Defense-related callose synthase *PMR4* promotes root hair callose deposition and adaptation to phosphate deficiency in *Arabidopsis thaliana*

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Summary

Plants acquire phosphorus (P) primarily as inorganic phosphate (Pi) from the soil. Under Pi deficiency, plants induce an array of physiological and morphological responses, termed phosphate starvation response (PSR), thereby increasing Pi acquisition and use efficiency. However, the mechanisms by which plants adapt to Pi deficiency remain to be elucidated. Here, we report that deposition of a β-1,3-glucan polymer called callose is induced in *Arabidopsis thaliana* root hairs under Pi deficiency, in a manner independent of PSR-regulating *PHR1/PHL1* transcription factors and *LPR1/LPR2* ferroxidases. Genetic studies revealed *PMR4* (*GSL5*) callose synthase being required for the callose deposition in Pi-depleted root hairs. Loss of *PMR4* also reduces Pi acquisition in shoots and plant growth under low Pi conditions. The defects are not recovered by simultaneous disruption of *SID2*, mediating defense-associated salicylic acid (SA) biosynthesis, unlike SA-dependent powdery mildew resistance previously described in *pmr4*. Grafting experiments and characterization of plants expressing *PMR4* specifically in root hair cells suggest that a PMR4 pool in the cell type contributes to shoot growth under Pi deficiency. Our findings thus suggest an important role for *PMR4* in plant adaptation to Pi deficiency.

Significance statement

We reveal that PMR4 callose synthase mediates callose deposition in root hairs under phosphate (Pi) deficiency, without requiring Pi starvation response regulators *PHR1/PHL1* or *LPR1/LPR2*. The loss of the callose deposition is accompanied by decreases in Pi acquisition and plant growth in *pmr4*. Root hair cell-specific *PMR4* expression restores callose deposition in root hairs and shoot growth under Pi deficiency, indicating a critical role for root hair callose in plant adaptation to Pi deficiency.

Keywords: root hair, cell wall, Glucan Synthase Likes (GSLs), phosphate acquisition, phosphate transport, plant nutrition, nutrient deficiency, salicylic acid (SA), pattern triggered immunity

Introduction

As an essential macronutrient, plants acquire and distribute P from the rhizosphere to the whole tissues, primarily in the form of inorganic phosphate (Pi). To maintain Pi homeostasis under Pi deficiency, plants induce an array of adaptive responses, *e.g.* increasing Pi uptake, remobilization and utilization, termed Pi starvation response (PSR), and involve beneficial microbes in mutualistic associations (Péret et al., 2011; Chien et al., 2018; Isidra-Arellano et al., 2021). PSR is accompanied by extensive transcriptional reprogramming, mainly through the transcription factor *PHR1* and related *PHL1*, which leads to increases in Pi transporter expression, root hair formation and anthocyanin accumulation (Rubio et al., 2001, Bustos et al., 2010). *PHR1/PHL1* mediate systemic coordination of these responses according to the internal Pi availability. In addition, Pi depletion leads to remodeling root system architecture, a decrease in primary root growth while increasing lateral root formation and growth, through the ferroxidases *LPR1* and *LPR2* (Svistoonoff et al., 2007; Péret et al., 2014; Balzergue et al., 2017). *LPR1/LPR2* are required for arresting primary root growth when the external Pi levels are reduced surrounding the root tips (Svistoonoff et al., 2007). Recent studies illuminated an intimate relationship between PSR and biotic interactions, but the molecular basis for their coordination remains poorly understood (Chan et al., 2021).

Callose deposition on the plasma membrane through callose synthases *Glucan Synthase Likes* (*GSLs*) reinforces barrier functions of the cell wall and restricts plasmodesmatal permeability, in plant development and biotic/abiotic stress responses (Jacobs et al., 2003; Nishimura et al., 2003; Dong et al., 2005; Enns et al., 2005; Nishikawa et al., 2005; Chowdhury et al., 2014; Ellinger & Voigt, 2014; Sager & Lee,

2014; Wu et al., 2018). In *Arabidopsis thaliana* (hereafter Arabidopsis), callose deposition is induced in the root phloem in response to excess iron, thereby reducing plasmodesmatal permeability (O'Lexy et al., 2018). Of 12 Arabidopsis *GSLs*, *GSL2* (*CALS5*) and *GSL5* (*PMR4*, *CALS12*) respectively contribute to basal accumulation and iron-induced deposition of callose in the root phloem (O'Lexy et al., 2018). Callose also accumulates in the stem cell niche (SCN) surrounding the quiescent center cells, thereby inhibiting cell-to-cell movement of the transcription factor SHORT ROOT (SHR) during PSR. This serves to suppress cell division and elongation in the root apical meristem and elongation zone, respectively, through the *LPR1/LPR2* pathway (Müller et al., 2015; Mora-Macías et al., 2017). *GSL5* (*PMR4*) also mediates callose deposition in response to pathogen challenge and microbe/damage-associated molecular patterns (MAMPs/DAMPs), and promotes fungal penetration resistance (Wang et al., 2021). However, the loss of *GSL5* function results in strong resistance to powdery mildew fungi through tightened SA-based defenses, demonstrated by *gsl5* mutants revealed as *powdery mildew resistance4* (*pmr4*) (Vogel & Somerville, 2000; Nishimura et al., 2003). This resistance is thought to represent a backup defense activated when *GSL5* (*PMR4*)-mediated callose deposition, a key output of MAMP/DAMP-triggered defense signaling, is disrupted. Although the significance of *PMR4* has been genetically shown, our knowledge is very limited for the mechanisms that involve and regulate PMR4 in callose deposition during defense responses (Wang et al., 2021). Our understanding is also limited for the possible role of PMR4 in different biotic and abiotic contexts.

Here, we report that callose deposition is induced in root hairs under Pi deficiency, in a manner independent of *PHR1/PHL1* and *LPR1/LPR2*. Our genetic studies revealed that the Pi depletion-induced callose deposition is impaired in *pmr4* (*gsl5*) root hairs.

The loss of *PMR4*-mediated callose deposition is associated with reduced plant growth as well as exaggerated stress symptoms under Pi deficiency, in addition with enhanced powdery mildew resistance. The results indicate *PMR4*-mediated callose deposition in root hairs as an important step in plant adaptation to Pi-limiting conditions.

Results

Callose deposition occurs in root hairs under phosphate deficiency, independently of *PHR1/PHL1* and *LPR1/LPR2*

Under low Pi conditions (Pi at 0 or 50 µM), we noticed callose deposition in the epidermal cells, particularly in root hairs of the differentiation zone (Figure 1a, b; O'Lexy et al., 2018). However, callose deposition was not detected in root hairs under the tested N, Fe or Fe/Pi deficient conditions (Figure 1a). The results indicate that the callose deposition is specific to Pi deficiency and is dependent on Fe. The absence of callose staining in the epidermis of root hair-less *rhd6* mutants (Figure 1c) validated that callose deposition occurs in root hairs under low Pi. In contrast to previous study (Müller et al., 2015), we did not detect callose deposition in the root SCN and cortex cells of the elongation zone under our experimental conditions (Figure 1a, c).

We next assessed whether *PHR1/PHL1* and *LPR1/LPR2* play a role in the root hair callose deposition. Unlike *LPR1*/*LPR2*-mediated callose deposition in SCN (Müller et al., 2015), the root hair callose deposition was not affected in *lpr1 lpr2* (Figure 1c). Furthermore, wild type (WT)-like callose deposition was retained in root hairs of *phr1 phl1 lpr1 lpr2* under low Pi conditions (Figure 1c). The results indicate that callose deposition is induced in root hairs under Pi deficiency, independently of these PSR regulators and callose deposition in the root SCN.

PMR4 callose synthase contributes to callose deposition under Pi deficiency

To unravel genetic requirements for the callose deposition in root hairs under Pi deficiency, we examined *glucan synthase like* (*gsl*) mutants for alternations in callose deposition under phosphate deficiency. Of the tested *gsl* mutants, *gsl5* (hereafter, *pmr4*) mutants specifically lacked callose deposition in root hairs (Figure S1). We also screened a total of 18,550 ethyl methane sulfonate (EMS)-mutagenized individuals in Col-0 and Ler backgrounds. This revealed two mutants in the Col-0 background, designated *callose alteration under Pi starvation 1 and 2* (*caps1 and caps2*, Figure 2a). *caps1* and *caps2* mutants were severely defective in callose deposition of root hairs under low Pi conditions (Figure 2a). DNA Sequencing on the *PMR4* genomic region revealed a DNA substitution within the coding region, resulting in a non-synonymous amino acid substitution, in both *caps1* and *caps*2 alleles (Figure 2b). The substitution in *caps*1 converts Tryptophan to Stop codon at the 260th amino acid (PMR4^{W260*}), resulting in PMR4 truncation, consistent with the severely impaired callose deposition in *caps1* as well as *pmr4-1* (PMR4^{W687*}). In *caps2*, glutamic acid is substituted with lysine at the 1181st amino acid (PMR4^{E1181K}), located within the cytosolic side of a predicted glucan-synthase domain (residue 575 to 1558, orange-marked domain, Figure 2b). The results suggest that dysfunction of PMR4 fails to induce root hair callose deposition under Pi deficiency in these *caps* alleles. We validated that leaf callose deposition in response to the bacterial MAMP flg22, a *PMR4*-dependent output in pattern-triggered immunity (PTI), is abolished in these *caps* alleles (Figure S2). Moreover, we showed strong resistance to powdery mildew fungi in these *caps* plants

(Figure 2c), as previously described in *pmr4* (Vogel & Somerville, 2000; Nishimura et al., 2003). The results confirm that PMR4 functions are impaired in both *caps* alleles.

PMR4 influences root development traits induced under Pi deficiency

By using *pmr4-1* as a representative mutant allele, we further investigated the functional significance of *PMR4* under Pi deficiency. The loss of *PMR4* leads to an increase in salicylic acid (SA) biosynthesis and defense under biotic stress (Vogel & Somerville, 2000; Nishimura et al., 2003), in a manner requiring *SID2 isochorismate synthase* that mediates a critical step in SA biosynthesis during defense activation (Wildermuth et al., 2001). Given the negative impact of SA defense activation on plant growth (Mateo et al., 2006; Zhang et al., 2010), we assessed the possible SA dependence of the observed *pmr4* phenotypes, by examining genetic interactions between *pmr4* and *sid2*. Callose deposition was induced in *sid2* root hairs but not induced in *pmr4 sid2* under low Pi conditions (Figure 3a), demonstrating that *SID2*-mediated SA biosynthesis is not required for *PMR4*-mediated callose deposition in root hairs. The results also exclude over-activation of SA defense as the cause of the callose defects in *pmr4* root hairs.

In Arabidopsis, root hair cells specifically occur at the junction of two underlying cortical cells, whereas non-hair cells occur in the other epidermal cells (Dolan et al., 1994; Schiefelbein et al., 2009). Root hair density was increased under low Pi conditions compared to normal Pi conditions, and the increase was exaggerated in *pmr4* as well as *pmr4 sid2*, without ectopic root hair formation (Figures 3b-d, S3a). Likewise, root hair length was also increased in *pmr4* as well as *pmr4 sid2* under both Pi conditions (Figure 3c, d). The results suggest that *PMR4* is not required for the

differentiation or outgrowth of root hair cells, but that *PMR4* contributes to restricting root hair formation and elongation, in the presence or absence of *SID2*. Conversely, primary root length was reduced in *pmr4* compared with WT in both low and normal Pi conditions (Figure 3b, e). We then determined the ratio of the primary root length under normal Pi versus low Pi as an indicator for the primary root growth inhibition under Pi deficiency. The ratio (normal Pi/low Pi) was increased in *pmr4* compared with WT, indicating that primary root growth inhibition, a Pi deficiency-characteristic response, is exaggerated in *pmr4* (Figure 3f). We also observed a decrease in the root meristem region in *pmr4* compared with WT under Pi deficiency, suggesting that the meristem exhaustion coincides with the inhibition of primary root growth in *pmr4* (Figure S3b). The primary root growth inhibition under low Pi was alleviated in *sid2*, but it was again exaggerated in *pmr4 sid2* compared to *sid2* (Figure 3e, f), indicating that the loss of *PMR4* enhances the Pi deficiency response, with or without *SID2*.

Since the primary root growth inhibition under Pi deficiency relies on *LPR1* and *LPR2* (Svistoonoff et al., 2007; Müller et al., 2015), we examined the possible relationship between *LPR1/LPR2* and *PMR4* in the regulation of primary root growth. The primary root length was reduced in *pmr4 lpr1 lpr2* compared with *lpr1 lpr2* under both Pi conditions, whereas it was increased in *pmr4 lpr1 lpr2* compared with *pmr4* specifically under low Pi (Figure S4a, b). The results indicate that *PMR4* serves to sustain the primary root growth irrespective of the Pi availability or *LPR1/LPR2* activity, whereas the *LPR1/LPR2* inhibition is specific under low Pi with or without *PMR4*. The ratio of the primary root length under normal Pi versus low Pi was lowered to a similar extent in *lpr1 lpr2* and *pmr4 lpr1 lpr2* compared with WT (Figure S4c), despite the

decrease in the primary root length of *pmr4* (Figure S4b), indicating that *lpr1 lpr2* is epistatic to *pmr4*.

We next determined the convex hull root area as a proxy of overall root system growth, which was correlated with Pi availability (Figure 3g). It was reduced in *pmr4* and *sid2* compared with WT under both normal and low Pi conditions, albeit to a lesser extent in *sid2* than *pmr4* under low Pi (Figure 3g), suggesting that *PMR4* and *SID2* both contribute to expanding the root area. Notably, only in the presence of *SID2*, the root area was reduced under both Pi conditions and its ratio under normal Pi versus low Pi was increased, in *pmr4* relative to WT (Figure 3g, h). The results suggest that *PMR4*-mediated expansion of the root foraging area relies on *SID2*. Conversely, the root area was greater in *pmr4 sid2* compared to *pmr4* under low Pi (Figure 3g), suggesting that *SID2* negatively impacts root expansion specifically in the absence of *PMR4* under Pi deficiency. There results thus suggest that *PMR4* directs *SID2*-mediated SA biosynthesis to root growth expansion in Pi-depleted plants, consistent with a previously described uncoupling of SA induction from defense activation under Pi deficiency (Gulabani et al. 2021).

PMR4 serves to sustain shoot growth and restrict anthocyanin accumulation under Pi deficiency.

We next examined whether *PMR4* influences shoot growth, by determining shoot fresh weight (FW) under Pi deficiency and sufficiency. Shoot FW was lowered in *pmr4* compared with WT, and it was not even partially restored by the simultaneous loss of *SID2* in *pmr4 sid2* under low-Pi media or soil (Figures 3i-j and S5). Together with the root phenotypes described above, the results exclude *SID2* as the major cause of the

growth inhibition in *pmr4*. The results suggest that *PMR4* positively influences shoot growth in the presence of *SID2*. Although high SA accumulation was previously described to reduce plant growth (Miura et al., 2010), we did not detect negative impacts of *SID2* on shoot growth under low Pi conditions (Figure 3j). *SID2* even positively influenced shoot growth under the normal Pi conditions, indicated by the reduced shoot biomass in *sid2* compared to WT (Figure 3j).

Anthocyanin content was increased in *pmr4* compared with WT, in the presence or absence of *SID2* (Figure 3k, l). Although SA was described to enhance anthocyanin accumulation under Pi deficiency (Morcillo et al., 2019), *sid2* was largely indistinguishable from WT in the anthocyanin contents under our low Pi conditions (Figure 3k, l). The results imply that PMR4-mediated Pi acquisition and allocation serves to alleviate Pi starvation causing anthocyanin accumulation under Pi deficiency in a manner independent of *SID2*.

PMR4 promotes the acquisition and root-to-shoot translocation of Pi

The *PMR4* effects on shoot and root growth prompted us to examine its possible role in Pi acquisition from the rhizosphere and Pi mobilization within the plant. We found that shoot Pi concentrations were greatly reduced in *pmr4* compared with WT, without alterations in root Pi concentrations, under Pi deficiency (Figure 4a, b). This indicates that *PMR4* contributes to shoot Pi accumulation in Pi-depleted plants. It was described that *PHR1/PHL1* promotes shoot Pi accumulation (Wang et al., 2018) and that *PHO1* Pi transporter mediates Pi mobilization from roots to shoots through the root xylem vessels (Stefanovic et al., 2007). We validated that Pi contents were lowered in the shoots but increased in the roots, in *phr1 phl1* and *pho1* compared with WT under low Pi (Figure

4c). A decrease in shoot Pi contents was also observed in *phr1 phl1* and *pho1* under normal Pi (Figure 4d). Since root hairs contribute to Pi accumulation under Pi deficiency (Tanaka et al., 2014; Robertson-Albertyn et al., 2017; Holz et al., 2018), we next determined Pi contents in a root hair less mutant, *rhd6* (Figure 4c, d). As seen in *pmr4, phr1 phl1* and *pho1*, Pi contents were lowered in the shoots but not in the roots of *rhd6* compared to WT, specifically under Pi deficiency (Figure 4c, d). As illustrated in Figure 4e, all these mutants displayed a decrease in shoot Pi contents without affecting or reducing root Pi contents under Pi deficiency. We inferred from these results that *PMR4*-mediated callose deposition in root hairs contributes to shoot Pi accumulation, possibly through increasing the uptake or root-to-shoot translocation of Pi. To test the hypothesis, we determined the rates of Pi uptake and Pi translocation by applying and tracing radio-labelled P. The rate of ^{32}P uptake per time and the ratio of ^{32}P in the shoot to whole seedling, standing for Pi translocation rate, were both lowered in *pmr4* and *pmr4 sid2* compared to WT and *sid2*, respectively, under low Pi (Figure 4f, g). In the presence of *SID2*, *PMR4* contribution to the two proxies was significant only under low Pi (Figure 4f, g). In the absence of *SID2*, however, *PMR4* contribution to P uptake was apparent under normal Pi as well (Figure 4f). Moreover, the ^{32}P uptake rate was reduced in *sid2* compared to WT, whereas *pmr4 sid2* was not distinguishable from *pmr4* under low Pi (Figure 4f), suggesting that SID2 positively influences Pi uptake under Pi deficiency in the presence of PMR4. These results suggest that PMR4 promotes Pi uptake and root-to-shoot Pi translocation under Pi deficiency and that this is in part achieved through diverting SID2 to promote these processes.

We further assessed the possible *PMR4* contributions to the acquisition and tissue mobilization of Pi and other elements, by inductively coupled plasma mass

spectrometry (ICP-MS) analyses for P, K, Mg, Ca, Mn, Fe, Zn, Mo and B contents in Pi-depleted and Pi-replete plants. Under Pi sufficiency, although the contents of these elements were not distinguishable between WT and *pmr4* roots, the contents of K and the rest were increased and lowered, respectively, in *pmr4* compared to WT shoots (Figure S6). Under low Pi, Mn, Zn and Mo contents were greater in *pmr4* compared with WT roots, whereas the contents of all the tested elements except K were lowered in *pmr4* compared with WT shoots (Figure 5). The results validated the role for *PMR4* in shoot P accumulation without affecting root P contents under both Pi conditions (Figures 4a-b, 5, S6). Additionally, the results suggest that *PMR4* also positively influences shoot accumulation of Mg, Ca, Mn, Fe, Zn, Mo and B (Figures 5, S6). WT-like and increased K accumulation in *pmr4* shoots compared to WT under Pi deficiency and sufficiency, respectively, show that the root-to-shoot transport is not entirely reduced for all elements in *pmr4*, pointing to a certain degree of element specificity in *PMR4* dependence (Figures 5, S6).

The role of *PMR4* in the shoot accumulation of different elements prompted us to examine whether *PMR4* influences plant responses to different nutrient deficiency. We examined *pmr4* phenotypes under B scarcity, as B scarcity induces inhibition of the primary root and transcriptional reprogramming related to ROS and defense in roots (Miwa et al., 2013; Das & Purkait, 2020). Although the primary root length was reduced in *pmr4* compared to WT over the tested range of B conditions, its ratio under B sufficiency (100 µM) versus the examined low B conditions was not affected in *pmr4* (Figure S7). The results point to specific contribution of *PMR4* to plant growth under Pi deficiency.

Root *PMR4* contributes to shoot growth under Pi deficiency

We next examined the possible significance of the root *PMR4* pool in promoting shoot growth, through micrografting between WT and *pmr4* plants. We validated that the micrografted plants of WT/WT (shoot/root genotypes) and *pmr4/pmr4* showed the corresponding WT and *pmr4* phenotypes in root growth and SFW (Figure 6a-c). The root hair callose was detected only when *PMR4* was present in the roots, although the callose signals were not as strong as in intact plants, possibly reflecting the stresses imposed by the grafting procedure (Figure 6d). Remarkably, SFW was greatly reduced in WT/*pmr4* whereas it was less affected in *pmr4*/WT (Figure 6b), pointing to a critical role for a root *PMR4* pool in shoot growth under Pi deficiency. However, the root growth was partially restored in WT/*pmr4* (Figure 6c), pointing to a role for a shoot *PMR4* pool in sustaining the primary root growth in Pi-depleted plants.

To further examine the role of a PMR4 pool in root hairs, we generated transgenic plants expressing *PMR4* under the control of a root hair-specific *LRX1* promoter (*pLRX1*, Baumberger et al., 2003) in the *pmr4-1* background (Figure 7a, b). *pLRX1::PMR4* complemented *pmr4* in terms of the callose deposition in root hairs and shoot growth under low Pi conditions (Figure 7a-c). By contrast, the primary root growth remained suppressed in the *pLRX1::PMR4* transgenic lines as well as in *pmr4* (Figure 7b, d). The results indicate that PMR4 in root hairs is sufficient to restore shoot growth, but not the primary root growth, under Pi deficiency.

Discussion

Our studies on callose deposition in root hairs induced under Pi deficiency revealed *PMR4* as a positive regulator of plant adaptation to Pi deficiency (Figure 8). *PMR4*

mediates defense-induced callose deposition in pattern triggered immunity and pre-invasion fungal resistance (Jacobs et al., 2003; Chowdhury et al., 2014), and that its loss is linked to enhanced SA-based pathogen resistance (Nishimura et al., 2003). Therefore, *PMR4* seems to play a critical role in an array of biotic and abiotic stress responses, possibly through callose deposition at different sites.

The present study suggests that the root hair callose contributes to root Pi uptake and shoot Pi accumulation under Pi deficiency. Importantly, the loss of *PMR4* affects both *PHR1*-dependent and *LPR1*-dependent processes (Figures 3, 4) (Bates & Lynch 1996; Thibaud et al., 2010), which have been genetically uncoupled in PSR (Wang et al., 2010; Balzergue et al., 2017). Pi accumulation and anthocyanin accumulation are suppressed in *phr1 phl1* (Rubio et al., 2001; Bustos et al., 2010; Wang et al., 2018). The shoot Pi contents, and plant biomass are also reduced in *pmr4*, implying that *PMR4* positively influences *PHR1/PHL1*-mediated Pi acquisition and plant growth under internal Pi deficiency. However, over-accumulation of anthocyanin in *pmr4* (Figure 3k, l) in contrast to *phr1 phl1* (Rubio et al., 2001; Bustos et al., 2010; Wang et al., 2018), suggests that *PMR4* contribution is specific to the former two outputs of *PHR1/PHL1* and that its loss results in exaggeration of Pi starvation symptom. The loss of *PMR4* also strengthens inhibition of the primary root growth under Pi deficiency, only in the presence of *LPR1/LPR2* (Figure S4). The LPR1 pathway mediates callose deposition at the root SCN to inhibit the cell-to-cell movement of the transcription factor SHR, thereby suppressing the cell division and subsequent root elongation (Müller et al., 2015; Mora-Macías et al., 2017). *PMR4*-mediated retention of primary root growth is prominent only in the presence of *LPR1/LPR2*, whereas *LPR1/LPR2*-mediated growth inhibition is significant even in the absence of *PMR4* under low Pi (Figure S4). Our

results suggest a role for PMR4 in antagonizing LPR1/LPR2-dependent root growth inhibition.

The failure of root hair PMR4 to restore primary root growth (Figure 7b, d) implies the involvement of PMR4 from another cell type(s) than root hairs in this regulation. Consistently, shoot Pi accumulation is partially and essentially retained in *rhd6* under low and normal Pi conditions, respectively, despite the lack of root hairs (Figure 4c-e). Although our callose detection was limited to root hairs, it is conceivable that PMR4-mediated callose deposition in different cell types serves to sustain the primary root growth against LPR1/LPR2-mediated attenuation following a decrease in external Pi levels. The mechanisms underlying antagonistic interactions between *PMR4* and *LPR1/LPR2* require further studies including the identification of the SCN callose synthase.

Shoot Pi accumulation depends on root Pi uptake and subsequent Pi translocation. *PHT1;1* and *PHT1;4* provide two major Pi transporters in roots, contributing greatly to Pi acquisition under Pi deficiency (Shin et al., 2004). *PHO1* mediates Pi translocation from roots to shoots through Pi loading into the root xylem (Poirier et al., 1991; Hamburger et al., 2002; Liu et al., 2012). Notably, the loss of *PMR4* results in a decrease in the shoot Pi contents without affecting the root Pi contents (Figures 4a-b, 5). Likewise, the loss of *PHR1/PHL1* and *PHO1* also lowers shoot Pi contents without reducing root Pi contents (Figure 4c, d; Liu et al., 2012; Wang et al., 2018). In *pht1;1 pht1;4*, a decrease of Pi uptake in roots is associated with that of Pi translocation to the shoots and shoot Pi accumulation (Shin et al., 2004). An intimate linkage between root uptake and root-to-shoot translocation is also seen for K^+ (Nieves-Cordones et al., 2019). It is thus conceivable that the lowered Pi accumulation in *pmr4* shoots is caused by a

decrease in the rates of the root Pi uptake and the root-to-shoot Pi translocation, possibly through the compromised root hair function in the absence of *PMR4*. Indeed, ^{32}P tracer analyses indicate that *PMR4* promotes both root Pi uptake and root-to-shoot Pi translocation, without affecting the root Pi contents (Figure 4a, b, f, g). This is consistent with the view that the root-to-shoot Pi translocation is driven by the Pi uptake in the roots (Shin et al., 2004). Leaf transpiration affects root-to-shoot solute translocation (Nieves-Cordones et al., 2019), but it might be ignorable under high humidity in the closed plate assays. Notably, *PMR4* also contributes to the accumulation of different elements except K in shoots under Pi deficiency (Figures 5, S6). This might largely attribute to secondary effects of *PMR4*-dependent root growth and expansion (Figure 3e-h), but the separate regulation of K (Figures 5, S6) cannot be explained by this model. It was previously described that the disruption of multiple *PHT1* members (*PHT1;1/1;2/1;3* silencing in the *phf1 pht1;4* mutant background, designated mut5) results in a decrease in Na as well as P contents compared to WT (Ayadi et al., 2015). Cross-regulation among Fe, Zn and Pi homeostasis relies on *PHR1* (Briat et al., 2015). Further studies will be required to unravel the mechanisms underlying the complex coordination of the acquisition and tissue distribution among different elements and the observed role of PMR4 in the regulation (Figures 5, S6).

Root hairs increase the root surface area to promote root anchorage, nutrient acquisition, and metabolite exudation, thereby facilitating the interactions with the soil environments (Datta et al., 2011; Poole et al., 2018; Vissenberg et al., 2020; Robe et al., 2021; Han et al., 2023). The elongation and increased densities of root hairs in the absence of PMR4-mediated callose deposition therein (Figure 3c, d) can be inferred that the root hairs formed in *pmr4* are less functional in Pi acquisition or Pi allocation to the

other tissues, thereby leading to further formation of root hairs. Our findings point to a positive role for *PMR4* in the root hair functions under low Pi. The mechanisms by which callose deposited in the root hairs promotes Pi acquisition require further investigation.

Root hairs also provide a key interface to soil-borne microbes and are intimately associated with root defense regulation. Root hair formation and elongation are facilitated by jasmonate (JA) (Zhu et al., 2006), which mediates the expression of key transcription factors for root hair differentiation and the de-repression of RHD6 from the interference by JAZ proteins (Han et al., 2020). In beneficial interactions with the epiphytic bacterium *Pseudomonas simiae* WCS417, activation of defense-related genes is increased in hair cell lineages compared to non-hair cell lineages (Verbon et al., 2023). The endogenous elicitor-active Pep peptides induce root hair formation (Okada et al., 2021; Jing et al., 2024), and Pep receptor signaling from the root hairs can activate defense responses without root growth inhibition (Okada et al., 2021). These findings suggest a specific capacity of root hair cells in root immunity. How these properties of root hairs are influenced by *PMR4*-mediated callose deposition under Pi deficiency also represents an interesting topic for future studies.

Experimental procedures

Plant materials and growth conditions

The *Arabidopsis thaliana* accessions used were Col-0 and Ws-2, including the mutants examined (Table S1) and the wild type used was Col-0 unless otherwise stated. We generated ethyl methanesulfonate (EMS)-mutagenesis (M2) seeds for screening in both Col-0 and Ler backgrounds. All seeds were sterilized using 6% sodium hypochlorite

and 0.2% Trion X-100 for 15 min, and then kept at 4°C for 2 d before sowing a half-strength Murashige and Skoog (MS) medium with 25 mM sucrose and 0.8% Agar (WAKO, Japan). If not stated otherwise, the normal Pi media contained 625 μM KH_2PO_4, whereas the low Pi media contained 0 μM, 10 μM, 50 μM or 150 μM KH_2PO_4 as indicated. The low N and low Fe media used contained 0 μM N and 0 μM Fe, respectively. Plants were grown in a growth chamber under 12 h: 12 h, light: dark cycles at 22°C. Six-day (d)-old seedlings were transferred to the indicated Pi liquid media or agar media with 1% Difco Agar, Bacteriological (BD, Franklin Lakes, NJ, USA).

Construction and plant transformation

The genomic DNA sequences of the *POWDERY MILDEW RESISTANCE 4* (*PMR4*, AT4G03550) loci, together with a regulatory DNA sequence for root hair specific expression (below), were cloned into the Gateway binary vector pGWB504 (Nakagawa et al., 2007; primer information Table S2) to express a C-terminally GFP-fused PMR4 under its native regulatory DNA sequence. The root hair-specific promoter sequence used was derived from *LRR-EXTENSIN 1* (*LRX1*, AT1G12040) (Baumberger et al., 2003). Transgenic lines were generated by Agrobacterium-mediated transformation of *pmr4-1* with the DNA construct and subsequently selected on hygromycin resistance and DNA genotyping.

Aniline blue and PI staining

Callose staining with aniline blue was conducted as described in Schenk & Schikora (2015). Plants were examined at 3 d after the transfer of 5-d-old seedlings to the

indicated Pi, N or Fe liquid media (Figure 1a, c). Plant tissues were fixed in a solution containing acetic acid and ethanol (1:3 v/v) overnight. The fixation solution was exchanged to a 150 mM K_2HPO_4 solution before callose staining with 0.1% aniline blue in the K_2HPO_4 solution for 2 h. The data presented are representative of three independent experiments (Figures 1a, 1c, 2a, 3a), two independent experiments (Figures 1b, 6d, S1, S2) and one experiment (Figures 7a, S3). To localize the root meristematic regions, the primary roots were stained with 20 μg/ml propidium iodide (PI). Callose or PI signals were examined under microscopy using a fluorescence microscope DMI 6000B (Leica Microsystems, Wetzlar, Germany) and a confocal microscope SP8 (Leica Microsystems, Wetzlar, Germany).

Identification of mutants altered in the root hair callose deposition

We examined callose deposition in *gsl* mutants (Table S1) after aniline blue staining of the roots transferred to the low Pi liquid media (50 μM) for 3 d. In the EMS mutagenic screening, we subjected a region ~5 cm from the root tip to aniline blue staining, after cutting out of the M2 individuals transferred to the Pi -depleted liquid media (0 μM) for 3 d. This enabled to keep the plant alive for subsequent characterization and seed propagation. The *PMR4* genomic region was sequenced in the selected mutant candidates.

Inoculation analyses with powdery mildew fungi

Inoculation of *Golovinomyces orontii* and determination of fungal versus plant DNA ratio were conducted essentially by following Weßling & Panstruga (2012). Eighteen-d-old plants grown in 10 h: 14 h, light: dark cycles at 22°C were inoculated

with *G. orontii* spores at approximately 750 spores cm^{-2}. The inoculated leaves were harvested and frozen in liquid nitrogen before determining fungal biomass at 10 d after inoculation. Genomic DNA was extracted essentially by following Brouwer et al. (2001). Quantitative real-time PCR was performed with primers (Table S2) and Brilliant Sybr Green QPCR Reagent Kit (Stratagene, Waldbronn, Germany). The ratio of *G. orontii* to Arabidopsis genomic DNA was calculated using the ΔΔCt method (Pfaffl, 2001). Data are representative of two independent experiments (Figure 2c).

Determination of root hair density, primary root length and root area

Six-day-old seedlings were transferred and further grown in the low (50 μM) or normal Pi media (625 μM) for 6 d, and then root hairs, primary root and root system area were examined under stereomicroscope M80 (Leica Microsystems, Wetzlar, Germany). The number of root hairs was determined in a region 2.0-3.0 mm from the root tip using Image J software (National Institutes of Health, Bethesda, MD, USA). Data are representative of similar results from three independent experiments (Figure 3c, d) and two independent experiments (Figure S4). The primary root length and convex hull area were determined using ImageJ software and Root nav software (Pound et al., 2013). Data are representative of three independent experiments (Figure. 3e-h). For growth under different Boron (B) conditions, plants were grown in MGRL solid media containing 1% sucrose and 1% gellan gum (WAKO, Japan). B concentrations were adjusted with boric acid. Plants were grown under 16 h: 8 h, light: dark cycles at 22°C. The data presented are from one independent experiment (Figure S7).

Determination of shoot fresh weight, anthocyanin and Pi contents

In plant growth assays on media, six-day-old seedlings were transferred to the indicated media (10 μM, 50 μM or 625 μM Pi) and then further grown for the indicated days. Shoot fresh weight (SFW) was determined 16 d after transfer to the indicated media (50 μM or 625 μM Pi; Figure 3i, j). In plant growth assays with *pLRX1::PMR4* lines, SFW was determined 10 d after the transfer to the low Pi media (10 μM ; Figure 6a, c). In plant growth assay on soil, organic black soil (Maruishi Engei, Osaka, Japan) and Ezo sand (Kobayashi Sangyo, Kita-Nagoya, Japan) mixed at 1:1 (v/v) were used as low Pi soil, and Sakata-Super-Mix-A (Sakata Seed, Kyoto, Japan) and Vermiculite G20 (Maruishi Engei, Osaka, Japan) mixed at 1:1 (v/v) were used as normal Pi soil (Figure S5). SFW of seedlings grown on soil was determined 14 d after the transfer of 10-d-old seedlings to the low and normal Pi soils. Data are representative of two independent experiments (Figures 3i, 3j, S5) and one independent experiment (Figure 7b, c).

In the determination of anthocyanin content, six-day-old seedlings were transferred to and further grown on low Pi media (10 μM) for 10 d (Figure 3k, l). The shoots were subject to the procedure described in Mita et al. (1997). In brief, a HCl: methanol mixture at 1:99 (v/v) was added to the ground tissues with 1 ml per sample, and then the resultant homogenates were incubated at 4°C overnight. A_{530} and A_{657} of the supernatants were determined with a spectrometer UV-1800 (Shimadzu, Kyoto, Japan). Anthocyanin content was calculated following the formula: $(A_{530} - 0.25 \times A_{657}) \times$ fresh weight $(FW)^{-1}$. Data are representative of two independent experiments with the same conclusions (Figure 3k, l).

In the determination of Pi content, six-d-old seedlings were transferred to and further grown on the low Pi (150 μM) or normal Pi (625 μM) media for 9 d (Figure 4a-d). Pi content was determined by following Ames (1966). In brief, the extraction buffer (10

mM Tris, 1 mM EDTA, 100 mM NaCl, 1 mM phenylmethylsulfonyl fluoride pH8.0) was added with 10 µl per sample mg FW. The homogenate 100 µl was mixed with 1% glacial acetic acid 900 µl, and then incubated at 42°C for 30 min. The supernatant was recovered to be incubated in a solution (0.35% NH_4MoO_4, 0.86 N H_2SO_4, and 1.4% ascorbic acid) at 42°C for 30 min. A_{820} of the aliquot was measured under Eppendorf BioSpectrometer® basic (Eppendorf, Hamburg, Germany). Then, Pi content was calculated by a P calibration curve prepared with a P standard solution. Data are representative of three independent experiments (Figure 4a, b) and two independent experiments (Figure 4c, d) with the same conclusions.

Scanning electron microscope

The maturation zone of primary roots 10 d after transfer of 5-d-old seedlings to low Pi (50 µM) liquid media was fixed in half-Karnovsky's fixative overnight at 4°C, and then postfixed in 1% osmium tetroxide for 1 h at room temperatures (Karnovsky, 1965). The specimens were dehydrated in an ethanol series (30%, 50%, 70%, 80%, and 90%, and 100% at three times) for 10 min each at RT, and then the specimens were dipped in t-butyl alcohol and dried with a vacuum evaporator. They were coated with vapor of 1% osmium tetroxide and then observed with a scanning electron microscope (JIB-4601F, JEOL, Tokyo, Japan).

Inductively coupled plasma-mass spectrometer (ICP-MS) analyses

Five-day-old seedlings were transferred to and further grown on the growth media (10 µM or 625 µM KH_2PO_4) for 4 d (Figures 5, S6). Shoots and roots were harvested and dried at 60°C. After determining the dry weight, the dried tissues were digested with 2

ml nitric acid and 0.5 ml hydrogen peroxide (Wako Pure Chemicals) at 110°C. Digested samples were then dissociated with 5 ml of 2% (w/v) nitric acid and subjected to inductively coupled plasma mass spectrometry (NexION 2000, Perkin-Elmer). The contents of P, K, Mg, Ca, Mn, Fe, Zn, Mo and B were determined. The data were obtained from one experiment (Figures 5, S6).

Determination of ^{32}P uptake rate and translocation ratio

Seven-day-old seedlings were transferred to and further grown on the growth media (50 or 625 μM KH_2PO_4) for 10 d. The seedlings were exposed to the liquid media containing ^{32}P-labeled Pi ($^{32}PO_4^{3-}$ 20 kBq / ml) for 30 min. The plant tissues were fixed to an Imaging Plate (GE Healthcare co., Chicago, USA) for 3 h and then scanned by Typhoon FLA 9000 (GE Healthcare co., Chicago, USA). ^{32}P uptake rate was calculated based on a calibration curve for ^{32}P. For determining the Pi translocation rate into shoots, the seedlings were exposed to the Pi-containing liquid media containing ^{32}P-labeled Pi for 10 min, and then transferred to non-labelled agar media for 120 min. The translocation rate of Pi was determined by calculating ^{32}P ratio of the shoot to the whole seedlings. The data presented are representative of two independent experiments with the same conclusions (Figure 4f, g).

Grafting experiments

Grafting of WT and *pmr4* was performed with a supportive micro-device as described in Tsutsui et al. (2020). Four-day-old seedlings grown on a 1/2 MS medium containing 0.5% sucrose and 1% agar under constant light at 22°C were subject to grafting. Both hypocotyls of stock and scion were cut horizontally with a knife and assembled on the

device using forceps. The grafted plants were grown in a 1/2 MS medium containing 0.5% sucrose and 2% agar at 27°C for 6 d. In callose staining, the grafted plants transferred and grown on low Pi media (50 μM) under constant light at 22°C for 9 d were examined. In plant growth assays, the grafted plants transferred and grown on low Pi media (150 μM) under constant light at 22°C for 14 d were examined. The data presented are representative of two independent experiments with the same conclusions (Figure 6).

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Author contributions

YS conceived the study. KO and YS designed the experiments. KO, KY, TANN, SK, SY, HT, CT, THL, UN, KI, KM, TM and MN developed the methods, performed the experiments, and/or analyzed the data. SM, MN and KH advised on the experiments. KO and YS wrote the manuscript with contributions from the other authors.

Supporting Information

Figure S1. Callose deposition in *gsl* mutants under phosphate deficiency.

Figure S2. flg22-induced callose deposition is impaired in the *caps* and *pmr4* alleles.

Figure S3. Meristematic region is reduced in *pmr4* mutants under phosphate deficiency.

Figure S4. *PMR4* restricts *LPR1/LPR2*-mediated inhibition of the primary root growth in response to low Pi.

Figure S5. Plant growth is lowered in *pmr4* on soil.

Figure S6. Plant element contents in shoot and root under phosphate sufficiency.

Figure S7. Primary root growth inhibition under B deficiency.

Table S1. *Arabidopsis thaliana* mutants used in this study.

Table S2. DNA Primers used in DNA vector construction and qPCR analyses.

References

Ames, B.N. (1966) Assay of inorganic phosphate, total phosphate and phosphatases. *Methods Enzymol*, **8**, 115–118.

Ayadi, A., David, P., Arrighi, J.F., Chiarenza, S., Thibaud, M.C., Nussaume, L. and Marin, E. (2015) Reducing the genetic redundancy of Arabidopsis phosphate transporter1 transporters to study phosphate uptake and signaling. *Plant Physiol*, **167**, 1511–1526.

Balzergue, C., Dartevelle, T., Godon, C., et al. (2017) Low phosphate activates STOP1-ALMT1 to rapidly inhibit root cell elongation. *Nat Commun*, **8**.

Bates, T.R. and Lynch, J.P. (1996) Stimulation of root hair elongation in *Arabidopsis thaliana* by low phosphorus availability. *Plant Cell Environ*, **19**, 529–538.

Baumberger, N., Ringli, C. and Keller, B. (2001) The chimeric leucine-rich repeat/extensin cell wall protein LRX1 is required for root hair morphogenesis in *Arabidopsis thaliana*. *Genes Dev*, **15**, 1128–1139.

Briat, J.F., Rouached, H., Tissot, N., Gaymard, F. and Dubos, C. (2015) Integration of P, S, Fe, and Zn nutrition signals in *Arabidopsis thaliana*: Potential involvement of PHOSPHATE STARVATION RESPONSE 1 (PHR1). *Front Plant Sci*, **6**.

Brouwer, M., Lievens, B., Hemelrijck, W. Van, Ackerveken, G. Van Den, Cammue, B.P.A. and Thomma, B.P.H.J. (2003) Quantification of disease progression of several microbial pathogens on *Arabidopsis thaliana* using real-time fluorescence PCR. *FEMS Microbiol Lett*, **228**, 241–248.

Bustos, R., Castrillo, G., Linhares, F., Puga, M.I., Rubio, V., Pérez-Pérez, J., Solano, R., Leyva, A. and Paz-Ares, J. (2010) A central regulatory system largely controls transcriptional activation and repression responses to phosphate starvation in Arabidopsis. *PLoS Genet*, **6**.

Chan, C., Liao, Y.Y. and Chiou, T.J. (2021) The Impact of Phosphorus on Plant Immunity. *Plant Cell Physiol*, **62**, 582–589.

Chien, P.S., Chiang, C.P., Leong, S.J. and Chiou, T.J. (2018) Sensing and Signaling of Phosphate Starvation: From Local to Long Distance. *Plant Cell Physiol*, **59**, 1714–1722.

Chowdhury, J., Henderson, M., Schweizer, P., Burton, R.A., Fincher, G.B. and Little, A. (2014) Differential accumulation of callose, arabinoxylan and cellulose in nonpenetrated versus penetrated papillae on leaves of barley infected with *Blumeria graminis* f. sp. *hordei*. *New Phytologist*, **204**, 650–660.

Das, A.K. and Purkait, A. (2020) Boron dynamics in soil: classification, sources, factors, fractions, and kinetics. *Commun Soil Sci Plant Anal*, **51**, 2778–2790.

Datta, S., Kim, C.M., Pernas, M., Pires, N.D., Proust, H., Tam, T., Vijayakumar, P. and Dolan, L. (2011) Root hairs: development, growth and evolution at the plant-soil interface. *Plant Soil*, **346**, 1–14.

Dolan, L., Duckett, C.M., Grierson, C., Linstead, P., Schneider, K., Lawson, E., Dean, C., Poethig, S. and Roberts, K. (1994) Clonal relationships and cell patterning in the root epidermis of *Arabidopsis*. *Development*, **120**, 2465–2474.

Dong, X., Hong, Z., Sivaramakrishnan, M., Mahfouz, M. and Verma, D.P.S. (2005) Callose synthase (CalS5) is required for exine formation during microgametogenesis and for pollen viability in Arabidopsis. *Plant Journal*, **42**, 315–328.

Ellinger, D. and Voigt, C.A. (2014) Callose biosynthesis in arabidopsis with a focus on pathogen response: What we have learned within the last decade. *Ann Bot*, **114**, 1349–1358.

Enns, L.C., Kanaoka, M.M., Torii, K.U., Comai, L., Okada, K. and Cleland, R.E. (2005) Two callose synthases, GSL1 and GSL5, play an essential and redundant role in plant and pollen development and in fertility. *Plant Mol Biol*, **58**, 333–349.

Gulabani, H., Goswami, K., Walia, Y., et al. (2022) Arabidopsis inositol polyphosphate kinases IPK1 and ITPK1 modulate crosstalk between SA-dependent immunity and phosphate-starvation responses. *Plant Cell Rep*, **41**, 347–363. Available at: https://doi.org/10.1007/s00299-021-02812-3.

Hamburger, D., Rezzonico, E., Petétot, J.M.D.C., Somerville, C. and Poirier, Y. (2002) Identification and characterization of the Arabidopsis PHO1 gene involved in phosphate loading to the xylem. *Plant Cell*, **14**, 889–902.

Han, G., Wei, X., Dong, X., et al. (2020) Arabidopsis ZINC FINGER PROTEIN1 Acts Downstream of GL2 to Repress Root Hair Initiation and Elongation by Directly Suppressing bHLH Genes. *Plant Cell*, **32**, 206–225.

Han, X., Kui, M., He, K., Yang, M., Du, J., Jiang, Y. and Hu, Y. (2023) Jasmonate-regulated root growth inhibition and root hair elongation. *J Exp Bot*, **74**, 1176–1185.

Holz, M., Zarebanadkouki, M., Kuzyakov, Y., Pausch, J. and Carminati, A. (2018) Root hairs increase rhizosphere extension and carbon input to soil. *Ann Bot*, **121**, 61–69.

Isidra-Arellano, M.C., Delaux, P.M. and Vald°s-L°pez, O. (2021) The Phosphate Starvation Response System: Its Role in the Regulation of Plant–Microbe Interactions. *Plant Cell Physiol*, **62**, 392–400.

Jacobs, A.K., Lipka, V., Burton, R.A., Panstruga, R., Strizhov, N., Schulze-Lefert, P. and Fincher, G.B. (2003) An Arabidopsis Callose Synthase, GSL5, Is Required for Wound and Papillary Callose Formation. *Plant Cell*, **15**, 2503–2513.

Jing, Y., Zhao, F., Lai, K., et al. (2024) Plant elicitor Peptides regulate root hair development in *Arabidopsis*. *Front Plant Sci*, **15**.

Karnovsky, M.J. (1964) A Formaldehyde-Glutaraldehyde Fixative of High Osmolality for Use in Electron Microscopy. *The Journal of Cell Biology*, **27**, 1A-149A. https://www.researchgate.net/publication/244955881.

Liu, T.Y., Huang, T.K., Tseng, C.Y., Lai, Y.S., Lin, S.I., Lin, W.Y., Chen, J.W. and Chioua, T.J. (2012) PHO2-dependent degradation of PHO1 modulates phosphate homeostasis in *Arabidopsis*. *Plant Cell*, **24**, 2168–2183.

Mateo, A., Funck, D., Mühlenbock, P., Kular, B., Mullineaux, P.M. and Karpinski, S. (2006) Controlled levels of salicylic acid are required for optimal photosynthesis and redox homeostasis. *J Exp Bot*, **57**, 1795–1807.

Mita, S., Murano N., Akaike M. and Nakamura K. (1997) Mutants of *Arabidopsis thaliana* with pleiotropic effects on the expression of the gene for β-amylase and on the accumulation of anthocyanin that are inducible by sugars. *Plant Journal*, **11**, 841-851.

Miura, K., Lee, J., Miura, T. and Hasegawa, P.M. (2010) SIZ1 controls cell growth and plant development in Arabidopsis through salicylic acid. *Plant Cell Physiol*, **51**, 103–113.

Miwa, K., Wakuta, S., Takada, S., Ide, K., Takano, J., Naito, S., Omori, H., Matsunaga, T. and Fujiwara, T. (2013) Roles of BOR2, a boron exporter, in cross linking of rhamnogalacturonan II and root elongation under boron limitation in Arabidopsis. *Plant Physiol*, **163**, 1699–1709.

Mora-Macías, J., Ojeda-Rivera, J.O., Gutiérrez-Alanís, D., et al. (2017) Malate-dependent Fe accumulation is a critical checkpoint in the root developmental response to low phosphate. *Proc Natl Acad Sci USA*, **114**, E3563–E3572. https://doi.org/10.1073/pnas.1701952114

Morcillo, R.J., Singh, S.K., He, D., et al. (2020) Rhizobacterium□derived diacetyl modulates plant immunity in a phosphate□dependent manner. *EMBO J*, **39**, 1–15.

Müller, J., Toev, T., Heisters, M., Teller, J., Moore, K.L., Hause, G., Dinesh, D.C., Bürstenbinder, K. and Abel, S. (2015) Iron-Dependent Callose Deposition Adjusts Root Meristem Maintenance to Phosphate Availability. *Dev Cell*, **33**, 216–230.

Nakagawa, T., Suzuki, T., Murata, S., et al. (2007) Improved gateway binary vectors: High-performance vectors for creation of fusion constructs in transgenic analysis of plants. *Biosci Biotechnol Biochem*, **71**, 2095–2100.

Nieves-Cordones, M., Lara, A., Ródenas, R., Amo, J., Rivero, R.M., Martínez, V. and Rubio, F. (2019) Modulation of K^+ translocation by AKT1 and AtHAK5 in Arabidopsis plants. *Plant Cell Environ*, **42**, 2357–2371.

Nishikawa, S.I., Zinkl, G.M., Swanson, R.J., Maruyama, D. and Preuss, D. (2005) Callose (β-1,3 glucan) is essential for Arabidopsis pollen wall patterning, but not tube growth. *BMC Plant Biol*, **5**, 1–9.

Nishimura, M.T., Stein, M., Hou, B.H., Vogel, J.P., Edwards, H. and Somerville, S.C. (2003) Loss of a callose synthase results in salicylic acid-dependent disease resistance. *Science (1979)*, **301**, 969–972.

Okada, K., Kubota, Y., Hirase, T., Otani, K., Goh, T., Hiruma, K. and Saijo, Y. (2021) Uncoupling root hair formation and defence activation from growth inhibition in response to damage-associated Pep peptides in *Arabidopsis thaliana*. *New Phytologist*, **229**, 2844–2858.

O'Lexy, R., Kasai, K., Clark, N., Fujiwara, T., Sozzani, R. and Gallagher, K.L. (2018) Exposure to heavy metal stress triggers changes in plasmodesmatal permeability via deposition and breakdown of callose. *J Exp Bot*, **69**, 3715–3728.

Omasits, U., Ahrens, C.H., Müller, S. and Wollscheid, B. (2014) Protter: Interactive protein feature visualization and integration with experimental proteomic data. *Bioinformatics*, **30**, 884–886.

Pasternak, T., Groot, E.P., Kazantsev, F. V., Teale, W., Omelyanchuk, N., Kovrizhnykh, V., Palme, K. and Mironova, V. V. (2019) Salicylic acid affects root meristem patterning via auxin distribution in a concentration-dependent manner. *Plant Physiol*, **180**, 1725–1739.

Péret, B., Clément, M., Nussaume, L. and Desnos, T. (2011) Root developmental adaptation to phosphate starvation: Better safe than sorry. *Trends Plant Sci*, **16**, 442–450.

Péret, B., Desnos, T., Jost, R., Kanno, S., Berkowitz, O. and Nussaume, L. (2014) Root architecture responses: In search of phosphate. *Plant Physiol*, **166**, 1713–1723.

Pfaffl, M.W. (2001) A new mathematical model for relative quantification in real-time RT-PCR. *Nucleic Acids Research*, **29**, 2003-2007

Poirier, Y., Thoma, S., Somerville, C. and Schiefelbein, J. (1991) A mutant of *Arabidopsis* deficient in xylem loading of phosphate. *Plant Physiol*, **97**, 1087–1093.

Poole, J., Day, C.J., Itzstein, M. Von, Paton, J.C. and Jennings, M.P. (2018) Glycointeractions in bacterial pathogenesis. *Nat Rev Microbiol*, **16**, 440–452. http://dx.doi.org/10.1038/s41579-018-0007-2.

Pound, M.P., French, A.P., Atkinson, J.A., Wells, D.M., Bennett, M.J. and Pridmore, T. (2013) RootNav: Navigating images of complex root architectures. *Plant Physiol*, **162**, 1802–1814.

Robe, K., Conejero, G., Gao, F., et al. (2021) Coumarin accumulation and trafficking in *Arabidopsis thaliana*: a complex and dynamic process. *New Phytologist*, **229**, 2062–2079.

Robertson-Albertyn, S., Terrazas, R.A., Balbirnie, K., et al. (2017) Root hair mutations displace the barley rhizosphere microbiota. *Front Plant Sci*, **8**, 1–15.

Rubio, V., Linhares, F., Solano, R., Martín, A.C., Iglesias, J., Leyva, A. and Paz-Ares, J. (2001) A conserved MYB transcription factor involved in phosphate starvation signaling both in vascular plants and in unicellular algae. *Genes Dev*, **15**, 2122–2133.

Sager, R. and Lee, J.Y. (2014) Plasmodesmata in integrated cell signalling: Insights from development and environmental signals and stresses. *J Exp Bot*, **65**, 6337–6358.

Schenk, S.T. and Schikora, A. (2015) Staining of Callose Depositions in Root and Leaf Tissues. *Bio-protocol*, **5**. http://www.bio-protocol.org/e1429

Schiefelbein, J., Kwak, S.H., Wieckowski, Y., Barron, C. and Bruex, A. (2009) The gene regulatory network for root epidermal cell-type pattern formation in *Arabidopsis*. *J Exp Bot*, **60**, 1515–1521.

Shin, H., Shin, H.S., Dewbre, G.R. and Harrison, M.J. (2004) Phosphate transport in *Arabidopsis*: Pht1;1 and Pht1;4 play a major role in phosphate acquisition from both low- and high-phosphate environments. *Plant Journal*, **39**, 629–642.

Stefanovic, A., Ribot, C., Rouached, H., Wang, Y., Chong, J., Belbahri, L., Delessert, S. and Poirier, Y. (2007) Members of the *PHO1* gene family show limited functional redundancy in phosphate transfer to the shoot, and are regulated by phosphate deficiency via distinct pathways. *Plant Journal*, **50**, 982–994.

Sun, L., Song, L., Zhang, Y., Zheng, Z. and Liu, D. (2016) Arabidopsis PHL2 and PHR1 act redundantly as the key components of the central regulatory system controlling transcriptional responses to phosphate starvation. *Plant Physiol*, **170**, 499–514.

Svistoonoff, S., Creff, A., Reymond, M., Sigoillot-Claude, C., Ricaud, L., Blanchet, A., Nussaume, L. and Desnos, T. (2007) Root tip contact with low-phosphate media reprograms plant root architecture. *Nat Genet*, **39**, 792–796.

Tanaka, N., Kato, M., Tomioka, R., Kurata, R., Fukao, Y., Aoyama, T. and Maeshima, M. (2014) Characteristics of a root hair-less line of *Arabidopsis thaliana* under physiological stresses. *J Exp Bot*, **65**, 1497–1512.

Thibaud, M.C., Arrighi, J.F., Bayle, V., Chiarenza, S., Creff, A., Bustos, R., Paz-Ares, J., Poirier, Y. and Nussaume, L. (2010) Dissection of local and systemic transcriptional responses to phosphate starvation in Arabidopsis. *Plant Journal*, **64**, 775–789.

Verbon, E.H., Liberman, L.M., Zhou, J., Yin, J., Pieterse, C.M.J., Benfey, N., Stringlis, I.A. and Jonge, R. De (2023) Cell-type-specific transcriptomics reveals that root hairs and endodermal barriers play important roles in beneficial plant-rhizobacterium interactions. *Mol Plant*, **16**, 1160-1177. https://doi.org/10.1016/j.molp.2023.06.001.

Vissenberg, K., Claeijs, N., Balcerowicz, D. and Schoenaers, S. (2020) Hormonal regulation of root hair growth and responses to the environment in Arabidopsis. *J Exp Bot*, **71**, 2412–2427.

Vogel, J. and Somerville, S. (2000) Isolation and characterization of powdery mildew-resistant *Arabidopsis* mutants. *Proc Natl Acad Sci USA*, **97**, 1897–1902. https://doi.org/10.1073/pnas.030531997

Wang, Xuemin, Du, G., Wang, Xuming, Meng, Y., Li, Y., Wu, P. and Yi, K. (2010) The function of LPR1 is controlled by an element in the promoter and is independent of SUMO E3 ligase SIZ1 in response to Low Pi stress in *Arabidopsis thaliana*. *Plant Cell Physiol*, **51**, 380–394.

Wang, Y., Li, X., Fan, B., Zhu, C. and Chen, Z. (2021) Regulation and function of defense-related callose deposition in plants. *Int J Mol Sci*, **22**, 1–15.

Wang, Z., Zheng, Z., Song, L. and Liu, D. (2018) Functional characterization of Arabidopsis PHL4 in plant response to phosphate starvation. *Front Plant Sci*, **9**, 1–19.

Weßling, R. and Panstruga, R. (2012) Rapid quantification of plant-powdery mildew interactions by qPCR and conidiospore counts. *Plant Methods*, **8**.

Wildermuth, M.C., Dewdney, J., Wu, G. and Ausubel, F.M. (2002) Isochorismate synthase is required to synthesize salicylic acid for plant defence. *Nature*, **417**, 571–571.

Wu, S.W., Kumar, R., Iswanto, A.B.B. and Kim, J.Y. (2018) Callose balancing at plasmodesmata. *J Exp Bot*, **69**, 5325–5339.

Zhang, X., Chen, S. and Mou, Z. (2010) Nuclear localization of NPR1 is required for regulation of salicylate tolerance, isochorismate synthase 1 expression and salicylate accumulation in *Arabidopsis*. *J Plant Physiol*, **167**, 144–148. http://dx.doi.org/10.1016/j.jplph.2009.08.002.

Zhu, C., Gan, L., Shen, Z. and Xia, K. (2006) Interactions between jasmonates and ethylene in the regulation of root hair development in *Arabidopsis*. *J Exp Bot*, **57**, 1299–1308.

Figure Legends

Figure 1. Callose deposition is induced in root hairs under phosphate deficiency, independently of *PHR1*, *PHL1*, *LPR1* and *LPR2*.

Aniline blue staining of the primary roots 3 d after transfer of 5-d-old *Arabidopsis thaliana* seedlings to low Pi (P 50 µM), low N (N 0 µM), low Fe (Fe 0 µM), low Fe low Pi (Fe 0 µM, P 0 µM) and normal Pi (P 625 µM) liquid media. (a, c, left) Bright field (BF) images. (a, c, right) Aniline blue staining images. White arrowheads indicate root hair tips. (b) Aniline blue and propidium iodide (PI) staining of root hairs under low Pi and normal Pi conditions. (c) The wild-type (WT) used for *rhd6* was Ws-2. Magnified images of the upper panels (a, c, lower panel). Bars, 500 µm (a, c, upper panel), 50 µm (a, c lower panel) and 25 µm (b).

Figure 2. Callose synthase *PMR4* is required for callose induction in root hairs under phosphate deficiency.

(a) Aniline blue staining of the primary roots 3 d after transfer of 5-d-old *Arabidopsis thaliana* plants to low Pi (0 µM) liquid media. Left, bright filed (BF) images. Right, aniline blue staining images. Two *caps* mutants are defective in inducing root hair *callose under phosphate starvation.* Lower panels, magnified images of upper panels. White arrowheads indicate root hair tips. Bars, 500 µm (upper panel) and 50 µm (lower panel). (b) Positions of mutated sites in the *PMR4* locus of *caps* mutants, illustrated

with Protter (Omasits *et al*., 2014). An orange region corresponds to the catalytic domain of PMR4. Asterisks indicate stop codons. (c) Ratios of *G. orontii* genomic DNA to Arabidopsis genomic DNA were determined by qPCR with the primers for Plasma membrane ATPase 1 (Go_V1_Conting3757) and RNA-binding family protein (AT3G21215), respectively. Data show means ± SD (n=3). Asterisks (** $P < 0.01$) indicate statistically significant differences relative to WT according to Student's *t*-test.

Figure 3. *PMR4* modulates root and shoot growth under phosphate deficiency. (a) Aniline blue staining of the primary roots 3 d after transfer of 5-d-old *Arabidopsis* seedlings to low Pi (50 µM) liquid media. Left, bright filed (BF) images. Right, aniline blue staining images. Lower panels, magnified images of upper panels. White arrowheads indicate root hair tips. Bars, 100 µm (upper panel) and 50 µm (lower panel). (b) Six-d-old seedlings were exposed to low Pi (50 µM) or normal Pi (625 µM) media for 6 d. Bar, 10 mm. (c) The primary roots 6 d after transfer of 6-d-old seedling to low Pi (50 µM) and normal Pi media. Bar, 1 mm. (d) Root hair density and root hair length in (c). (e) Primary root length in (b). (f) Ratio of the primary root length under normal Pi relative to low Pi of (e). (g) Convex hull area of the roots in (b). (h) Ratio of convex hull area under normal Pi relative to low Pi of (g). (i) Six-d-old seedlings were exposed to low Pi (50 µM) or normal Pi for 16 d. Bar, 10 mm. (j) SFW in (i). (k) Six-d-old seedlings were exposed to low Pi (10 µM) media for 10 d. Bar, 10 mm. (l) Anthocyanin contents in (k). (d-h, j, l) Data show means ± SD (n = 7~30). Statistical significance was assessed by one-way ANOVA and Tukey honestly significant different (HSD) test for each medium dataset. Different letters indicate significant differences ($P < 0.05$). NS, not significant.

Figure 4. *PMR4* contributes to shoot growth and Pi accumulation under phosphate deficiency.

(a-d) Pi contents per root or shoot fresh weight (FW) of 6-d-old *Arabidopsis* seedlings exposed to low Pi (150 µM) or normal Pi (625 µM) for 9 d. The WT used for *rhd6* was Ws-2 (c,d). (e) Illustration of Pi accumulation in the indicated genotypes, referred to (a-d). (f) ^{32}P uptake rate determined in the whole seedlings exposed to the indicated media (low Pi and normal Pi, 50 and 625 µM Pi, respectively) supplemented with ^{32}P-labelled Pi for 30 min. (g) Ratio of ^{32}P in the shoots relative to the whole plants after pulse-chase labeling. Seedlings grown on low and normal Pi media (50 and 625 µM, respectively) were exposed to ^{32}P-labeled Pi for 10 min, and then transferred to non-labelled Pi media for 120 min (pulse chase). The ^{32}P ratio of the shoots to the whole seedlings was calculated for the translocation rate of Pi. Data show means ± SD (n = 3~4) in (a-d, f-g). Statistical significance was assessed by one-way ANOVA and Tukey HSD test for each medium dataset (a-b, f-g). Different letters indicate significant differences ($P < 0.05$). Asterisks (** $P < 0.01$, * $P < 0.05$) indicate statistically significant differences relative to WT according to Student's *t*-test (c, d). NS, not significant.

Figure 5. Plant element contents in shoot and root under low phosphate conditions.

Element contents (P, K, Mg, Ca, Mn, Fe, Zn, Mo and B) per root or shoot dry weight (DW) of 5-d-old *Arabidopsis thaliana* seedlings exposed to low Pi (10 µM) for 4 d. Data show means ± SD (n=4~8). Asterisks (** $P < 0.01$, * $P < 0.05$) indicate

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Figure 6. Root *PMR4* contributes to shoot growth under phosphate deficiency.

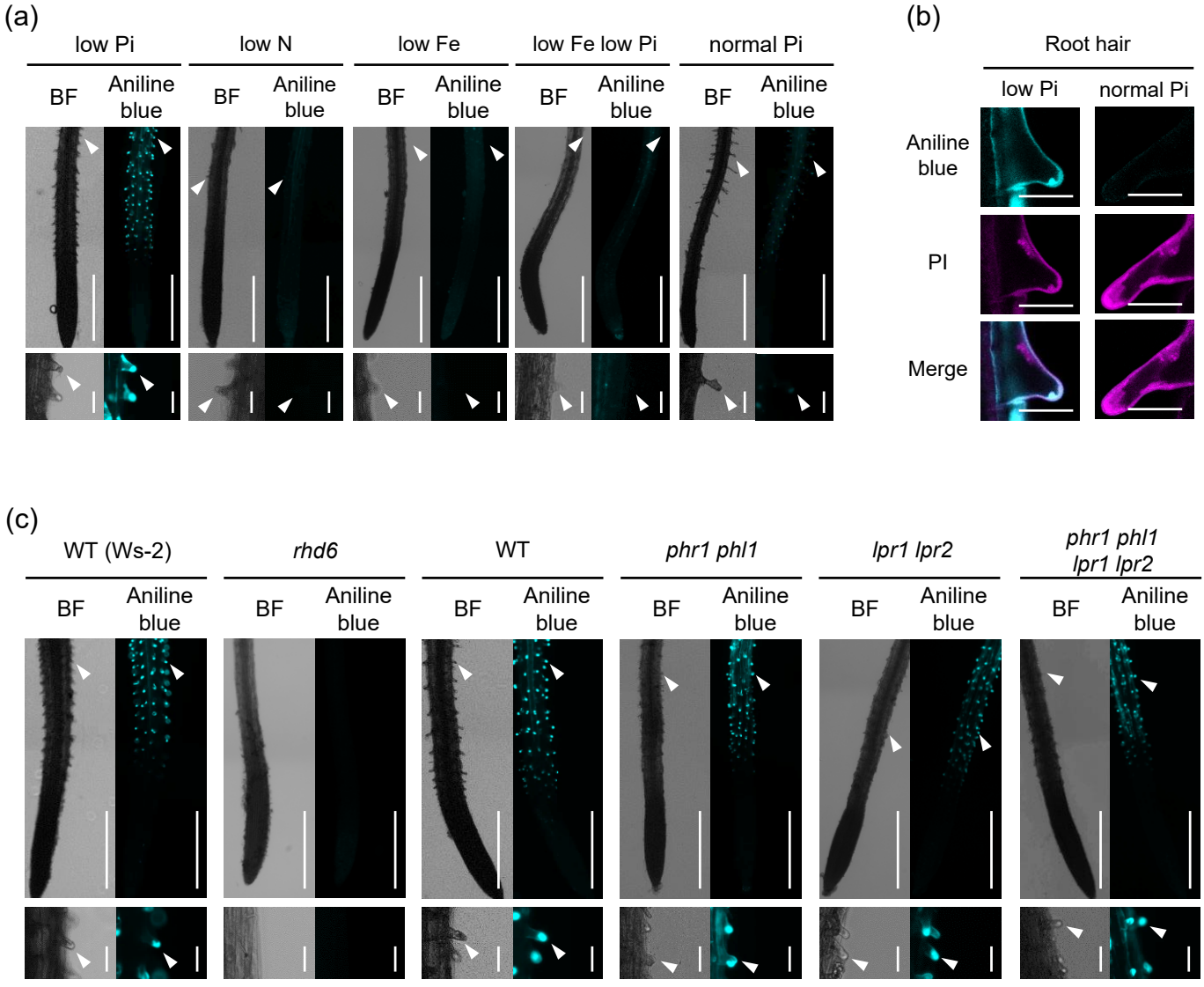
(a) Reciprocal micrografting of *Arabidopsis* WT and *pmr4* mutants. Images were obtained at 14 d after transfer to low Pi (150 μM) media. Bar, 10 mm. (b) Shoot fresh weight (SFW) of (a). (c) The primary root length of (a). (d) Aniline blue staining of grafted-seedlings at 9 d after transfer to low Pi (50 μM). (d, left) Bright filed (BF) images. (d, right) Aniline blue staining images. (d, lower panel) Magnified images of (d, upper panel). White arrows indicate root hair tips. Bar, 100 μm (c, upper panel) or 20 μm (c, lower panel). (b, c) Data show means ± SD (n=19-26). Statistical significance was assessed by one-way ANOVA and Tukey HSD test for each medium dataset. Different letters indicate significant differences ($P < 0.05$).

Figure 7. PMR4 in root hairs contributes to shoot growth under phosphate deficiency.

(a) Aniline blue staining of the primary roots 3 d after transfer of 5-d-old *Arabidopsis* seedlings to low Pi (10 μM). White arrowheads indicate root hair chips. Bars, 100 μm (upper panel) and 50 μm (lower panel). (b) Five-d-old seedlings were exposed to low Pi (50 μM) for 10 d. Bar, 10 mm. (c) SFW of (b). (d) Primary root length of (b). (c, d) Data show means ± SD (n = 22-48). Asterisks (** $P < 0.01$, * $P < 0.05$) indicate statistically significant differences relative to *pmr4* according to Student's *t*-test. NS, not significant.

Figure 8. Model for PMR4-mediated root hair callose deposition in plant adaptation to Pi deficiency.

PMR4 induces callose deposition in root hairs under phosphate deficiency, and thus contributes to shoot Pi accumulation and growth through enhancing Pi uptake and root-to-shoot transfer. This contribution alleviates phosphate starvation symptoms, such as root system area reduction and anthocyanin accumulation.



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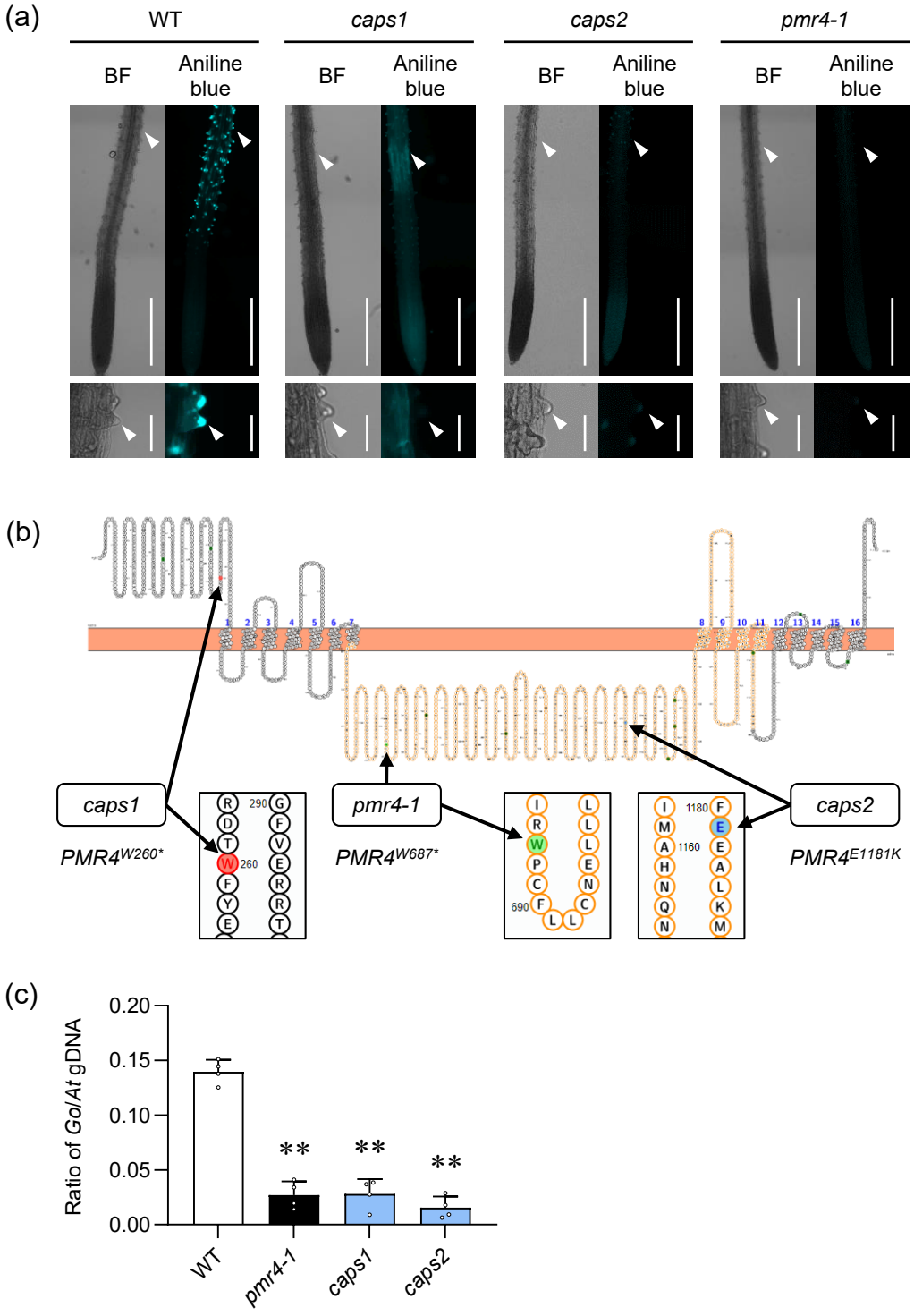

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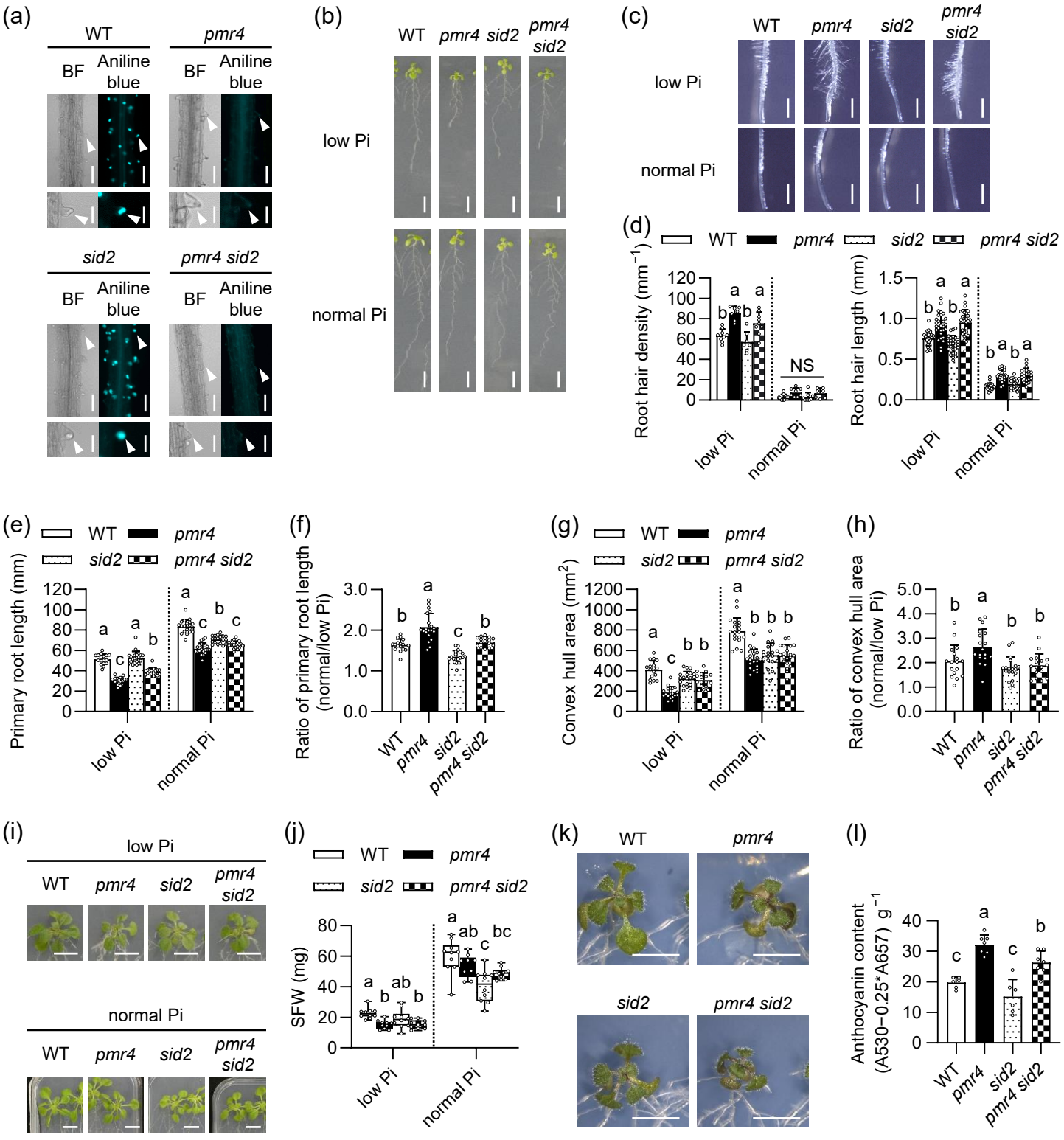

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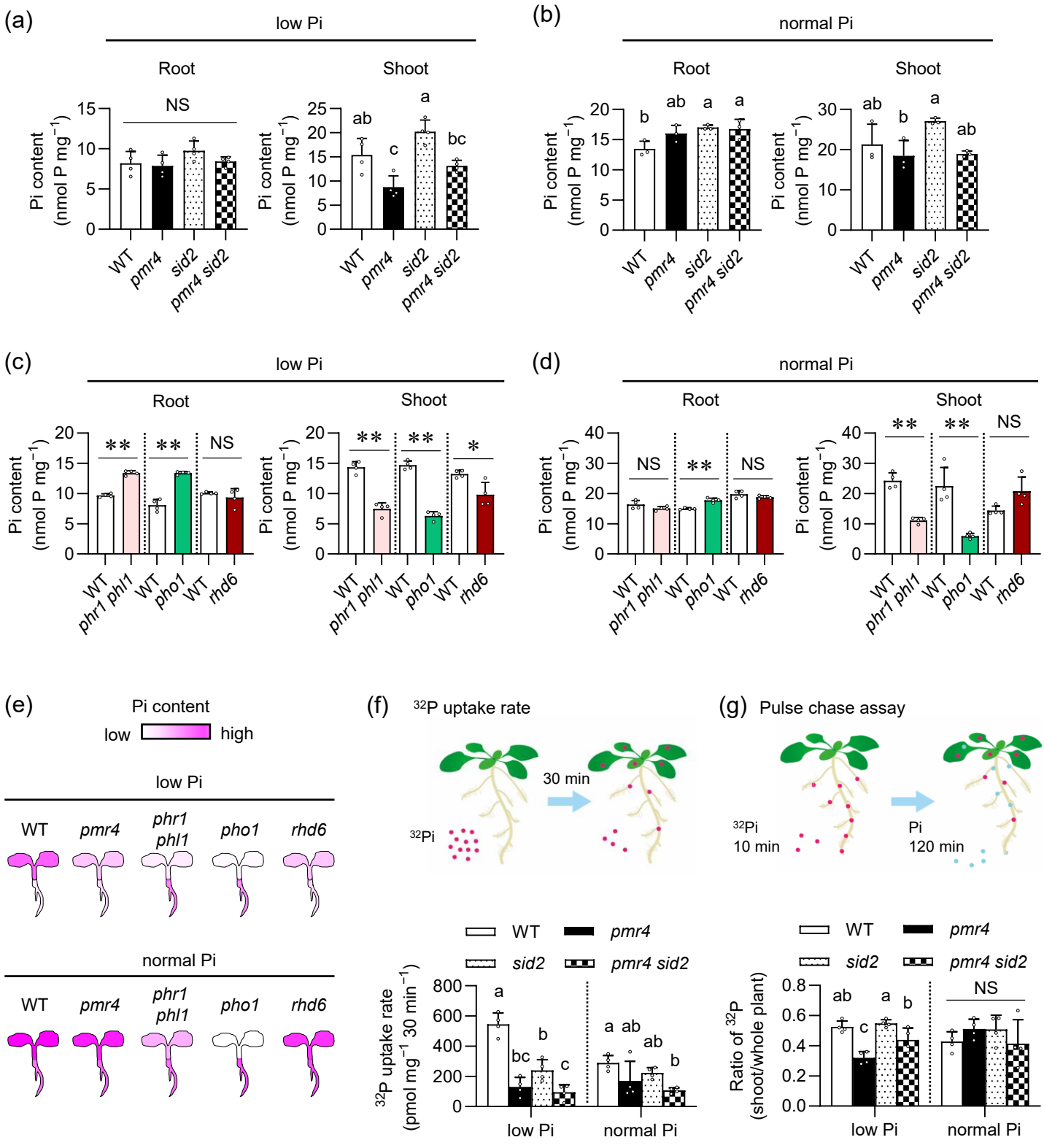

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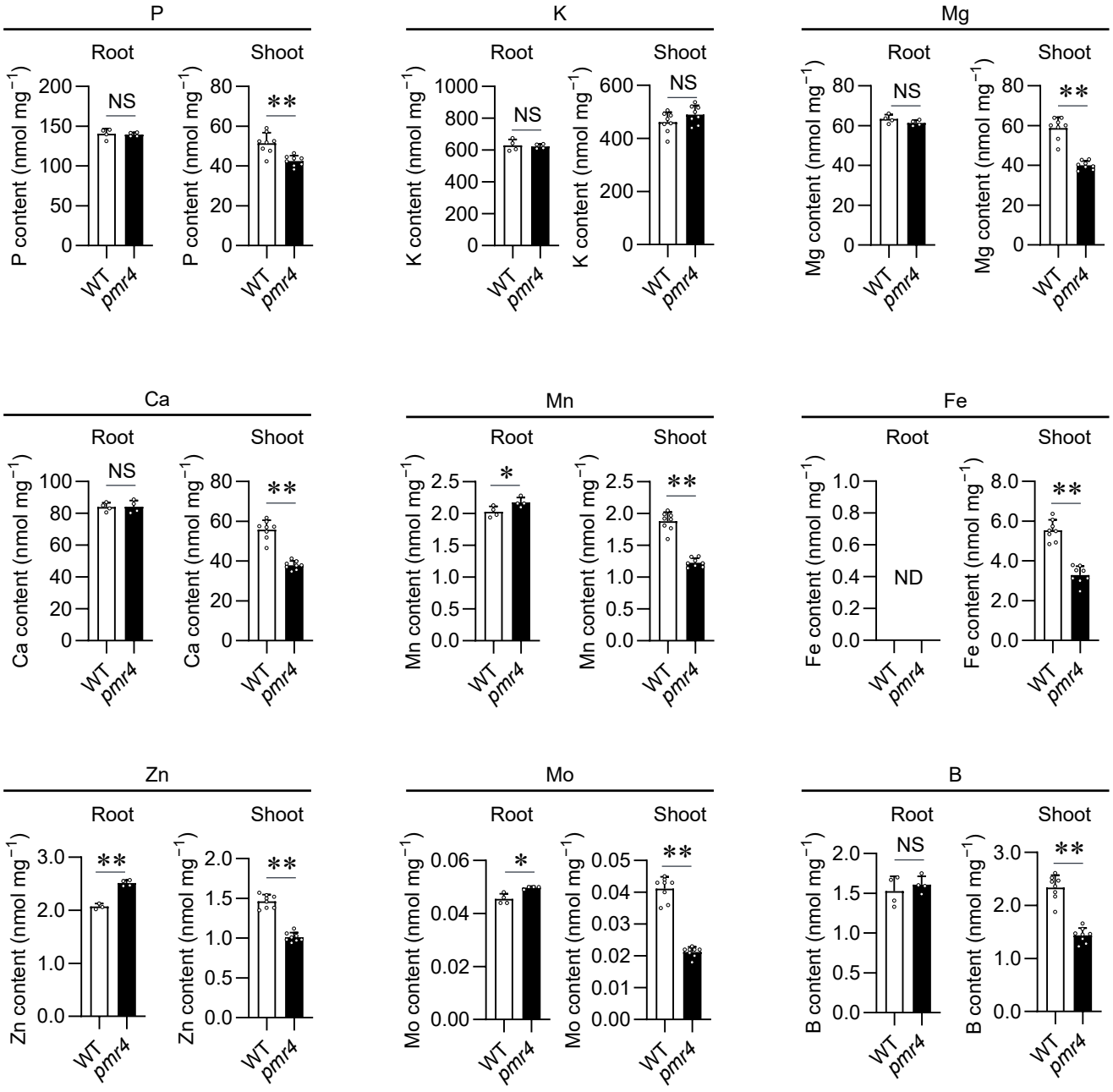

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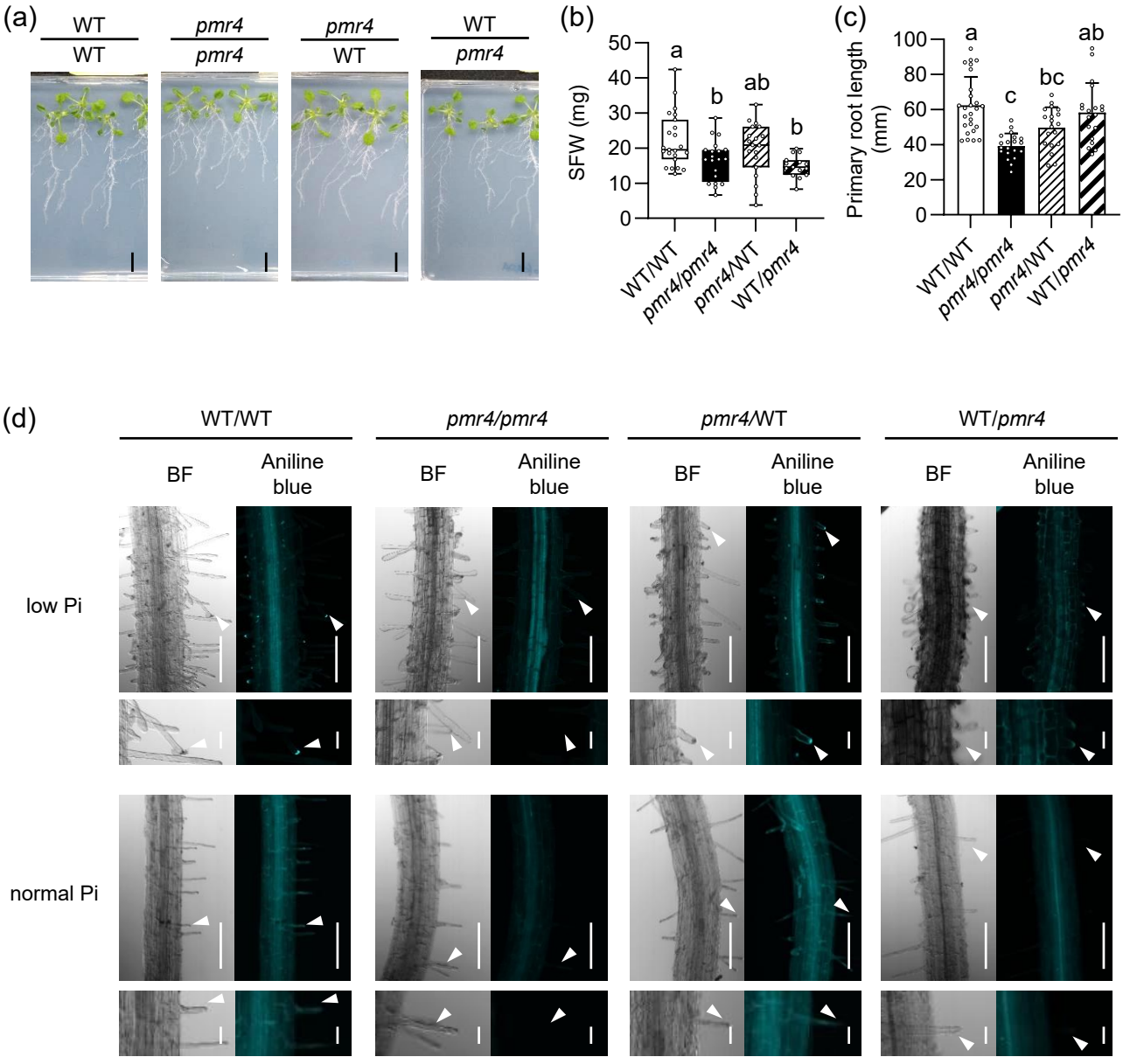

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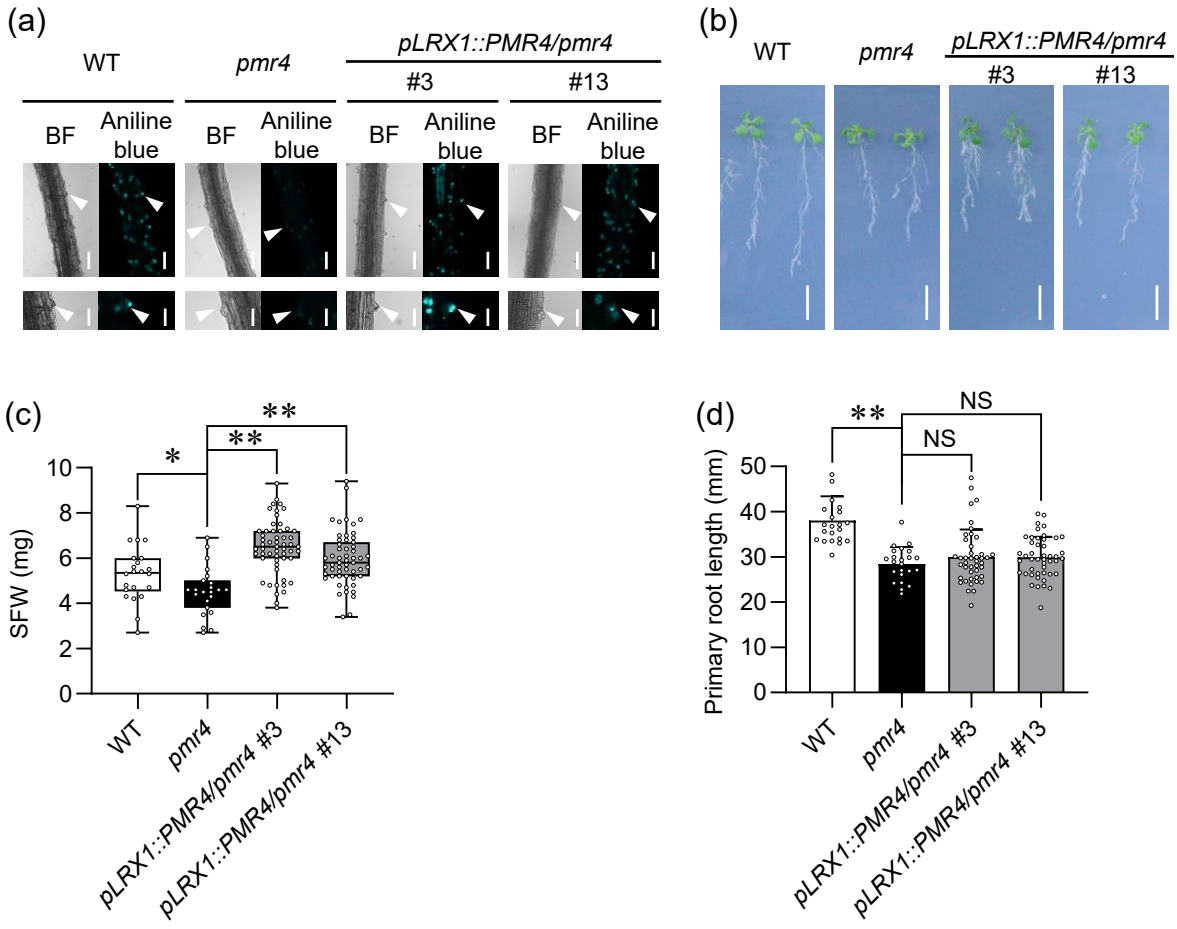

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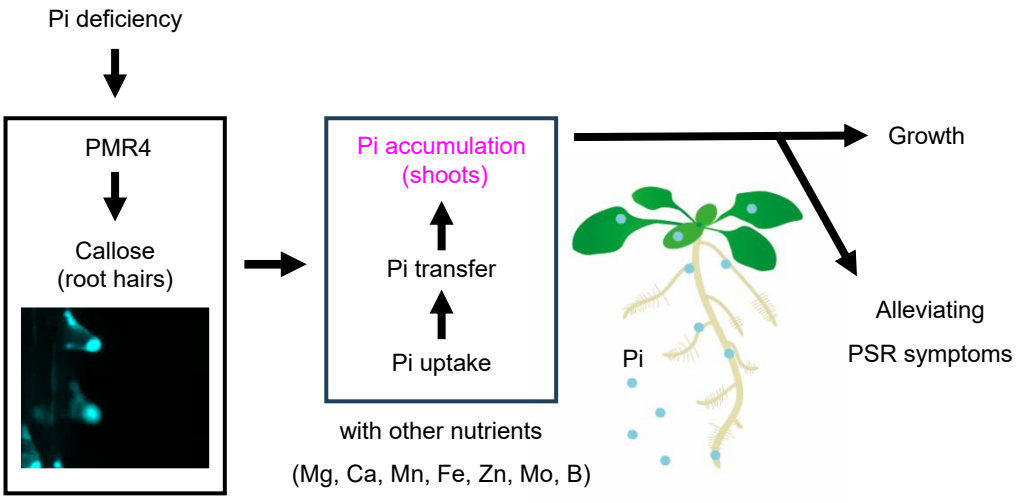

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