two atrichoblasts (numbered 2 and 4) could be quantified (Fig. 2A).

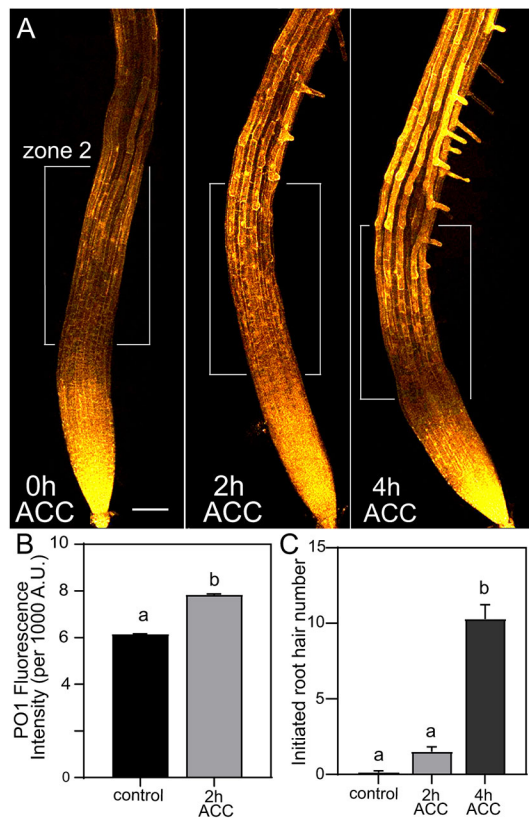
We found that PO1 fluorescence in trichoblasts in zone 2 of untreated seedlings was slightly, but not significantly, higher than in atrichoblasts. In trichoblast cells of ACC- and ethylene-treated seedlings, there was a significant increase in PO1 fluorescence compared with trichoblasts of untreated seedlings. In contrast, there were no changes in PO1 fluorescence in the atrichoblasts, suggesting that ethylene and ACC treatment increased ROS levels in only the cells that formed root hairs. PO1 fluorescence intensity values in trichoblasts after ethylene treatment showed the same magnitude increase as roots treated with ACC (Fig. 2C). These results are consistent with this short-term and low-dose treatment with ACC leading to efficient conversion to ethylene, which in turn produces elevated ROS in trichoblast cells (Fig. 2B). This is in contrast with some studies in which other developmental processes are altered by ACC acting directly, rather than conversion to ethylene (Li et al., 2022).

#### ROS accumulation increases prior to root hair emergence

To determine whether ethylene-induced ROS accumulation drives root hair emergence, we investigated whether ACC-induced ROS increases were detectable prior to the first ACC-induced root hair initiation. Wild-type Col-0 seedlings were treated with ACC for either 2 or 4 h and PO1 fluorescence was visualized in trichoblasts that did not have a root hair bulge (stages below +1) (Fig. 3A). This PO1 signal is only reported in the 2-h ACC treatment, as there was an insufficient number of cells without root hair bulges to quantify in



**Fig. 2. ACC and ethylene gas increase H<sub>2</sub>O<sub>2</sub> accumulation in trichoblasts after 4 h.** (A) Representative roots illustrating the alternating PO1 fluorescence between trichoblast and atrichoblast epidermal cell files of cells in Col-0 roots with and without ACC and ethylene gas treatment for 4 h. White brackets indicate zone 2 of the root. Scale bar: 100  $\mu\text{m}$ . (B) Examples of hair cells (1, 3, 5) and non-hair cells (2, 4) from roots treated with ethylene. Scale bar: 50  $\mu\text{m}$ . (C) Quantification of PO1 fluorescence intensity in trichoblasts (orange bars: cells 1, 3 and 5) and atrichoblasts (black bars: cells 2, 4). Data are mean  $\pm$  s.e.m. of three independent experiments ( $n=18-24$  seedlings/experiment). Columns with different letters indicate statistical significance ( $P<0.0001$ ) compared with Col-0 untreated cells as determined by two-way ANOVA followed by Tukey's multiple comparisons test. A.U., arbitrary units.

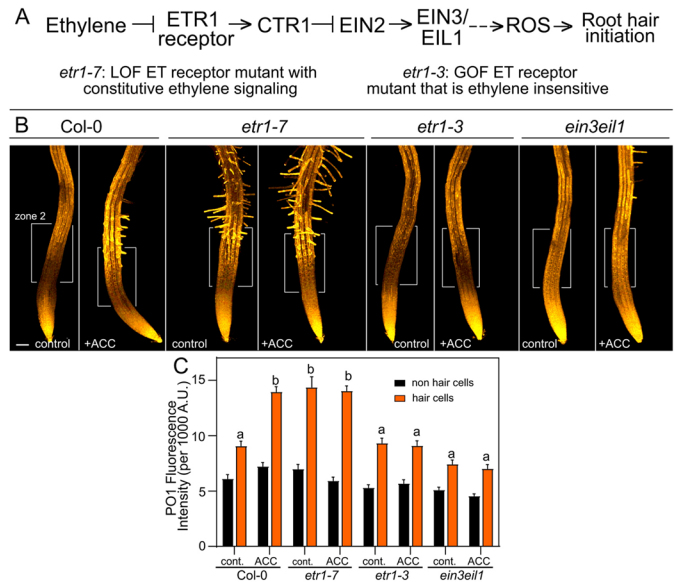


**Fig. 3. ACC-increased ROS accumulation precedes root hair initiation.** (A) Representative images of epidermal PO1 fluorescence in untreated Col-0 and Col-0 treated with ACC for 2 or 4 h. White brackets indicate zone 2 of the root. Scale bar: 100  $\mu$ m. (B) Quantification of trichoblasts that had not yet formed root hair bulges of control and 2 h ACC-treated seedlings. (C) Quantification of the number of initiated root hairs in zone 2 of the root (500  $\mu$ m–1000  $\mu$ m from root tip) in untreated and 2 and 4 h ACC-treated seedlings. Data are mean+s.e.m. of individual cells from three independent experiments ( $n=12$ –18 seedlings/experiment). Columns with different letters indicate statistical significance compared with untreated hair cells as determined by unpaired two-tailed Student's *t*-test ( $P<0.0001$ ) (B) or one-way ANOVA ( $P<0.0001$ ) followed by Tukey's multiple comparisons test (C). A.U., arbitrary units.

roots treated for 4 h. Total PO1 accumulation was measured in five individual trichoblasts per root with and without ACC treatment. The signal was quantified across the area of the entire cell and the average PO1 intensity of 24–30 individual cells was reported (Fig. 3B). Atrichoblast signal was not quantified as ROS levels did not change in those cells (Fig. 2C). These data showed a 1.3-fold increase in PO1 accumulation in hair cells of seedlings treated with ACC for 2 h compared with untreated controls, when trichoblast cells that had not yet begun to initiate root hairs were examined. These data are consistent with the hypothesis that ROS acts as a driver of root hair initiation downstream of ethylene signaling.

### Ethylene signaling mutants show altered ROS accumulation patterns and root hair phenotypes

The ethylene signaling pathway is well defined (Fig. 4A) and mutants in key signaling proteins, including receptors and transcription factors, are available. ROS accumulation was examined in response to ACC treatment in these ethylene signaling mutants to determine whether this response was dependent on the ethylene signaling pathway and downstream transcriptional responses. The number of root hairs and average length have been previously reported in the



**Fig. 4. The ETR1 receptor and EIN3/EIL1 transcription factors are required for ethylene-induced ROS accumulation and root hair proliferation.** (A) Schematic of the ethylene signaling pathway and explanation of the character of mutants in the *ETR1* (ethylene resistant) gene. ETR1 is the ethylene receptor controlling root hair formation, CTR1 (CONSTITUTIVE TRIPLE RESPONSE 1) is a kinase, EIN2 (ETHYLENE INSENSITIVE 2) is a signaling protein, and EIN3 and EIL1 (EIN3-like 1) are transcription factors. GOF, gain of function; LOF, loss of function. (B) Representative images of PO1 epidermal fluorescence in Col-0, *etr1-7*, *etr1-3* and *ein3-eil1-1* with and without ACC treatment for 4 h; 18–20 seedlings from each genotype and treatment were imaged. White brackets indicate zone 2 of the root. Scale bar: 100  $\mu$ m. (C) Quantification of PO1 fluorescence intensity in hair cells and non-hair cells. For each root, PO1 fluorescence was quantified in two hair cells and two non-hair cells and the values were pooled to obtain this average. Data are mean+s.e.m. of three experiments ( $n=18$ –20 seedlings/experiment). Columns with different letters indicate statistical significance ( $P<0.0001$ ) in PO1 signal compared with other hair cells as determined by two-way ANOVA followed by Tukey's multiple comparisons test. A.U., arbitrary units.

loss-of-function and gain-of-function ethylene receptor mutants *etr1-7* and *etr1-3*, and the transcription factor mutant *ein3eil1* in the presence and absence of ACC treatment (Harkey et al., 2018). *etr1-7* is a LOF mutant in which the ETR1 receptor is inactive, therefore the ethylene signaling pathway is constitutively active and there are increased numbers of root hairs independent of ACC addition, whereas *etr1-3* is a gain-of-function mutant in which the ETR1 receptor is always active leading to the ethylene signaling pathway being inhibited (Harkey et al., 2018; Hua and Meyerowitz, 1998). The double mutant *ein3eil1* has mutations in genes encoding EIN3 and EIL1 TFs (Chao et al., 1997; Solano et al., 1998). Both *etr1-3* and *ein3eil1* have reduced root hair initiation in response to ACC treatment (Harkey et al., 2018). We examined the PO1 distribution patterns in root hair cells with and without ACC treatment in these three mutants.

We visualized PO1 fluorescence by LSCM and saw that the constitutive ethylene signaling mutant *etr1-7* had increased  $H_2O_2$  in root hair cells and an increased number of trichoblasts with emerged root hairs regardless of ACC treatment (Fig. 4B). The PO1 signal in *etr1-7* in the absence of ACC was significantly elevated over untreated Col-0, but was equivalent to ACC-treated Col-0, and was not significantly changed by ACC treatment (Fig. 4C). The opposite response was seen in the ethylene-insensitive *etr1-3* and *ein3eil1* mutants, as they showed no change in PO1 signal in response to ACC treatment, resulting in a significantly lower level than ACC-

treated Col-0 and no induction of root hair initiation (Fig. 4C). Together, these results suggest that the ETR1 receptor and EIN3/EIL1 TFs are required for ACC-induced ROS accumulation and root hair initiation and suggest that, in these experiments, ACC is acting via conversion to ethylene.

### Mutants with altered root hair formation also show altered ROS accumulation

We also examined the effect of mutations that alter root hair formation on ROS levels. The *werewolf* (*wer*) mutant has increased root hair formation, whereas the *caprice* (*cpc*) mutant exhibits decreased root hair formation. ROS accumulation was significantly higher in root hairs along the root in *wer* compared with both Col-0 and *cpc*, whereas *cpc* had significantly less ROS accumulation along the root compared with Col-0 (Fig. 5).

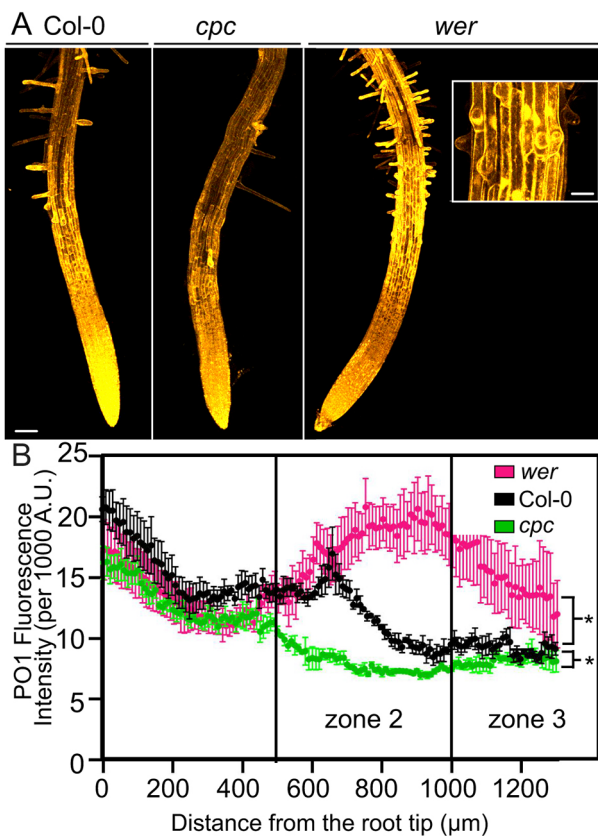
### The RBOHC knockout mutant *rhd2-6* shows decreased ROS accumulation after ACC treatment

It has been previously reported that RBOHC is involved in ROS production and subsequent root hair elongation (Foreman et al., 2003). We have also shown that there is decreased ROS in root hairs of the *rhd2-6* mutant, which has an insertion mutation in

the *RBOHC* gene (Gayomba and Muday, 2020). Therefore, we investigated whether RBOHC is required for ACC-induced ROS increases and root hair initiation. We examined root hair numbers by light microscopy in ACC-treated *rhd2-6* (*rbohc*) seedlings (Fig. 6). Root hair number in the ACC-treated *rhd2-6* mutant was significantly lower than in ACC-treated Col-0, suggesting that RBOHC contributes to ethylene-induced root hair initiation (Fig. 6B). We also quantified the length of all root hairs in zone 2 using the segmented line tool in ImageJ. A histogram showing the distribution of lengths of root hairs (reported as percentage in each length bin out of total number of roots) revealed that the length of root hairs increased after ACC treatment and that the *rhd2-6* mutant has reduced root hair elongation in the presence of ACC relative to Col-0 (Fig. 6C).

We also visualized the PO1 fluorescence by LSCM in the *rhd2-6* mutant with and without ACC treatment (Fig. 6D). We report the average of two trichoblast cells (1 and 5 from Fig. 2) and two atrichoblast cells (2 and 4) in Fig. 6E. This reveals a significant increase in H<sub>2</sub>O<sub>2</sub> fluorescence accumulation in Col-0 hair cells, whereas in *rhd2-6* there was no significant increase after ACC treatment (Fig. 6E), suggesting that RBOHC contributes to ethylene-induced ROS accumulation. The data is reported for all five cell files in Fig. S2. These images revealed slight increases in the numbers of root hairs in ACC-treated *rhd2-6* roots, suggesting that ACC also acts to induce root hairs in an RBOHC-independent mechanism.

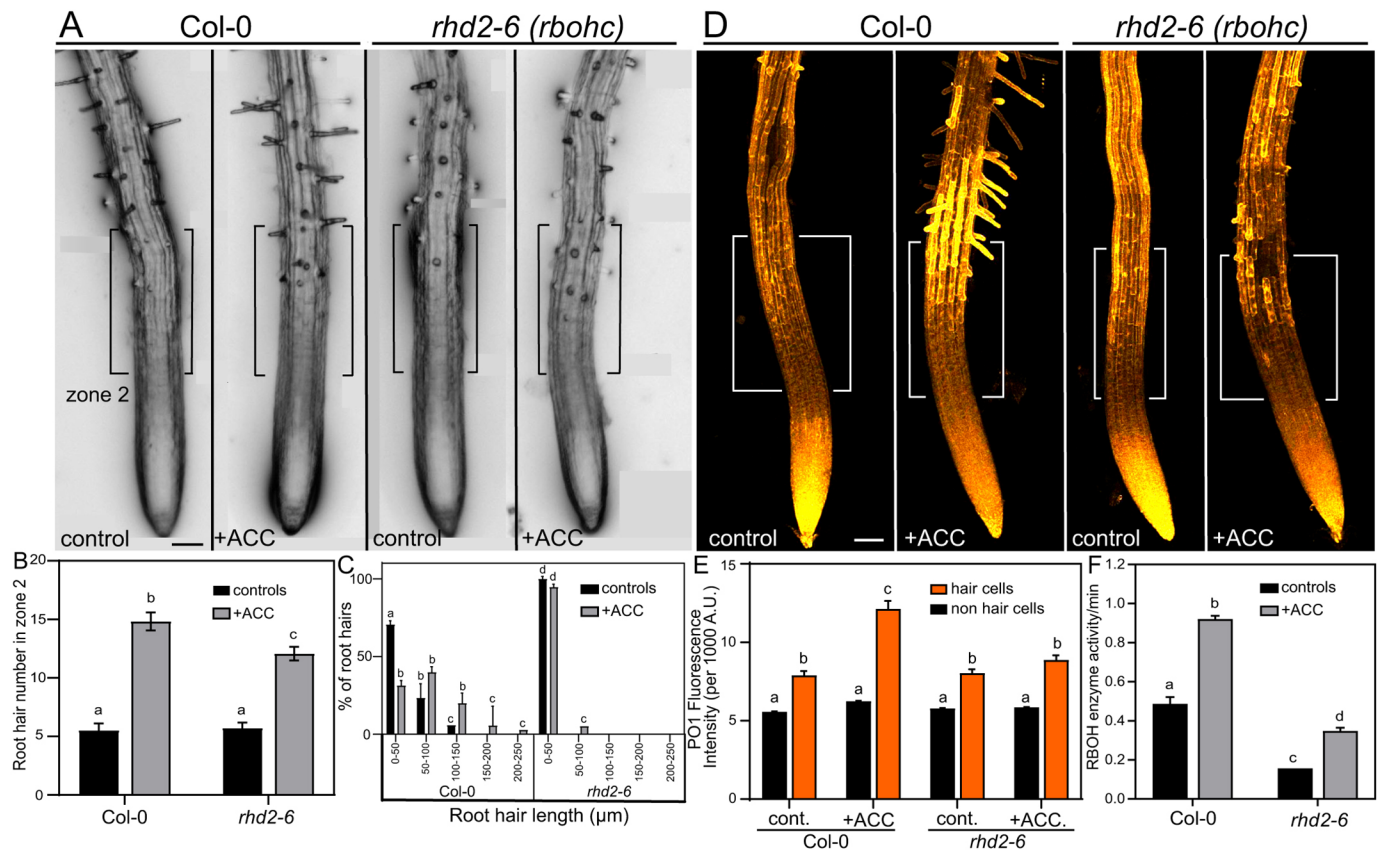
As there are also other RBOH enzymes expressed in roots, we examined whether these other enzymes affect root hair formation. RBOHD and RBOHF are highly expressed in roots and have been shown to be involved in regulation of root elongation and lateral root development (Chapman et al., 2019). Therefore, to determine whether these other root-expressed RBOHs contribute to ethylene-induced ROS accumulation, PO1 accumulation patterns were examined in the *rbohdf* double mutant and compared with *rhd2-6*. No change was seen in ROS levels in hair cells of *rbohdf*, suggesting that these RBOHs do not contribute to ethylene-induced ROS synthesis in root hair cells (Fig. S2).



**Fig. 5. ROS accumulation increases in *werewolf* and decreases in *caprice* compared with Col-0.** (A) Representative images of 5-day-old untreated Col-0, *cpc* and *wer* roots stained with PO1. Inset shows a high-magnification image of a *wer* root to illustrate ectopic root hair formation. Scale bars: 100  $\mu$ m (main panels); 25  $\mu$ m (inset). (B) Quantification of total PO1 accumulation along the root starting approximately 200  $\mu$ m from the root tip and ending at 1300  $\mu$ m. This region encompasses both zones 2 and 3. Data are mean  $\pm$  s.e.m. and are representative of two independent experiments (9–15 seedlings per experiment). Asterisks ( $P < 0.0001$ ) indicate statistically significant differences from Col-0 as determined by one-way ANOVA. A.U., arbitrary units.

### RBOHC transcript abundance and enzyme activity increases in response to ethylene signaling

Elevated ethylene may either increase *RBOHC* transcript abundance or enzyme activity. We examined several transcriptomic datasets with ACC- or ethylene-treated roots and found a subtle change in *RBOHC* transcripts, but not significant enough to pass the filtering on this transcriptomic analysis (Harkey et al., 2018, 2019) (Table S1). However, we also performed qRT-PCR in Col-0 and *ein3eil1* treated with and without ACC to examine changes in abundance in the transcript encoding RBOHC. *RBOHC* transcript abundance was twofold higher in Col-0 treated with ACC compared with Col-0 controls. The levels of *RBOHC* transcript abundance in untreated *ein3eil1* were equivalent to Col-0, and the increase by ACC treatment observed in Col-0 was lost in *ein3eil1* (Fig. S3B). To determine whether ethylene signaling regulates RBOH enzyme activity, a spectrophotometric assay using nitro blue tetrazolium (NBT) dye as an electron acceptor was performed. NBT is reduced by superoxide to monoformazan and this reduction can be detected at 530 nm. This experiment was performed in protein extracts from 7-day-old roots of Col-0 and *rhd2-6* with and without 4-h ACC treatment. Older roots were used in this experiment to obtain an adequate amount of protein for these assays. There was a significant twofold increase in monoformazan production in protein extracts of Col-0 roots treated with ACC for 4 h compared with controls



**Fig. 6. RBOHC activity contributes to ACC-induced ROS accumulation.** (A) Representative images of root hairs of Col-0 and *rhd2-6* with and without ACC treatment for 4 h. Black brackets indicate zone 2 of the root. (B) Root hair quantification of untreated and 4 h ACC-treated seedlings. Data are mean±s.e.m. of total root hairs in zone 2 from three experiments ( $n=18-24$  seedlings per experiment). Columns with different letters indicate statistically significant differences ( $P<0.0001$ ) determined by two-way ANOVA followed by Tukey's multiple comparisons test. (C) Quantification of the percentage of root hairs within a certain micron length of Col-0 and *rhd2-6* with and without ACC treatment. Columns with different letters indicate statistically significant differences ( $P<0.0001$ ) determined by two-way ANOVA followed by Tukey's multiple comparisons test. (D) Representative images of PO1 epidermal fluorescence in Col-0 and *rhd2-6* with and without ACC treatment for 4 h. White brackets indicate zone 2 of the root. (E) Quantification of PO1 fluorescence intensity in hair cells and non-hair cells. For each root, PO1 fluorescence was quantified in two hair cells (cells 1 and 5 from Fig. 2) and two non-hair cells (cells 2 and 4 from Fig. 2) and the values pooled to obtain this average. Data are mean±s.e.m. of three experiments ( $n=18-24$  seedlings/experiment). Columns with different letters indicate statistical significance ( $P<0.0001$ ) in PO1 signal compared with other hair cells as determined by two-way ANOVA followed by Tukey's multiple comparisons test. (F) Quantification of RBOH activity as reported as changes in formazan concentration per minute in roots of Col-0 and *rhd2-6* with and without ACC treatment for 4 h. Data are mean±s.e.m. of three experiments. Columns with different letters indicate statistical significance ( $P<0.0001$ ) as determined by two-way ANOVA followed by Tukey's multiple comparisons test. Scale bars: 100  $\mu\text{m}$ . A.U., arbitrary units.

(Fig. 6F). The enzyme activity of roots of *rbohC/rhd2-6* decreased by 70% in both control and ACC-treated conditions than in roots of similarly treated Col-0 (Fig. 6F). These data are consistent with previous results indicating that the RBOHC enzyme constitutes the majority of the RBOH activity in roots (Chapman et al., 2019; Gayomba and Muday, 2020).

#### Transcription of *PRX44* increases with ACC treatment, driving ROS accumulation and ethylene-induced root hair initiation

The *ein3/eil1* transcription factor mutant has reduced root hair initiation and a reduction in PO1 signal in trichoblasts, suggesting that there is transcriptional regulation of ROS-producing enzymes that drive root hair initiation. We examined a previously published microarray time-course experiment in roots treated with ACC (Harkey et al., 2018) to identify candidate transcriptional targets profiling the expression pattern of transcripts encoding both ROS-producing enzymes and proteins linked to trichoblast cell specification. ROP2, ROPGEF3, ROPGEF4, GL2, RSL4 and RHD6, which are linked to

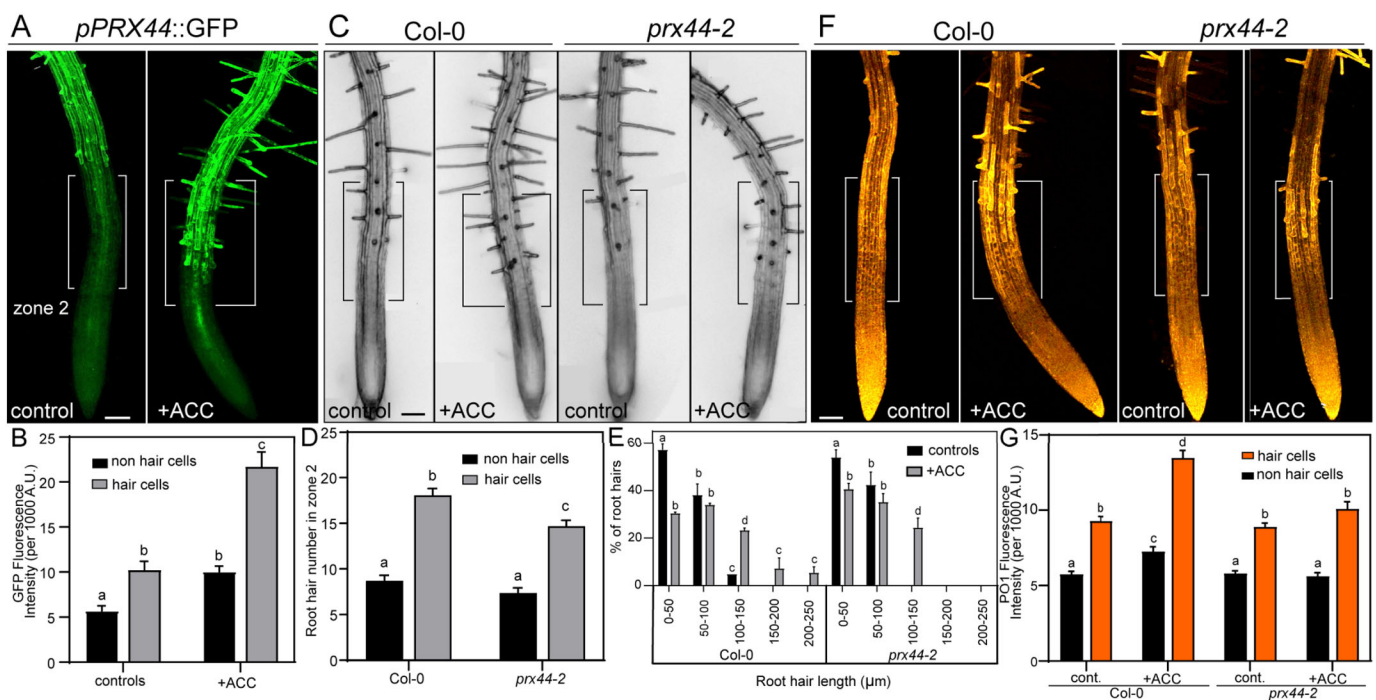
root hair initiation (Denninger et al., 2019), showed no transcriptional response to ACC treatment (Table S1). Primers that recognize RSL1 and RSL2 are not present on the microarray and could not be examined in this dataset. This transcriptomic analysis did not identify the previously reported ethylene induction of *RSL4* (Feng et al., 2017), which was revealed with a high-dose treatment of ethylene that led to ectopic root hair formation from non-hair cells, suggesting that *RSL4* transcript changes may be dose dependent or ethylene specific. A number of transcripts encoding class III peroxidases (PRXs) changed in abundance in response to ACC, including the transcript encoding *PRX44*, which increased threefold. Class III peroxidases are specific to plants and exist in large multigene families. They have been implicated in root hair tip growth, as null mutants have shorter root hairs compared with wild type (Mangano et al., 2017). Recent work has shown that auxin induces expression of genes encoding four class III PRXs, which results in an increase in both root hair length and ROS accumulation (Mangano et al., 2017); however, the role of ethylene signaling in regulating class III PRX expression to modulate root hair initiation or elongation has not been reported.

We examined control and ACC-treated seedlings harboring the *PRX44* promoter driving GFP and examined GFP expression by LSCM. Seedlings treated with ACC showed a statistically significant greater than twofold increase in GFP fluorescence, suggesting that the *PRX44* promoter is induced downstream of ethylene signaling (Fig. 7A,B). qRT-PCR was also performed in Col-0 and *ein3eil1* with and without ACC treatment to examine changes in abundance in the transcript encoding PRX44. We found that there was a twofold increase in *PRX44* transcript abundance in Col-0 treated with ACC compared with Col-0 controls. We also found that there was a significant decrease in *PRX44* transcript abundance in untreated and ACC-treated *ein3eil1* compared with Col-0 (Fig. S3A). We also investigated whether these PRXs participate in ACC-regulated root hair initiation. We examined root hair numbers by light microscopy in Col-0 and *prx44-2* seedlings with and without ACC treatment. The number of root hairs formed in the *prx44-2* mutant treated with ACC was significantly lower than that in ACC-treated Col-0 (Fig. 7C,D). The root hairs in the ACC-treated mutant were also significantly shorter compared with Col-0 (Fig. 7E), which is consistent with the phenotype that has been reported in response to auxin treatment (Mangano et al., 2017). When PO1 accumulation patterns were examined by LSCM, we observed no significant increase in PO1 accumulation in hair cells of *prx44-2* seedlings treated with ACC (Fig. 7F,G). We also examined *prx73-4*, because transcripts encoding PRX73 have been reported

to be induced by auxin treatment to drive root hair elongation (Mangano et al., 2017) and these transcripts increased sixfold in response to ACC treatment (Table S1). When treated with ACC, the *prx73-4* mutant exhibited root hair initiation and ROS accumulation patterns that were similar to those of wild-type seedlings, suggesting that this class III PRX is not involved in ethylene-induced ROS synthesis and root hair initiation (Fig. S4). These combined data indicate that both PRX44 and PRX73 are transcriptionally regulated by ethylene, but only PRX44 contributes to ethylene-induced ROS accumulation in root hairs and increases in root hair initiation.

## DISCUSSION

Ethylene is a key hormonal regulator of root hair initiation and elongation (Guzman and Ecker, 1990; Kieber et al., 1993). Although the proteins that drive ethylene signaling are well characterized, the downstream proteins that control the root hair developmental processes have received less attention. ROS are involved in both root hair initiation and elongation (Foreman et al., 2003; Gayomba and Muday, 2020). In root hairs, one source of ROS is the NADPH oxidase (NOX)/RESPIRATORY BURST OXIDASE HOMOLOG C (RBOHC), which is localized to root hairs to facilitate cell wall loosening and subsequent tip-focused  $\text{Ca}^{2+}$  accumulation leading to root hair cell elongation (Foreman et al., 2003; Monshausen et al., 2007). The plant hormone auxin also increases ROS accumulation to



**Fig. 7. *pPRX44::GFP* expression increases in roots treated with ACC and PRX44 contributes to ethylene-induced root hair formation and  $\text{H}_2\text{O}_2$  accumulation in root hair cells.** (A) Representative images of *pPRX44::GFP* roots with and without ACC treatment for 4 h. White brackets indicate zone 2 of the root. (B) Quantification of GFP signal in hair and non-hair cells of *pPRX44::GFP* with and without ACC treatment. Data are mean+s.e.m. (C) Representative images of Col-0 and *prx44-2* roots with and without ACC treatment for 4 h. Black brackets indicate zone 2 of the root. (D) Root hair quantification of untreated and 4 h ACC-treated seedlings. Data are mean+s.e.m. of three independent experiments ( $n=18-24$  seedlings per experiment). Columns with different letters indicated statistically significant differences ( $P<0.0001$ ) determined by two-way ANOVA followed by Tukey's multiple comparison test. (E) Quantification of the percentage of root hairs within a certain length for Col-0 and *prx44-2* with and without ACC treatment. Columns with different letters indicate statistically significant differences ( $P<0.0001$ ) determined by two-way ANOVA followed by Tukey's multiple comparisons test. (F) Representative images of Col-0 and *prx44-2* with and without ACC treatment for 4 h and stained with PO1. White brackets indicate zone 2 of the root. (G) Quantification of PO1 accumulation in hair and non-hair cells of Col-0 and *prx44-2* with and without ACC treatment for 4 h ( $n=12-18$  seedlings per experiment). For each root, PO1 fluorescence was quantified in two hair cells (cells 1 and 5 from Fig. 2) and two non-hair cells (cells 2 and 4 from Fig. 2) and the values pooled to obtain this average. Data are mean+s.e.m. of three experiments. Columns with different letters indicate statistical significance ( $P<0.0001$ ) compared with all other hair cells as determined by two-way ANOVA followed by Tukey's multiple comparisons test. Scale bars: 100  $\mu\text{m}$ . A.U., arbitrary units.

drive root hair elongation (Mangano et al., 2017), suggesting that ROS could also act as a signaling molecule in ethylene-induced root hair initiation and elongation (Foreman et al., 2003; Gayomba and Muday, 2020; Jones et al., 2007; Takeda et al., 2008). In this study, we examined the effects of ethylene signaling on root hair initiation and ROS accumulation to determine whether increased ethylene drives root hair initiation through regulation of ROS-producing enzymes and subsequent ROS accumulation in trichoblasts from which root hairs form.

We examined root hair number and ROS accumulation in Col-0 treated with low concentrations of ethylene gas (0.5 ppm) or the ethylene precursor ACC (0.7  $\mu$ M) for 4 h. There was an increase in the number of trichoblasts that formed root hairs in ethylene- and ACC-treated seedlings compared with untreated controls, which was most pronounced in a 500  $\mu$ m region starting 500  $\mu$ m from the root tip, which we call zone 2. Using a ratiometric reporter of ROS-induced gene expression, ZAT12p-ROS, we found that in ACC-treated roots, ROS-dependent gene expression was increased in zone 2 in ACC-treated roots compared with untreated controls. We also examined the effect of ethylene and ACC on H<sub>2</sub>O<sub>2</sub> accumulation using a boronate-based H<sub>2</sub>O<sub>2</sub>-selective sensor, PO1. Both ethylene and ACC led to elevated H<sub>2</sub>O<sub>2</sub> accumulation in the trichoblast cells in zone 2 after 4 h of treatment, but PO1 signal was at constant lower level in atrichoblast cells. There was a significant increase in PO1 signal within 2 h after ACC treatment in cells that had not yet formed root hair bulges, suggesting that ethylene-induced ROS acts to drive root hair initiation.

To determine whether canonical ethylene signaling controls ACC-dependent ROS synthesis driving root hair formation, we examined these responses in several ethylene signaling mutants. In the ethylene-insensitive *ein3eil1* and *etr1-3* mutants, the effect of ACC treatment on ROS synthesis and root hair formation was lost. In contrast, the *etr1-7* mutant allele, which has constitutive signaling, had increased ROS and root hair formation. These findings are consistent with our prior reports that showed that effects of this ACC dose and under these growth conditions on root elongation, gravitropism and lateral root formation are lost in the ethylene-insensitive *ein2-5* and *etr1-3* mutants (Harkey et al., 2018; Lewis et al., 2011; Negi et al., 2010). These findings contrast with other processes for which higher doses of ACC can affect development without conversion to ethylene, especially when the ACC oxidase enzyme is limited (Li et al., 2022; Polko and Kieber, 2019; Van de Poel, 2020; Vanderstraeten et al., 2019). We also did not observe ectopic root hair formation in response to these short-term and low-dose treatments with ACC and ethylene, as reported recently when seedlings were treated with 10 ppm ethylene for 2 days (Qiu et al., 2021), which is a 12-fold longer and 20-fold higher concentration.

If ROS is driving root hair formation, then it is predicted that mutants with impaired root hair initiation may have altered ROS levels. We examined the PO1 signal in the *caprice* (*cpc*) mutant, which forms no root hairs, and the *werewolf* (*wer*) mutant, which forms ectopic root hairs, both of which have mutations in MYB-like DNA-binding proteins (Lee and Schiefelbein, 1999; Wada et al., 1997). The *cpc* mutant showed a significant decrease in PO1 signal when it was measured using line profiles along the root whereas *wer* showed a significant increase, suggesting that mutants with altered root hair formation also show altered ROS accumulation patterns. Similarly, the *rhd2-6* mutant, which was isolated for altered root hair elongation (Foreman et al., 2003) and has impaired root hair formation (Gayomba and Muday, 2020), also had reduced levels of ROS, suggesting a positive relationship between root hair formation and ROS levels using root hair mutants.

Based on previous work detailing the role of RBOHC in root hair initiation and growth (Foreman et al., 2003; Gayomba and Muday, 2020), we also investigated whether this enzyme was contributing to ethylene-induced ROS accumulation and subsequent root hair initiation and/or elongation. ACC-treated *rhd2-6*, which is an *RBOHC* null mutant, had a reduced number of root hairs, less root hair elongation, and decreased ROS accumulation in trichoblasts compared with Col-0. We also observed a more than 200% increase in RBOH activity in response to ACC and a 70% reduction in activity in *rhd2-6* (*rbohc*). These data suggest that ethylene signaling induces ROS production in roots via increases in RBOHC enzyme activity to drive root hair initiation and elongation, although increased ROS production could also come from enhanced RBOHC synthesis. We examined several transcriptomic datasets (Harkey et al., 2018, 2019) and found evidence of changes in *RBOHC* transcripts in response to ACC treatment, which were confirmed by qRT-PCR. The ACC induction was lost in the *ein3eil1* mutant.

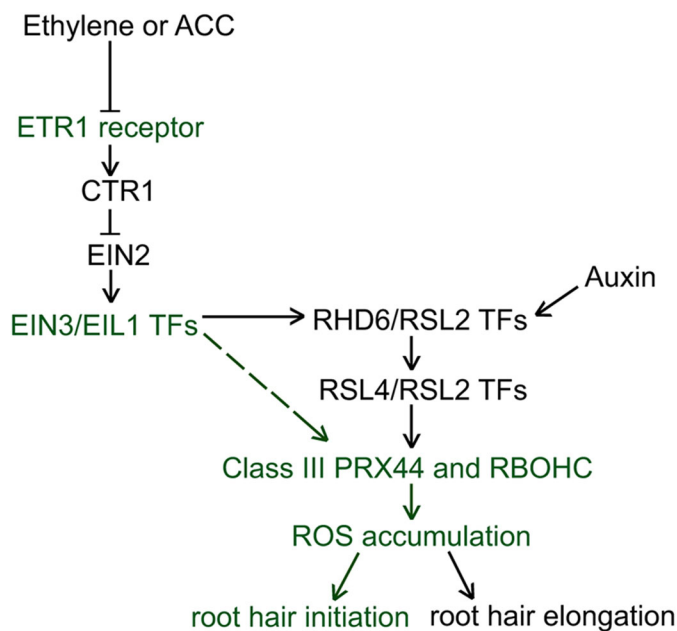
To determine whether other root hair-specific proteins and/or ROS producers are regulated by ethylene, we widened our analysis of these previously published ACC and ethylene transcriptomic datasets. Although ACC did not substantially change the abundance of transcripts encoding RHD2 (RBOHC), ROP2, ROPGEF3, ROPGEF4, GL2, RHD6 or RSL4, we identified several transcripts that were substantially increased in response to ACC. Of particular interest were increases in transcripts encoding two class III peroxidase enzymes, PRX44 and PRX73, which showed a significant increase in abundance after 4 h of ACC treatment. ACC treatment of seedlings harboring a *pPRX44::GFP* construct led to significant increases in GFP fluorescence intensity in zone 2 compared with untreated controls, consistent with transcriptional regulation of this gene. We also performed qRT-PCR and found a twofold increase in *PRX44* transcripts in response to ACC treatment. Additionally, this ACC effect was lost in the *ein3eil1* mutant and the abundance of *PRX44* transcripts was significantly lower in *ein3eil1* even in the absence of ACC treatment.

ACC treatment of the null mutant *prx44-2* led to decreased ROS accumulation in root hair-forming trichoblasts, whereas there was no effect in *prx73-4* compared with Col-0. The decreased PO1 signal in *prx44-2* was accompanied by significant reductions in the number of root hairs formed in this same root region and the length of these root hairs. Together, these experiments implicate RBOHC and class III PRXs in ethylene-regulated ROS accumulation and root hair initiation.

These class III PRXs have also been implicated as regulators of root hair growth in response to environmental cues. Two PRXs are involved in auxin-dependent root hair elongation, as null mutants treated with auxin have shorter root hairs and less ROS compared with wild type (Mangano et al., 2017). Another report demonstrated that PRXs function to modify the localization of extensin proteins to control cell wall structure to allow increased elongation (Marzol et al., 2022). Recent work has also indicated a role for PRXs as positive regulators of root hair growth at low temperatures (Pachecho et al., 2021). These results showed that low temperature-induced root hair growth required PRX activity and upregulated expression of PRXs that modulate ROS homeostasis and extensin stabilization to drive root hair growth (Pachecho et al., 2022). Prior publications and these results are consistent with a role for PRXs as drivers of root hair development and modulation of the apoplastic ROS pool.

To summarize our findings on ethylene-dependent root hair initiation via increased ROS synthesis and to tie these findings in with other studies, we present a model in Fig. 8. Low-dose and short (4-h) treatments with ACC led to efficient conversion of ACC to ethylene





**Fig. 8. Summary of the mechanisms by which ethylene and auxin modulate ROS and root hair formation.** In hair cells, ethylene, acting through the ETR1 receptor and the canonical ethylene signaling pathway, induces accumulation of EIN3/EIL1 TF proteins. Previous work has shown that EIN3/EIL1 physically interact with RHD6/RSL2 TFs to induce expression of *RSL4/RSL2* transcripts leading to root hair elongation (Feng et al., 2017). It has also been shown that RSL4/RSL2 is induced by auxin signaling and that RSL4/RSL2 bind to the promoters of RBOH and class III PRX genes to induce transcript expression and root hair elongation (Mangano et al., 2017). Here, in green text, we have shown that ethylene signaling through ETR1 and EIN3/EIL1 induces ROS accumulation through RBOHC enzyme activity and *PRX44* transcript abundance to drive root hair initiation.

by roots to initiate an ethylene signaling cascade. Ethylene acts through the ETR1 receptor and the canonical signaling pathway to increase the activity of EIN3/EIL1 TFs. EIN3/EIL1 binds to the root hair-specific TF RHD6 to induce RSL4 expression and root hair elongation (Feng et al., 2017). Root hair growth is controlled by several proteins leading to activation of the RHD6 TF. Downstream of RHD6, the RSL4 TF acts to define final root hair length based on its level of expression (Datta et al., 2015; Mangano et al., 2017). Previous work has shown that EIN3 physically interacts with RHD6 to form a transcriptional complex that co-activates RSL4 to promote root hair elongation (Feng et al., 2017), and that EIN3 and RSL4 both act to downregulate *GL2* expression to drive ectopic root hair formation with long-term treatments with high levels of ethylene (Qiu et al., 2021). It has also been shown that auxin treatment results in increased RSL4 transcript abundance, which binds to the promoters of RBOH and four class III PRX genes, including *PRX44*, to drive ROS synthesis in root hair cells. RSL4 activation via auxin induces class III PRX and RBOH expression, which leads to the ROS accumulation required to drive root hair elongation (Mangano et al., 2017). We find that ACC treatment increases the activity of RBOHC enzymes, leading to increased ROS and root hair elongation, without changing the abundance of *RBOHC* transcripts. Additionally, ACC treatment leads to the transcriptional regulation of class III PRXs and RBOHC, resulting in ROS production and root hair initiation. Although the reduction in ROS in *rh2-6* and *prx44-2*, in both the presence and absence of ACC was expected, we did not predict that there would still be an increase in root hair number in response to ACC. These data suggest that the ROS-selective dyes that we have

used may not be sensitive enough to detect the small changes in ROS that occur prior to root hair initiation in these mutants or that there are additional mechanisms that facilitate ethylene-induced root hair formation, including redundant action of these two enzymes. Further investigation regarding the mechanisms of ethylene-regulated ROS synthesis is required. For example, one important question to be answered is how ethylene signaling regulates RBOH activity. This could occur through a number of different mechanisms, such as calcium binding or phosphorylation, which are known to regulate RBOH enzymes (Postiglione and Muday, 2020).

This work supports the role of ROS as a signaling molecule, whereby ROS are produced in response to hormonal cues to drive developmental processes. We provide new evidence for regulation by ethylene of the ROS-producing enzymes RBOHC and the class III PRX44. However, there are other ROS-producing and scavenging enzymes, such as SODs, that require additional study to understand the numerous developmental mechanisms in which ROS may play a role. It is also crucial to understand the molecular mechanisms by which ROS regulates the activity of proteins that drive root hair initiation and elongation. ROS can oxidize cysteine residues in proteins to change their conformation and activity (Couturier et al., 2013) so an important next step is to identify those root hair proteins that are modified by ROS to drive this important developmental process.

## MATERIALS AND METHODS

### Growth conditions and root hair quantification

All *Arabidopsis* mutants were in the Col-0 background. The *etr1-7*, *etr1-3* (AT1G66340), *ein3-1* (AT3G20770) *eil1-1* (AT2G27050) double and *rh2-6* (AT5G51060) mutants have all been described previously (Binder et al., 2007; Gayomba and Muday, 2020; Harkey et al., 2018; Hua and Meyerowitz, 1998). The ZAT12p-ROS construct was generously provided by Won-Gyu Choi (Lim et al., 2019). *prx44-2* (AT4G06010) and *prx73-4* (AT5G67400) were obtained from the *Arabidopsis* SALK center (SALK\_057222C and SALK\_020724, respectively), the mutations were verified by PCR, and homozygous lines were isolated. The transcriptional reporter *pPRX44::GFP* transgenic line was described previously (Marzol et al., 2022). Seeds were sterilized in 75% ethanol for approximately 5 min and grown on 100×15 mm Petri dishes containing 25 ml media. Seedlings were grown on 1× Murashige and Skoog (MS) medium supplemented with 1% sucrose, vitamins (1 μg/ml thiamine, 0.5 μg/ml pyridoxine and 0.5 μg/ml nicotinic acid), 0.05% 2-ethanesulfonic acid (MES) (w/v) and 0.8% (w/v) agar. Media pH was 5.5. Micropore tape was used to seal the top of the Petri dish and plated seeds were stratified at 4°C in darkness for 2 days. Plants were grown under 24 h light from T5 fluorescent lights at 120–150 μmol photons m<sup>-2</sup> s<sup>-1</sup>.

### ACC and ethylene treatments

ACC stocks at 200 mM were prepared using ACC hydrochloride diluted in water. ACC was then added directly to media to yield a final concentration of 0.7 μM. Seeds were grown on control plates as described above for 5 days and then transferred to plates containing either 0.7 μM of ACC or transferred to new plates and placed in an air-tight tank containing 0.5 ppm ethylene gas. Seedlings were then left under the light conditions described above for 4 h and then used for experiments.

### Quantification of root hair number and root hair length

Seedlings were grown for 5 days on control media and then transferred to media containing 0.7 μM ACC and grown for 4 h or other indicated times. To examine root hairs, 5-day-old seedlings were imaged by bright-field microscopy on an Axio Zoom V16 stereomicroscope. Extended depth of focus was used to combine z-stack images. Root hairs that were at stage +1 or +2 (or later stages) as defined by Denninger et al. (2019) were quantified using Fiji/ImageJ software in three zones (0–500 μm, 500–1000 μm, 1000–1500 μm) starting from the root tip.

We also determined root hair length in the presence and absence of ACC treatment in Col-0 and mutants. All root hairs in zone 2 that had begun to elongate (were at stages above +2) were measured using Fiji/ImageJ software. The length of each root in microns was determined. To obtain the histogram of distribution, the roots were separated into bins of different lengths, and the percentage of root hairs in each length bin was determined by dividing by the total number of root hairs in zone 2 in that root. All statistics in this experiment and following experiments were performed using GraphPad Prism software.

### Confocal imaging of dyes and reporters of ROS levels

H<sub>2</sub>O<sub>2</sub> was visualized with PO1. PO1 was dissolved in DMSO to make a 500 μM stock and was further diluted in water to make a 50 μM working solution. Seedlings were incubated in PO1 for 15 min in the dark and were then rinsed with water and mounted in water for imaging. Control, ACC-treated and ethylene-treated seedlings were imaged on a Zeiss 880 laser-scanning confocal microscope using a 10× objective. PO1 was excited with a 488 nm laser at 0.25% power and emission was collected between 544 and 695 nm. Images were analyzed using Fiji/ImageJ software. Plot profiles were taken across the epidermal cell files of maximum intensity projections using a 20-pixel line and values were averaged within each cell file. All images were captured at levels below saturation, although images shown have been uniformly adjusted for brightness and contrast for clarity.

Five-day-old seedlings harboring the ZAT12p-ROS construct were mounted in water and excited with 488 and 561 nm lasers at 6% and 1.2% laser power, respectively. GFP emission was collected at 521 nm and mCherry emission was collected at 593 nm. Fiji/ImageJ software was used to generate two single-channel images to form individual GFP and mCherry channels. Plot profiles were taken using a 250-pixel-wide line to measure fluorescence of the entire root. Measurements were taken starting at 200 μm back from the root tip and ending at 1500 μm. Ratios were generated by dividing the GFP channel by mCherry using the Image Calculator tool (Process/Image Calculator).

Microscopic images were exported as CZI files and were opened in Fiji/ImageJ software. Scale bars (100 μm) were added to each image as well as brackets separating the three 500 μm sections from the root tip, referred to as zones 1, 2 and 3. Signal was quantified in each region. For creating images for this paper, these images were saved and uploaded into Photoshop where they were uniformly adjusted to increase brightness and clarity of images. Images edited via Photoshop were opened in Illustrator where the scale bars and initial brackets were covered with black boxes and new brackets redrawn so that all images were uniform and consistent in labels.

### PI staining and proPRX44::GFP imaging

Cell walls were stained with 0.5 μg/ml PI dissolved in water. Five-day-old seedlings with and without ACC treatment were incubated in PI for 4 min before imaging. Fluorescence was visualized using a 561 nm laser at 0.15% and emission spectra set to 561-695 nm. These settings were used for all images. Optical slices of the top section of the root to show epidermal cells and maximum intensity projections of z-stack images are shown in Fig. S1. Fiji software was used to measure cell length of three epidermal cells per root using optical slices of the top section of the root. Cells were chosen at the bottom of zone 2 (cell 1), in the middle (cell 2) and at the top (cell 3). Four cells of each of the three cell types were averaged and are shown in Fig. S1. Five-day-old seedlings were mounted in water and excited with a 488 nm laser at 5% and emission was collected between 490-606 nm. Seedlings were then stained with PI.

### DNA extraction and PCR for mutant genotype analysis

T-DNA insertion lines were grown for approximately 2 weeks on media described above, then transferred to soil and grown under 24 h light at 50-80 μmol photons m<sup>-2</sup> s<sup>-1</sup>. Leaves were harvested from 6-week-old plants and stored in Eppendorf tubes at -20°C prior to DNA extraction. Frozen leaves were ground in DNA extraction buffer [100 mM Tris-HCl pH 8.0, 10 mM Na<sub>2</sub>-EDTA, 100 mM LiCl<sub>2</sub>, 1% (w/v) SDS]. Samples were then washed once with isopropanol and three times with 80% (v/v) ethanol. Finally, DNA pellets were dried, resuspended in sterile water and stored at

-20°C. Each PCR reaction contained 1× GoTaq Polymerase, 1 μM of each primer, 1 μl of DNA and 5 μl water. To confirm that mutants were homozygous for the desired T-DNA insertion, one reaction was performed with primers (left primer, TTCAGAGTCTGCAGAAGGAATG; right primer, CATTGCAGTCTCTCCAGGAC) flanking the left and right sides of intact genes and one reaction was performed with the right primer and LBb1.3, which is a primer specific for the left border of the T-DNA insertion. PCR products were run on 0.8% (w/v) agarose gels containing 0.002% SERVA DNA stain G.

### RNA isolation and quantitative RT-PCR

RNA was isolated from seedlings grown on a nylon filter as described previously (Harkey et al., 2018). Five days after germination, seedlings were transferred to either control media or media containing 0.7 μM ACC and were grown for 4 h. Roots were then cut and flash-frozen in liquid nitrogen then ground using a mortar and pestle. RNA was isolated using the QIAGEN plant RNeasy kit protocol and RNA was DNase treated.

cDNA synthesis was performed using the RevertAid RT Reverse Transcription kit (Thermo Fisher). qRT-PCR analysis using this cDNA was performed on a QuantStudio real-time PCR machine using SYBR green detection chemistry. Primers specific to *PRX44* (FP: 5'-CAAGAG-ACTCGGTGCGATTAG-3'; RP: 5'-TTGTTGGTCCGGGTAAGTTC-3') and *RBOHC* (FP: 5'-CAAGGAACAAGCCCACTAAA-3'; RP: 5'-TTCT-ATTGGGTTACGCGTGAG-3') were used and relative transcript abundance was quantified using *ACT2* primers (FP, 5'-TGAGAGATTCAGATGCC-CAGAA-3' RP, 5'-GCAGCTTCCATTCCACAA-3') using comparative Ct analysis (ΔΔCt) to determine the relative quantity of target transcripts.

### NADPH oxidase/RBOH enzyme assays

Protein extract was isolated from seedlings grown on a nylon filter as described previously (Harkey et al., 2018). After 2 days of stratification and 7 days of growth under the conditions described above, the nylon was transferred to growth medium with and without 0.7 μM ACC for 4 h. Roots were then cut from seedlings and flash-frozen in liquid nitrogen. Frozen samples were ground in liquid nitrogen using a mortar and pestle. RBOH extraction buffer [50 mM Tris-HCl pH 7.5, 0.1 mM EDTA, 0.1% (v/v) Triton X-100, 1 mM MgCl<sub>2</sub>, 10% (v/v) glycerol] was then added to a plant material/buffer ratio of 1:3 (w/v). Samples were centrifuged at 16,000 g for 10 min and the supernatant was collected, desalted and concentrated using Amicon Ultra-0.5 Centrifugal Filter devices. RBOH reaction mixture (50 mM Tris-HCl pH 7.5, 1 mM CaCl<sub>2</sub>, 0.1 mM NBT, 0.1 mM NADPH) was then mixed with protein extract at a 1:1 ratio. The reduction of NBT to monoformazan was monitored spectrophotometrically at 530 nm. Monoformazan concentrations were calculated using a 12.8 mM<sup>-1</sup> cm<sup>-1</sup> extinction coefficient. This assay was adapted a previously described protocol using sweet peppers (Chu-Puga et al., 2019).

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### Competing interests

The authors declare no competing or financial interests.

### Author contributions

Conceptualization: R.E.M., G.K.M.; Methodology: G.K.M.; Formal analysis: R.E.M.; Investigation: R.E.M.; Resources: E.M., J.M.E.; Writing - original draft: R.E.M.; Writing - review & editing: G.K.M., J.M.E.; Visualization: R.E.M.; Supervision: G.K.M.; Project administration: G.K.M.; Funding acquisition: R.E.M., G.K.M., J.M.E.

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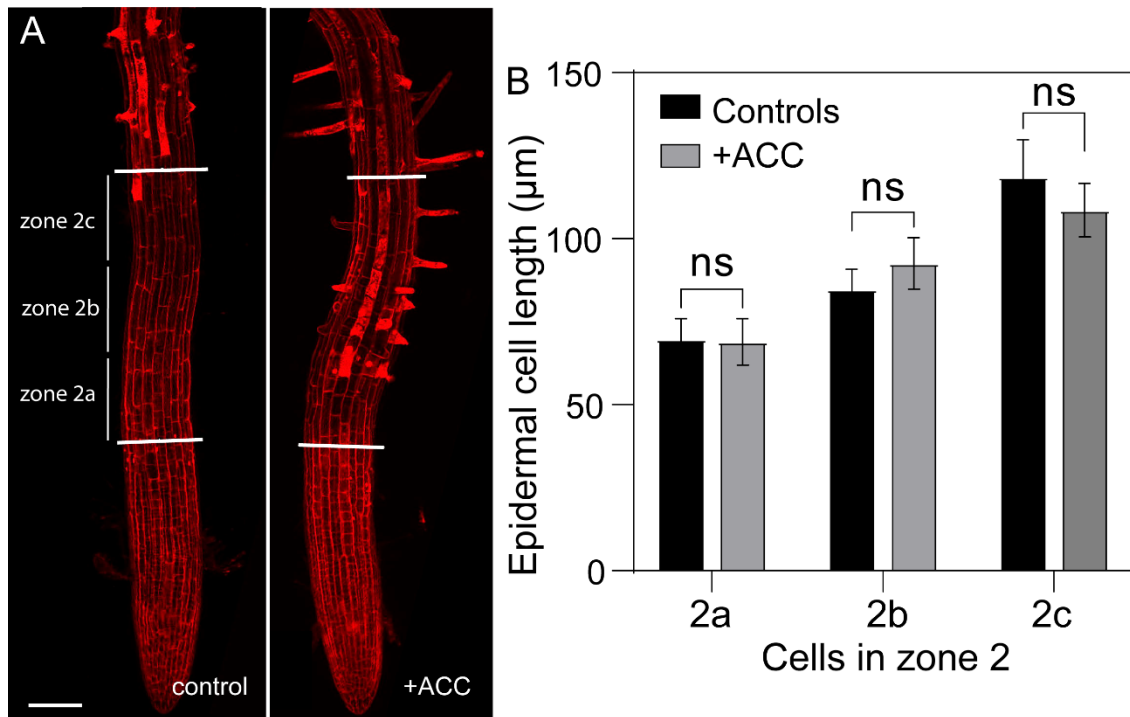
## Peer review history

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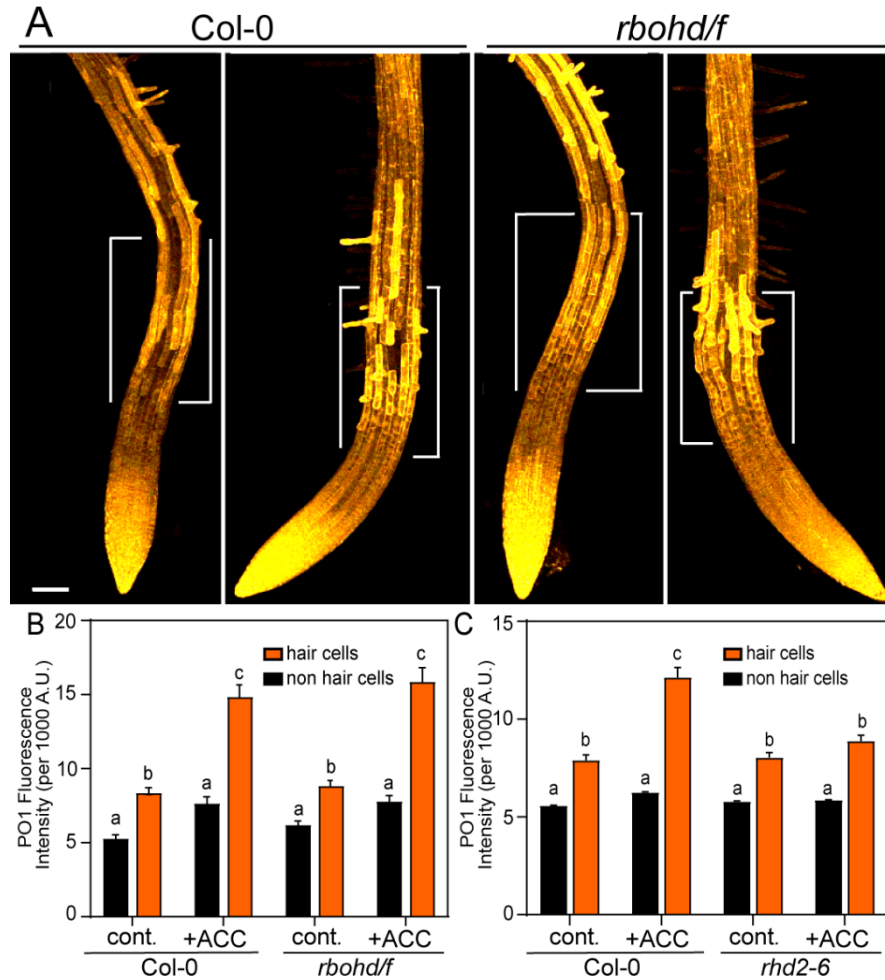
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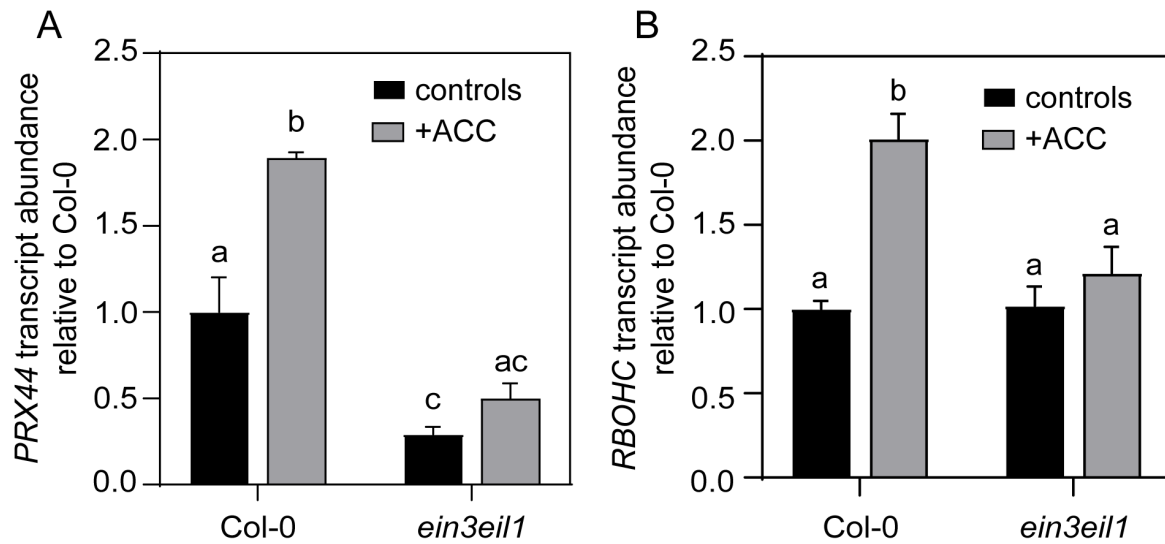
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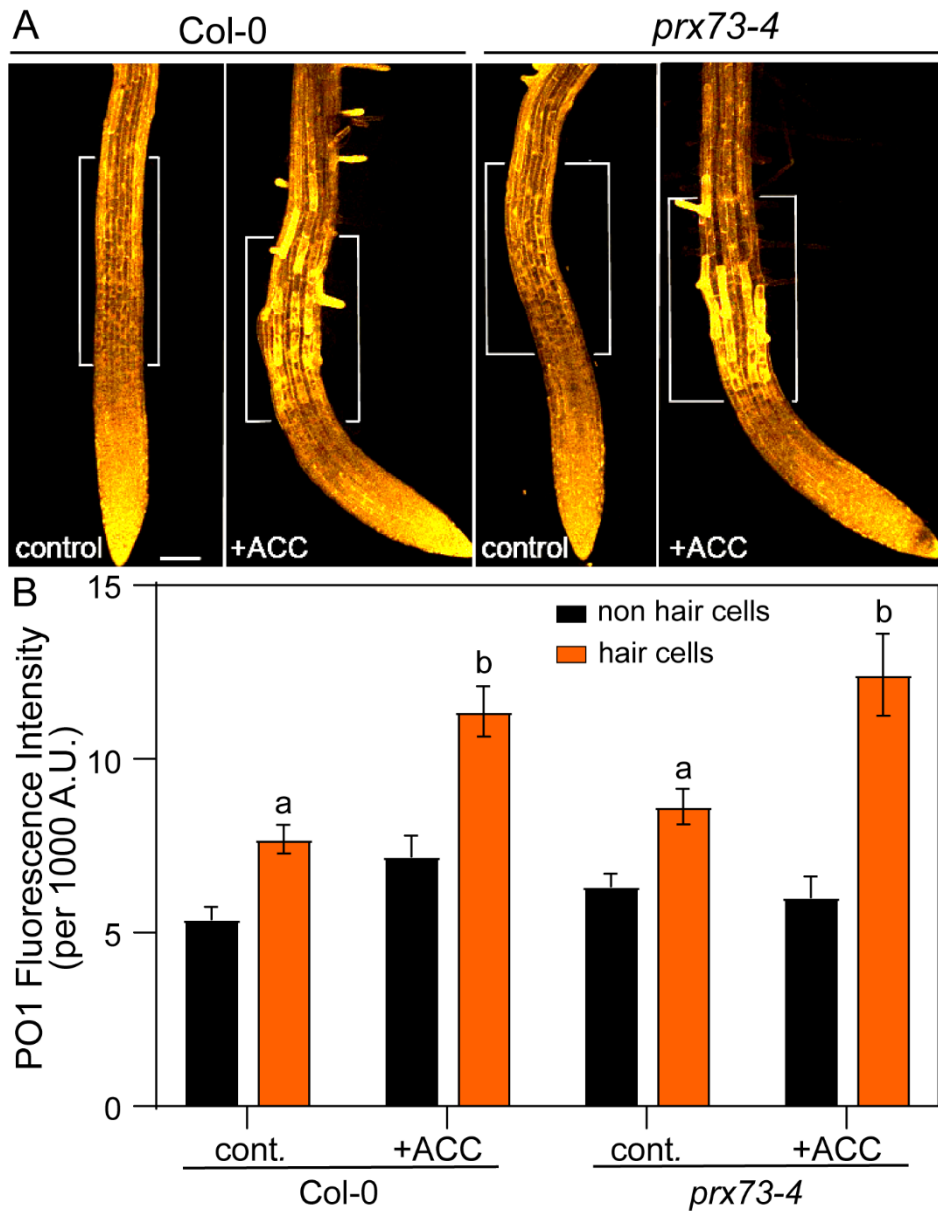
**Fig. S1. Epidermal cell length did not change after 4 hours of ACC treatment (A)** Representative images of Col-0 seedlings treated with and without ACC for 4 hours and stained with propidium iodide (PI). Scale bar is 100  $\mu\text{m}$ . (B) Quantification of an average length of 12-18 epidermal cells. 3 cells per zone were analyzed, a cell towards the bottom of the zone 2 (cell 2a), a cell in the middle (cell 2b) and a cell at the top of the zone (cell 2c). Data are  $\pm$  SEM of 3 independent experiments ( $n=12-15$  seedlings per experiment). Data was statistically analyzed using two-way ANOVA followed by Tukey's multiple comparisons test.



**Fig. S2. RBOHD and RBOHF do not contribute to ethylene-induced ROS synthesis in root hair cells** (A) Representative images of PO1 epidermal fluorescence in Col-0 and *rbohdf* treated with and without ACC for 4 hours. Scale bar is 100  $\mu$ m. (B) Quantification of PO1 fluorescence intensity in hair cells (1, 3, and 5) and non-hair cells (2, 4) of Col-0 and *rbohdf* treated with and without ACC. Data are means  $\pm$  SEM. of 3 experiments (n=12-18 seedlings/experiment). Columns with different letters indicate statistical significance compared to other hair cells as determined by two-way ANOVA followed by Tukey's multiple comparisons test. (C) For comparison, quantification of PO1 fluorescence intensity in hair cells and non-hair cells of Col-0 *rhd2-6* (*rbohdc*) treated with and without ACC from Figure 6E is shown. Data are means  $\pm$  SEM. of 3 experiments (n=12-18 seedlings/experiment). Columns with different letters indicate statistical significance compared to other hair cells as determined by two-way ANOVA followed by Tukey's multiple comparisons test.



**Fig. S3. *PRX44* and *RBOHC* transcript abundance is regulated by *EIN3/EIL1* (A) *PRX44* and (B) *RBOHC* transcript abundance in Col-0 and *ein3eil1* mutants treated with and without 0.7  $\mu$ M ACC for 4 hours as determined by qRT-PCR. Averages of 3 replicates are reported. Different letters represent statistical significant differences as determined by two-way ANOVA followed by Tukey's multiple comparisons test.**



**Fig. S4. PRX73 does not contribute to ACC-induced ROS synthesis in root hair cells** (A) Representative images of PO1 epidermal fluorescence in Col-0 and *prx73-4* treated with and without ACC for 4 hours. Scale bar is 100  $\mu$ m. (B) Quantification of PO1 fluorescence intensity in hair cells (1, 3, and 5) and non-hair cells. Data are means  $\pm$  SEM. of 3 independent experiments (n=15-18 seedlings per experiment). Columns with different letters indicate statistical significance compared to other hair cells as determined by two-way ANOVA followed by Tukey's multiple comparisons test.



Table S1. Transcript abundance changes in genes encoding RBOHC, class III peroxidases, and root hair patterning genes

		Signal log ratio values after ACC treatment (hours) <sup>a</sup>							
Gene Symbol	Locus ID	0	0.5	1	2	4	8	12	24
RHD2 (RBOHC) <sup>b</sup>	AT5G51060	-0.09	0.32	0.28	0.33	0.37	0.05	-0.06	0.16
PRX44 <sup>c</sup>	AT4G26010	0.40	-0.78	0.47	0.67	0.68	0.84	0.89	1.19
PRX69 <sup>c</sup>	AT5G64100	-0.09	0.51	0.80	0.96	1.23	1.32	1.50	0.80
PRX73 <sup>b</sup>	AT5G67400	0.6	-1.1	0.3	0.39	1.42	1.2	1.2	1.43
PRX39 <sup>c</sup>	AT4G11290	0.07	0.00	0.15	0.24	0.86	0.97	1.51	0.62
PRX25 <sup>c</sup>	AT2G41480	-0.78	0.60	0.20	-0.71	0.28	-0.20	-0.08	0.53
PRX72 <sup>c</sup>	AT5G66390	-0.09	0.11	-0.05	-0.12	-0.43	-0.94	-0.96	-1.02
PRX64 <sup>c</sup>	AT5G42180	0.08	-0.35	0.02	0.00	-0.44	-0.81	-0.84	-1.06
PRX66 <sup>c</sup>	AT5G51890	0.01	-0.02	0.00	0.21	-0.27	-0.45	-0.82	-1.32
ROP2 <sup>b</sup>	AT1G20090	0.08	-0.04	-0.13	-0.11	-0.12	-0.31	-0.22	-0.28
ROPGEF3 <sup>b</sup>	AT4G00460	0.03	0.00	0.28	-0.08	-0.06	-0.11	-0.12	-0.12
ROPGEF4 <sup>b</sup>	AT2G45890	0.26	-0.59	-0.08	0.21	-0.04	-0.22	-0.30	-0.34
GL2 <sup>d</sup>	AT1G79840	-0.06	-0.46	-0.36	-0.23	-0.42	-0.32	-0.44	-0.29
RHD6 <sup>b</sup>	AT1G66470	0.10	-0.56	0.06	-0.01	-0.17	-0.31	-0.24	-0.09
RSL4 <sup>b</sup>	AT1G27740	-0.25	0.00	-0.19	-0.10	0.04	-0.01	-0.30	-0.22

<sup>a</sup>Signal log ratio (SLR; log base 2 of fold change) was calculated for each transcript relative to an averaged time matched control (Harkey et al 2018); <sup>b</sup>transcripts that did not change significantly in response to ACC in either the DE or PA filtering; <sup>c</sup>transcripts that were differentially regulated in response to ACC; <sup>d</sup>transcripts that were present (above background) at some time points while absent at others (PA).

