Vacuolar Ca\textsuperscript{2+}/H\textsuperscript{+} Transport Activity Is Required for Systemic Phosphate Homeostasis Involving Shoot-to-Root Signaling in Arabidopsis\textsuperscript{[W]}\textsuperscript{[OA]}

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Calcium ions (Ca\textsuperscript{2+}) and Ca\textsuperscript{2+}-related proteins mediate a wide array of downstream processes involved in plant responses to abiotic stresses. In Arabidopsis (Arabidopsis thaliana), disruption of the vacuolar Ca\textsuperscript{2+}/H\textsuperscript{+} transporters CAX1 and CAX3 causes notable alterations in the shoot ionome, including phosphate (P\textsubscript{i}) content. In this study, we showed that the cax1/cax3 double mutant displays an elevated P\textsubscript{i} level in shoots as a result of increased P\textsubscript{i} uptake in a miR399/PHO2-independent signaling pathway. Microarray analysis of the cax1/cax3 mutant suggests the regulatory function of CAX1 and CAX3 in suppressing the expression of a subset of shoot P\textsubscript{i} starvation-responsive genes, including genes encoding the PHT1;4 P\textsubscript{i} transporter and two SPX domain-containing proteins, SPX1 and SPX3. Moreover, although the expression of several PHT1 genes and PHT1;1/2/3 proteins is up-regulated in the root of cax1/cax3, results from reciprocal grafting experiments indicate that the cax1/cax3 scion is responsible for high P\textsubscript{i} accumulation in grafted plants and that the pht1;1 rootstock is sufficient to moderately repress such P\textsubscript{i} accumulation. Based on these findings, we propose that CAX1 and CAX3 mediate a shoot-derived signal that modulates the activity of the root P\textsubscript{i} transporter system, likely in part via posttranslational regulation of PHT1;1 P\textsubscript{i} transporters.

Transient increases in cytoplasmic calcium concentrations ([Ca\textsuperscript{2+}]\textsubscript{cyt}) or the spatial and temporal dynamics of stimulus-induced alterations in [Ca\textsuperscript{2+}]\textsubscript{cyt} constitute a signal that mediates a wide array of downstream processes involved in plant responses to many developmental cues and environmental stresses (Knight, 2000; McAinsh and Pittman, 2009). The generation of such stimulus-specific Ca\textsuperscript{2+} signatures is associated with various Ca\textsuperscript{2+} channels, transporters, and pumps throughout the membrane system. In particular, tonoplast-localized Ca\textsuperscript{2+}/H\textsuperscript{+} exchangers and Ca\textsuperscript{2+}-ATPase pumps play a key role in the sequestration of Ca\textsuperscript{2+} into the vacuole, the primary pool for Ca\textsuperscript{2+} buffering and release, and are assumed to participate in resetting the [Ca\textsuperscript{2+}]\textsubscript{cyt} following stimuli (Hirschi, 2004; McAinsh and Pittman, 2009).

In yeast, the vacuolar Ca\textsuperscript{2+}/H\textsuperscript{+} exchanger VCX1, as a high-capacity and low-affinity Ca\textsuperscript{2+} transporter, functions to rapidly sequester cytosolic Ca\textsuperscript{2+} and supposedly attenuates the activation of Ca\textsuperscript{2+} signaling pathways, as the vcx1Δ strain displayed a transient and strongly elevated [Ca\textsuperscript{2+}]\textsubscript{cyt} followed by a slow and weak recovery from a Ca\textsuperscript{2+} shock (Miseta et al., 1999). In Arabidopsis (Arabidopsis thaliana), the cation/H\textsuperscript{+} exchangers CAX1 (the ortholog of VCX1), CAX3, and CAX4 are phylogenetically grouped into type IA, whereas CAX2, CAX5, and CAX6 belong to type IB (Shigaki et al., 2006). However, only CAX1 to CAX4 have been functionally characterized to possess a vacuolar Ca\textsuperscript{2+}/H\textsuperscript{+} exchange activity. Whereas CAX1 and CAX3 mediate specifically Ca\textsuperscript{2+} transport (Hirschi, 1999; Catala et al., 2003; Cheng et al., 2003; Mei et al., 2007; Zhao et al., 2008), CAX2 and CAX4 were documented to have high transport and selectivity for cadmium ions (Cd\textsuperscript{2+}) over Ca\textsuperscript{2+} in tonoplast vesicles (Hirschi et al., 2000; Cheng et al., 2002; Pittman et al., 2004; Korenkov et al., 2007).

Knockout of CAX1 in Arabidopsis increased the tolerance to high concentrations of various ions, Ca\textsuperscript{2+}-depleted conditions, and freezing after cold acclimation (Catala et al., 2003; Cheng et al., 2003). Conversely, transgenic tobacco plants overexpressing CAX1 were hypersensitive to ion imbalance and cold shock and...
exhibited Ca\(^{2+}\)-deficient symptoms in spite of increased accumulation of Ca\(^{2+}\) (Hirschi, 1999). Loss-of-function of CAX3, a close homolog of CAX1, increased the sensitivity to salt stress, lithium, and low pH (Zhao et al., 2008). While cax1 and cax3 single mutants displayed subtle phenotypes, the cax1/cax3 double mutant showed stunted growth with chlorosis on the leaf tips and a drastic reduction in silique size (Cheng et al., 2005; Zhao et al., 2008). Elemental analysis also revealed that impairment of CAX1 and CAX3 caused dramatic alterations in the shoot ionome, namely elevated levels of phosphate (PO\(_4\)\(^{3-}\); P), manganese (Mn\(^{2+}\)), and zinc (Zn\(^{2+}\)) and decreased Ca\(^{2+}\) and magnesium (Mg\(^{2+}\)) ion concentrations (Cheng et al., 2005). A recent study further showed that the leaf apoplastic free Ca\(^{2+}\) concentration ([Ca\(^{2+}\)]\(_{a}\)) of cax1/cax3 was 3-fold greater than that of the wild type, accounting for the phenotypes of reduced cell wall extensibility, stomatal aperture, transpiration, CO\(_2\) assimilation, and leaf growth rate (Conn et al., 2011b). Despite no evidence yet of a direct involvement of Ca\(^{2+}\) in P\(_7\) signaling, the observation of an increased accumulation of P\(_7\) in the cax1/cax3 mutant has provided the first link between Ca\(^{2+}\) and P\(_7\) homeostasis and thus drew our attention and interest to investigate the potential role of CAX1 and CAX3 in P\(_7\) signaling.

Phosphorus (P), one of the mineral nutrients essential for plant survival and productivity, is a major structural constituent of fundamental macromolecules such as nucleic acids and phospholipids and is involved in energy transfer, metabolic regulation, and protein activation. However, most of the P in the soil is unavailable for plant uptake because of adsorption, precipitation, or conversion to organic forms (Marschner, 1995). As a result, plants constantly encounter P\(_7\) limitation and have developed a number of adaptive strategies to maintain P\(_7\) homeostasis, including enhancing acquisition of P\(_7\), coordinating allocation of P\(_7\) among different organs, and remobilizing P\(_7\) from old to young tissues (Poirier and Bucher, 2002; Ticconi and Abel, 2004). Although many plant responses to P\(_7\) starvation have been extensively explored, the molecular mechanisms by which plants sense the P\(_7\) signal and elicit these responses remain largely unknown.

Identification and characterization of several mutants with aberrant responses to P\(_7\) starvation or with altered levels of P\(_7\) has advanced our understanding of the molecular components involved in P\(_7\) homeostasis (Lin et al., 2009; Chiu and Lin, 2011). For example, the Arabidopsis pho2 mutant accumulated high levels of P\(_7\) in the shoot and showed symptoms of P\(_7\) toxicity as a result of increased P\(_7\) uptake and translocation of P\(_7\) from roots to shoots (Delaize and Randall, 1995; Dong et al., 1998). The PHO2 gene was identified to encode an E2 ubiquitin-conjugating enzyme (UBC24), whose expression during P\(_7\) deficiency is posttranscriptionally suppressed by a specific microRNA, miR399 (Fujii et al., 2005; Aung et al., 2006; Bari et al., 2006; Chiu and Lin, 2011). In accordance with the inverse correlation between miR399 and PHO2 mRNA levels, transgenic plants overexpressing miR399 phenocopied the pho2 mutant and PHO2 T-DNA knockout lines (Aung et al., 2006; Chiu and Lin, 2011). Reciprocal micrografting experiments further demonstrated that a pho2 root genotype is sufficient and necessary for P\(_7\) accumulation in the shoot, whereas the shoot-to-root movement of mature miR399 is responsible for the degradation of PHO2 mRNA in roots (Bari et al., 2006; Lin et al., 2008; Pant et al., 2008). Moreover, up-regulation of miR399 by P\(_7\) deprivation is mediated by the PHOSPHATE STARVATION RESPONSE1 (PHR1) transcription factor, a key positive regulator of multiple P\(_7\) starvation-induced (PSI) genes (Rubio et al., 2001; Bari et al., 2006). These findings suggest that the P\(_7\) starvation signaling involving PHR1, miR399, and PHO2 is crucial for the maintenance of P\(_7\) homeostasis. Recently, a unique gene family in Arabidopsis (AtSPX1–AtSPX4) exclusively harboring the SPX (for SYG1/Pho81/XPR1) domain was identified to be regulated by P\(_7\) starvation, in part through PHR1 (Hamburger et al., 2002; Wang et al., 2004; Duan et al., 2008). Overexpression of SPX1 (At5g20150) increased the transcript levels of several PSI genes under both P\(_7\)-sufficient and P\(_7\)-deficient conditions, whereas RNA interference-mediated partial down-regulation of SPX3 (At2g45130) led to aggravated P\(_7\) deficiency symptoms, altered P\(_7\) allocation, and enhanced expression of a subset of PSI genes (Duan et al., 2008). In rice (Oryza sativa), OsSPX1 (Os06g40120), the ortholog of SPX3, acts via a negative feedback loop to adjust the expression of several PSI genes under P\(_7\)-limited conditions (Wang et al., 2009; Liu et al., 2010). These findings revealed that the plant SPX domain-containing proteins are new players in the regulatory network of P\(_7\) signaling.

Although Ca\(^{2+}\) and Ca\(^{2+}\)-related proteins are indispensable messengers in the signal transduction of many stress responses, the role of Ca\(^{2+}\) in P\(_7\) signaling and the cross talk between Ca\(^{2+}\) and P\(_7\) homeostasis are barely understood. The association between loss of CAX1 and CAX3 activities and increased P\(_7\) accumulation as seen in cax1/cax3 (Cheng et al., 2005) provides an opportunity to tackle this issue. It is also of interest to compare cax1/cax3 and pho2 mutants in terms of regulatory pathways, because both mutants show elevated shoot P\(_7\) concentrations. Here, our study of the cax1/cax3 mutant suggests that vacuolar Ca\(^{2+}\)/H\(^{+}\) transporters exert a negative regulation of P\(_7\) starvation responses, as revealed by suppression of the expression of a subset of shoot P\(_7\) starvation-responsive (PSR) genes and inhibition of P\(_7\) uptake activity in the root. Our results also suggest that the effects of CAX1/ CAX3- and PHO2-mediated signaling pathways on the suppression of P\(_7\) uptake are different. Moreover, results from reciprocal grafting experiments demonstrate that CAX1 and CAX3 mediate a shoot-derived
signal that modulates the activity of the root $P_i$ transporter system, likely in part via posttranslational regulation of PHT1;1.

RESULTS

cax1/cax3 Mutant Accumulates High Levels of $P_i$ in Shoot and Displays Increased $P_i$ Transport Activity

The shoot $P_i$ concentration has been shown to increase by 66% in the cax1/cax3 mutant but to remain unchanged in the single cax1 and cax3 mutants (Cheng et al., 2005). To confirm this finding, we grew cax1, cax3, and cax1/cax3 along with the pho2 mutant and wild-type controls in $P_i$-sufficient (+$P_i$) or $P_i$-deficient ($-P_i$) half-strength modified Hoagland hydroponic solution for $P_i$ concentration measurement. Under both +$P_i$ and $-P_i$ conditions, the shoot and root $P_i$ concentrations of cax1 and cax3 were similar to those of the wild type; by contrast, the cax1/cax3 double mutant accumulated high levels of $P_i$ in the shoot under both conditions and showed modestly increased $P_i$ concentrations in the root under $P_i$ deficiency (Fig. 1). Of note, cax1/cax3 did not accumulate $P_i$ in the shoot to a level as high as pho2 under +$P_i$ conditions but maintained a comparable level of shoot $P_i$ as pho2 under $-P_i$ conditions (Fig. 1A).

We then performed $P_i$ transport assays to determine whether $P_i$ accumulation in cax1/cax3 can be attributed to an enhanced $P_i$ uptake rate. As expected, cax1/cax3 exhibited a higher $P_i$ uptake activity than wild-type plants, regardless of external $P_i$ concentrations (Fig. 2, A and B). By contrast, pho2 exhibited a higher $P_i$ uptake activity than cax1/cax3 when the $P_i$ supply was adequate (Fig. 2A) but exhibited only a slightly enhanced $P_i$ uptake activity as compared with the wild type when $P_i$ was limited (Fig. 2B). Under +$P_i$ conditions, pho2 but not cax1/cax3 showed an increased shoot-to-root ratio of $P_i$ distribution (Fig. 2C), indicating that the $P_i$ translocation activity of cax1/cax3 from roots to shoots was not changed even though the $P_i$ uptake activity was increased. However, under $-P_i$ conditions, the shoot-to-root ratio of $P_i$ distribution in cax1/cax3 was increased (Fig. 2D). Taken together, cax1/cax3 displayed increased shoot $P_i$ accumulation, increased $P_i$ uptake activity regardless of external $P_i$ concentrations, and greater $P_i$ translocation from roots to shoots when $P_i$ was limited.

$P_i$ Accumulation in cax1/cax3 Mutant Increases in an Exogenous [Ca$^{2+}$]-Dependent Manner

It has been reported that cax1/cax3 showed a higher sensitivity to Ca$^{2+}$ stress when grown in medium supplemented with high [Ca$^{2+}$] (Cheng et al., 2005). To address whether the impaired cellular Ca$^{2+}$ homeostasis due to the loss of CAX1 and CAX3 plays a direct role in enhancing $P_i$ uptake activity, we examined the effect of exogenous Ca$^{2+}$ on $P_i$ accumulation of cax1/cax3 by growing mutants in +$P_i$ hydroponic medium containing different concentrations of Ca$^{2+}$ (0.05 mM, 0.25 mM, 1 mM, and 2 mM [Ca$^{2+}$]), contrasting with 2.5 mM [Ca$^{2+}$] in half-strength modified Hoagland solution. Interestingly, $P_i$ accumulation in the shoot of cax1/cax3 was increased in a [Ca$^{2+}$]-dependent manner (Fig. 3B) and displayed a negative correlation with the leaf size of mutants (Fig. 3A). Consistent with previous results showing that the growth defects of cax1/cax3 mutant were suppressed by supplemented exogenous Mg$^{2+}$ (Cheng et al., 2005), we also found that the exacerbated $P_i$ accumulation in cax1/cax3 mutants resulting from exogenous Ca$^{2+}$ stress could be attenuated by supplementing high concentrations of Mg$^{2+}$ (Fig. 3). These results suggest an antagonistic relationship between Ca$^{2+}$ and Mg$^{2+}$ regarding their interplay in $P_i$ accumulation of cax1/cax3.

Cross-Regulation of PHO2 and CAX1/CAX3 Expression in cax1/cax3 and pho2 Mutants

To investigate the role of CAX1 and CAX3 in $P_i$ signaling, we first used quantitative reverse transcrip-

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Figure 1. Analysis of $P_i$ concentrations in the cax1/cax3 mutant. The $P_i$ concentrations are shown in the shoot (A) and root (B) of 23-d-old wild-type (WT), cax1, cax3, cax1/cax3, and pho2 plants grown under +$P_i$ or $-P_i$ conditions. Values represent means ± SD of three to five biological replicates. Data significantly different from the corresponding wild-type controls are indicated (* $P < 0.05$, ** $P < 0.01$; Student’s t test). FW, Fresh weight.

Figure 2. Analysis of $P_i$ concentrations in the cax1/cax3 mutant. The $P_i$ concentrations are shown in the shoot (A) and root (B) of 23-d-old wild-type (WT), cax1, cax3, cax1/cax3, and pho2 plants grown under +$P_i$ or $-P_i$ conditions. Values represent means ± SD of three to five biological replicates. Data significantly different from the corresponding wild-type controls are indicated (* $P < 0.05$, ** $P < 0.01$; Student’s t test). FW, Fresh weight.
tion (qRT)-PCR to examine changes in the expression levels of CAX1 and CAX3 in wild-type plants subjected to P$_i$ deficiency. In the shoot, the transcript level of CAX1 was reduced by 60% after 5 d of P$_i$ deprivation (Fig. 4A). However, no difference was observed in the expression level of CAX3 between P$_i$-sufficient and P$_i$-deficient shoots (Fig. 4B). In the root, no significant change was found in the transcript level of CAX1 in response to P$_i$ starvation (Fig. 4A), while the CAX3 transcript level was increased (Fig. 4B). Regulation of the expression of CAX1 and CAX3 by P$_i$ deficiency implied the involvement of these genes in P$_i$ starvation responses.

Although cax1/cax3 and pho2 mutants showed distinct properties in terms of P$_i$ uptake and translocation (Fig. 2), we wondered whether there is a cross talk between PHO2 and CAX1/CAX3-mediated P$_i$ signaling pathways. Therefore, we examined the expression of CAX1 and CAX3 in pho2 and of PHO2 in cax1/cax3. Under both +P$_i$ and −P$_i$ conditions, the transcript levels of CAX1 in the shoot and root of pho2 were similar to that of wild-type plants (Fig. 4A). Under +P$_i$ conditions, the transcript levels of CAX3 in the shoot and root were higher in pho2 than in wild-type plants; however, under −P$_i$ conditions, the transcript level of CAX3 was higher in the shoot but not in the root of pho2 as compared with wild-type plants (Fig. 4B).

The complementary measurements revealed that the transcript level of PHO2 in the shoot of cax1/cax3 was higher than that of wild-type plants under both +P$_i$ and −P$_i$ conditions (Fig. 4C). As the pho2 rootstock genotype has been shown sufficient for P$_i$ accumulation in the scion (Bari et al., 2006; Lin et al., 2008), the role of PHO2 in the shoot and the implication of increased transcript levels of PHO2 in the shoot of cax1/cax3 remain to be resolved. On the other hand, the level of PHO2 mRNA in the root of cax1/cax3 was reduced by 36% under +P$_i$ conditions as compared with wild-type plants but was not as low as what detected in the root of wild-type plants under P$_i$ deficiency (Fig. 4C). The mature miR399 was not detectable in the shoot or root of cax1/cax3 under +P$_i$ conditions (Fig. 4D), indicating that the moderate reduction of PHO2 mRNA level in the root of cax1/cax3 under +P$_i$ conditions did not result from suppression by miR399. Furthermore, the PHO2 transcripts were reduced to a similar level in the root of cax1/cax3 and wild-type plants under −P$_i$ conditions (Fig. 4C), supporting the idea that the increased P$_i$ uptake activity in the root of cax1/cax3 under P$_i$ deficiency (Fig. 2B) is unlikely mediated by a PHO2-dependent signaling pathway. Of interest, we also found that the induction of mature miR399s under P$_i$ deficiency in the root of cax1/cax3 under −P$_i$ conditions did not result from suppression by miR399. Furthermore, the PHO2 transcripts were reduced to a similar level in the root of cax1/cax3 and wild-type plants under −P$_i$ conditions (Fig. 4C), supporting the idea that the increased P$_i$ uptake activity in the root of cax1/cax3 under P$_i$ deficiency (Fig. 2B) is unlikely mediated by a PHO2-dependent signaling pathway. Of interest, we also found that the induction of mature miR399s under P$_i$ deficiency in the root of cax1/cax3 was lower than that of wild-type controls; therefore, it is likely that miR399 induction is inhibited by the elevated concentrations of internal shoot P$_i$, as shown in Figure 1A. From these results, we conclude that the expression of CAX1/CAX3 and PHO2 appears to be cross-regulated, even though they function to suppress P$_i$ uptake and regulate root-to-shoot P$_i$ translocation in different modes.
Suppression of a Subset of Shoot PSR Genes by CAX1 and CAX3

To gain more insight into the molecular basis for Pi accumulation in cax1/cax3, we analyzed the expression profiles of cax1/cax3 and ecotype Columbia (Col-0) wild-type plants grown in hydroponic solutions under +Pi and -Pi conditions using the Affymetrix ATH1 chip. The numbers of differentially expressed genes (P ≤ 0.01, more than 2-fold change) were identified as follows: 854 and 1,047 under +Pi and -Pi conditions, respectively, between cax1/cax3 and wild-type shoots; and 84 and 655 under +Pi and -Pi conditions, respectively, between cax1/cax3 and wild-type roots (Supplemental Table S1, subgroups 1–4). We also determined the number of PSR genes with more than 2-fold change in expression (P ≤ 0.01) between the +Pi and -Pi conditions (Supplemental Table S1, subgroups 5–8) and compared the expression of the PSR genes between cax1/cax3 and wild-type plants. To evaluate the effects of our Pi starvation regimes on gene expression in wild-type plants, we plotted the fold change ratio of the differentially regulated shoot and root PSR genes from our microarray data against that reported by Morcuende et al. (2007), who subjected liquid culture-grown whole seedlings to Pi deprivation under continuous light. Despite different plant growth stages and growth conditions applied in these two studies, there is a clear positive correlation in the comparison as revealed by the $r^2$ value (Supplemental Fig. S1).

None of the wild-type shoot PSR genes (0 of 163) and only 2.6% (12 of 455) of the wild-type root PSR genes were regulated in the same fashion in cax1/cax3 (Fig. 5A). Strikingly, in the shoot of cax1/cax3 under +Pi conditions, 22.1% (32 of 145) of the wild-type PSi genes were up-regulated and 22.2% (4 of 18) of the wild-type Pi starvation-repressed genes were down-regulated (Fig. 5B), indicating that one-fifth of the PSR genes are constitutively activated in the shoot of cax1/cax3. Among these 36 differentially expressed PSR genes (Table I), we further validated the up-regulation of several PSi genes, including genes implicated in Pi signaling, SPX1 (At5g20150) and SPX3 (At2g45130), and PHT1;4 (At2g38940), a member of the P$_i$ high-affinity transporter (PHT1) gene family, by qRT-PCR (Fig. 6, A–C; Supplemental Fig. S3). Although our microarray data did not reveal gene expression changes of other PHT1 members, we were able to observe by qRT-PCR 4- to 5-fold increased transcript levels for PHT1;1 (At5g43350) and PHT1;3 (At5g43360) in the shoot of cax1/cax3 under +Pi conditions (data not shown). This discrepancy can be explained by the higher sensitivity of qRT-PCR, as the levels of PHT1;1 and PHT1;3 transcripts in the shoot were low. Up-regulation of these PHT1 genes may reflect an increased sink demand for P$_i$, likely a downstream event of the CAX1/CAX3 Pi signaling cascade. When the expression of the PSR genes was examined in cax1/cax3, most of them, such as SPX1, SPX3, and PHT1;4, were less responsive to Pi starvation compared with wild-type plants (Fig. 6, A–C; Supplemental Table S1, subgroups 5 and 6). This may have resulted from their suppression by the high shoot P$_i$ levels of cax1/cax3 (Fig. 1A). Given that a significant proportion of PSR genes were constitutively activated in the shoot of cax1/cax3, we conclude that the function of CAX1 and CAX3 is required for the suppression of a discrete subset of shoot PSR genes under +Pi conditions.

In contrast to the gene expression profile in the shoot of cax1/cax3 under +Pi conditions, only 2% (seven of 343) of the wild-type PSi genes and 1.8% (two of 112) of the wild-type P$_i$ starvation-repressed genes were up-regulated and down-regulated, respectively, in the root of cax1/cax3 under +Pi conditions (Fig. 5B; Table I). These results seem contradictory to our speculation that the increased P$_i$ uptake rate in cax1/cax3 may be caused by up-regulation of PHT1 genes in the root, as many members in the PHT1 P$_i$ transporter family are expressed preferentially in root epidermal or cortical cells and function in P$_i$ acquisition (Muchhal and Raghothama, 1996; Mudge et al., 2002). Both results of RT-PCR (Fig. 6D) and qRT-PCR (data not shown) analyses showed that the transcript levels of PHT1 genes were not increased under +Pi or under -Pi.
conditions in the root of \textit{cax1}/\textit{cax3} as compared with the wild-type controls, indicating that a posttranscriptional regulation of the \textit{PHT1} or other unidentified P\(_1\) transporters accounts for enhanced P\(_1\) transport activity in \textit{cax1}/\textit{cax3}.

Since the P\(_1\)-replete \textit{pht1;1} mutant showed reduced P\(_1\) uptake activity, \textit{PHT1;1} was suggested to play a primary role in P\(_1\) acquisition under +P\(_1\) conditions (Shin et al., 2004). Thus, we next examined whether \textit{PHT1;1} is up-regulated at the protein level in \textit{cax1}/\textit{cax3}. Because of the high homology in protein sequence (94\%–98\% identity) among \textit{PHT1;1}, \textit{PHT1;2}, and \textit{PHT1;3}, we raised an antibody against all three proteins. While the levels of \textit{PHT1;1/2/3} were increased in wild-type seedlings in response to low P\(_1\) availability, they were greatly decreased in the +P\(_1\) root of the \textit{pht1;1} knockout mutant (Fig. 7A). This validated the specificity of this antibody against \textit{PHT1;1/2/3} and supported the conclusion that \textit{PHT1;1} is the major P\(_1\) transporter accountable for P\(_1\) acquisition under +P\(_1\) conditions (Shin et al., 2004). Surprisingly, we did not observe much difference in the protein level of \textit{PHT1;1/2/3} in the +P\(_1\) root between \textit{cax1}/\textit{cax3} and the wild type (Fig. 7B). Taken together, the enhanced P\(_1\) uptake activity in \textit{cax1}/\textit{cax3} does not result from an up-regulation of \textit{PHT1;1/2/3} at the protein level.

**Shoot-Derived Signals Are Responsible for the High Accumulation of P\(_1\) in the \textit{cax1}/\textit{cax3} Mutant**

Since our microarray data revealed the activation of about 22\% of PSR genes in the shoot of \textit{cax1}/\textit{cax3} under +P\(_1\) conditions, we asked next whether a shoot-derived P\(_1\) starvation signal mediates the enhanced P\(_1\) uptake activity resulting in P\(_1\) accumulation of \textit{cax1}/\textit{cax3}. To address this issue, we performed reciprocal micrografting between \textit{cax1}/\textit{cax3} and wild-type plants. When grown under +P\(_1\) conditions, the grafted plants with \textit{cax1}/\textit{cax3} scions and wild-type rootstocks resembled the phenotype of \textit{PHT1;1} toxicity and stunted growth seen in the \textit{cax1}/\textit{cax3} mutant, whereas the grafted plants with wild-type scions and \textit{cax1}/\textit{cax3} rootstocks showed a wild-type phenotype (Fig. 8A). In accordance with the phenotype, while \textit{cax1}/\textit{cax3} scions grafted on wild-type rootstocks exhibited high shoot P\(_1\) levels (Fig. 8B) and P\(_1\) uptake activity (data not shown) as \textit{cax1}/\textit{cax3} self-grafts, wild-type scions grafted...
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**Figure 5.** The number of genes differentially expressed in cax1/cax3 and wild-type (WT) plants under P_i-sufficient and P_i-deficient conditions. A, Overlap of the PSR genes between cax1/cax3 and wild-type plants. B, Overlap of wild-type PSR genes and genes differentially expressed between cax1/cax3 and wild-type plants under +P_i conditions. Numbers designate the genes with significantly differential expression (P ≤ 0.01 and more than 2-fold change) between the indicated data sets derived from microarray analysis. The total number of genes in each data set is shown in parentheses. The numbers of induced and repressed genes are indicated in black and gray, respectively.

**DISCUSSION**

**CAX1- and CAX3-Mediated Regulation of Leaf Calcium Ion Homeostasis Is Required for Systemic P_i Homeostasis**

To maintain the [Ca^{2+}]_{cyt} levels in the micromolar range (Marty, 1999), plant cells transport Ca^{2+} out of the cytoplasm across the plasma membrane or into various organelles such as endoplasmic reticulum, chloroplast, and vacuole (Sze et al., 2000). CAX1 and CAX3 are tonoplast-localized Ca^{2+}/H^{+} antiporters that mediate the sequestration of Ca^{2+} into the vacuole (Cheng et al., 2003, 2005). Knockout or overexpression of CAX1 or CAX3 alone in planta has been shown to result in perturbations in ion homeostasis and altered responses to salinity and cold stresses, while loss of both CAX1 and CAX3 led to a severe reduction in growth, leaf tip and flower necrosis, and a pronounced sensitivity to exogenous Ca^{2+} and other ions (Hirschi, 1999; Catala et al., 2003; Cheng et al., 2003, 2005; Mei et al., 2007; Zhao et al., 2009). Intriguingly, alterations in transport properties resulting from overexpression of both CAXs in yeast could not be recapitulated by high-level expression of either transporter individually (Zhao et al., 2009). It was postulated that the differential stress sensitivities of cax mutants are due to specific responses by CAX1 or CAX3 to individual stresses or to distinct transport properties conferred by hetero-CAX complexes formed by CAX1 and CAX3 (Cheng et al., 2005; Zhao et al., 2009). If we suppose that loss of the putative CAX1-CAX3 heteromer is
responsible for the enhanced $P_i$ uptake in $cax1/cax3$, then lack of either CAX1 or CAX3 should have also rendered an increased $P_i$ uptake activity. However, our findings here suggest a functional redundancy of CAX1 and CAX3 regarding their role in regulating $P_i$ uptake, because neither the $cax1$ nor the $cax3$ single mutant exhibits an increased uptake rate of $P_i$. Similarly, no significant change in total leaf $[Ca^{2+}]$ in either the $cax1$ or the $cax3$ single mutant was observed (Cheng et al., 2005). Thus, one possible explanation for the impaired $P_i$ homeostasis in $cax1/cax3$ is that disruption of both vacuolar $Ca^{2+}$ transporters leads to...

**Table 1. Misregulated PSR genes in $cax1/cax3$**

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<tr>
<th>AGIa</th>
<th>Fold Change in Expression</th>
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</tr>
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<td></td>
<td>-$P_i$/$+$ in the Wild Type</td>
<td>$cax1/cax3$ Wild Type under $+$</td>
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<tr>
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<td>At5g20790</td>
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aArabidopsis Genome Initiative number. Genes validated by qRT-PCR are underlined.
elevated levels of \([\text{Ca}^{2+}]_{\text{cyt}}\) and/or aberrant subcellular compartmentation of \([\text{Ca}^{2+}]\) due to the inability to move excess cytosolic \([\text{Ca}^{2+}]\) into the vacuole. Although approximately 50% of wild-type vacuolar \([\text{Ca}^{2+}]_{\text{H}^+}\) transport activity was reported for \(cax1/cax3\) following pretreatment with exogenous \([\text{Ca}^{2+}]\) (Cheng et al., 2005), whether the \([\text{Ca}^{2+}]_{\text{cyt}}\) level is indeed increased in \(cax1/cax3\) has not yet been demonstrated.

Recently, Conn et al. (2011b) convincingly demonstrated that the severe phenotypes of \(cax1/cax3\), such as reduced leaf growth rate, mainly result from the increased leaf apoplastic free \([\text{Ca}^{2+}]\) rather than from the reduced vacuolar \([\text{Ca}^{2+}]\) of mesophyll cells. Their findings prompted us to suspect that the abrogated cellular \([\text{Ca}^{2+}]\) homeostasis in the leaf of \(cax1/cax3\) brings about misregulation of \(\text{Pi}\) homeostasis. As qRT-PCR analysis revealed that the transcript level of \(CAX1\) in shoot was decreased upon \(\text{Pi}\) limitation, it is likely that the leaf vacuolar sequestration of \([\text{Ca}^{2+}]\) by \(CAX1\) is down-regulated under \(\text{Pi}\) deficiency (Fig. 4A). We thus surmised that simultaneous loss of \(CAX1\) and \(CAX3\), a gene ectopically expressed in leaves upon knockout of \(CAX1\) (Cheng et al., 2003, 2005), may mimic the demands on reduction of the shoot vacuolar \([\text{Ca}^{2+}]_{\text{H}^+}\) transport activity under \(\text{Pi}\) deficiency, accompanied by activation of a subset of shoot PSR genes (Fig. 5B). By contrast, since \(CAX4\) is primarily expressed in root tissues and up-regulated in the \(cax1\) mutant (Cheng et al., 2003), the increased expression of \(CAX4\) may compensate for the functional role of \(CAX1\) and \(CAX3\) in the root of \(cax1/cax3\), hinting at why activation of the PSR genes was not observed in the root of \(cax1/cax3\) (Fig. 5B). In fact, overexpression of \(CAX4\) was able to partially suppress the \(cax1\) defect in vacuolar \([\text{Ca}^{2+}]_{\text{H}^+}\) transport (Zhao et al., 2009).

Notably, \(cax1/cax3\) displays 47% and 20% reductions, respectively, in \(V\)-ATPase and \(P\)-ATPase activities (Cheng et al., 2005; Zhao et al., 2008). As these \(H^+\)-ATPases generate a \(pH\) gradient across membranes that provides the driving force for the \(H^+\)-coupled transporters and contributes to the maintenance of the cytosolic \(\text{pH}\) homeostasis, whether the complex interaction of \(H^+\)-ATPases with other transporters and/or the resulting impaired \(\text{pH}\) homeostasis is associated with the activation of \(P_i\) transporters remains obscure. It is important to note that the \(\text{vha-a2/vha-a3}\) double mutant, which lacks the tonoplast \(V\)-ATPase, was shown to contain reduced \([\text{Ca}^{2+}]\) levels in leaves and to display symptoms of \(\text{Ca}^{2+}\) deficiency similar to \(cax1/cax3\) (Krebs et al., 2010). Moreover, it has been demonstrated that cytosolic \([\text{Ca}^{2+}]\) homeostasis is a constitutive function of the yeast \(V\)-ATPase. Cellular responses to a brief \([\text{Ca}^{2+}]\) challenge were affected not only by an acute loss of \(V\)-ATPase activity (in temperature-sensitive \(\text{vma}\) mutants or in wild-type cells treated with a \(V\)-ATPase inhibitor) but also by a permanent loss of \(V\)-ATPase activity in a \(\text{vma}\) deletion mutant ( Förster and Kane, 2000). In the future, it would be interesting to determine whether \(V\)-ATPase contributes to the impaired \(P_i\) homeostasis of the \(cax1/cax3\) mutant through the misregulation of \([\text{Ca}^{2+}]\) homeostasis.

Figure 6. Gene expression of SPX1, SPX3, and PHT1;4 \(P_i\) transporter in the \(cax1/cax3\) mutant. A to C, qRT-PCR analysis of SPX1 (A), SPX3 (B), and PHT1;4 (C) in the shoot of wild-type (WT) and \(cax1/cax3\) plants under +\(P_i\) (+) or −\(P_i\) (−) conditions. The value for each gene is presented as the fold change relative to the expression of wild-type plants under +\(P_i\) conditions. Error bars represent \(s\) of biological replicates from three independent experiments. Data significantly different from the corresponding controls are as indicated (mutant versus the wild type, * \(P < 0.05\), ** \(P < 0.01\); \(P_i\) sufficient versus \(P_i\) deficient, * \(P < 0.05\), ** \(P < 0.01\); Student’s \(t\) test). D, RT-PCR analysis of members in the PHT1 gene family in the root of wild-type and \(cax1/cax3\) plants under +\(P_i\) (+) or −\(P_i\) (−) conditions.

Figure 7. Analysis of PHT1;1/2/3 protein in the root of \(pht1;1\) seedlings (A) and \(cax1/cax3\) hydroponically grown plants (B) under +\(P_i\) (+) and −\(P_i\) (−) conditions. The bottom panels show the protein staining on the membrane. WT, Wild type.
Besides high Pi accumulation, disturbance of other ion homeostases, such as increased levels of Mn²⁺ and Zn²⁺ and decreased levels of Mg²⁺, has also been reported for cax1/cax3 (Cheng et al., 2005). Although an interplay between these various ions within cax1/cax3 cannot be excluded from participating in Pi signaling, our results that Pi accumulation of cax1/cax3 was exacerbated upon exogenous supplement of Ca²⁺ support the hypothesis that leaf Ca²⁺ homeostasis is directly involved in Pi signaling. It is unclear why the growth retardation and Pi accumulation of cax1/cax3 can be alleviated when high concentrations of Mg²⁺ are added to growth medium. However, it was argued that with a supplement of Mg²⁺ to growth medium, more Mg²⁺ is sequestered to the vacuole from the cytoplasm to compensate for the reduced vacuolar [Ca²⁺] in cax1/cax3 (Conn et al., 2011a).

Intriguingly, it is known that Ca²⁺ tends to precipitate with Pi, rendering the soil P unavailable for plant acquisition (Hinsinger, 2001). This aspect is noteworthy, considering that the excess cytosolic Ca²⁺ in the shoot of cax1/cax3 may potentially decrease the available shoot P, which may alternatively but not perfectly explain why only a subset of the shoot PSR genes are activated in cax1/cax3 under +Pi conditions. However, x-ray microanalysis showed that Arabidopsis Col-0 plants preferentially accumulate Ca in the vacuoles of mesophyll cells but P within vacuoles of the epidermis and bundle sheath (Conn et al., 2011b). Ce1-specific compartmentation of these two elements makes this possibility unlikely.

Loss of Function of CAX1 and CAX3 Disturbs the Regulation of PSR and Calcium-Related Gene Expression in the Shoot

Under +Pi conditions, one-fifth of the PSR genes are constitutively activated in the shoot of cax1/cax3,
including acid phosphatase type 5 (ACP5; At3g17790),
digalactosyl diacylglycerol synthase (DGD1; At3g11670),
phospholipase D2 (PLD2; At3g05630), SPX1, and
SPX3 (Table I). ACP5 has been proposed to be in-
volved in Pi mobilization (del Pozo et al., 1999),
whereas DGD1 and PLD2 participate in the biosyn-
thesis of nonphosphorus lipids during Pi-limited
growth (Härter et al., 2000; Cruz-Ramirez et al.,
2006). Loss of PLD2 and PLD2 reduces primary root
elongation under low-Pi conditions (Li et al., 2006),
but their roles in the shoot remain unknown. Nucleus-
localized SPX1 was suggested to be involved in transcrip-
tional activation of genes related to Pi mobilization
and scavenging of reactive oxygen species in response
to Pi starvation (Duan et al., 2008). By contrast, SPX3
was shown to localize to intracellular compartments.
SPX3 RNA interference lines exhibited higher total Pi
and Pi contents yet stronger induction of several PSR
genes, including SPX1, suggesting that SPX3 nega-
tively regulates Pi starvation signaling (Duan et al.,
2008). Up-regulation of both genes in the shoot of
cax1/cax3 under +Pi conditions may reflect the activation of
the SPX1/SPX3-mediated Pi signaling pathway. More-
over, a group of genes encoding Ca2+-ATPases, includ-
ing ACA2 (At4g357640), ACA10 (At4g29900), ACA11
(At3g57330), ACA12 (At3g63380), and ACA13 (At5g22100),
were up-regulated in the shoot but not in the root
of cax1/cax3 (Supplemental Table S2), implying that
Ca2+-ATPases are activated to balance the decreased
Ca2+ efflux in the shoot due to the dysfunction of
CAX1 and CAX3. Therefore, it seems that the disturbed

Cellular Ca2+ homeostasis and misregulated expres-
sion of Ca2+-related proteins caused by lack of CAX1
and CAX3, rather than CAX1 and CAX3 per se, are
involved in transcriptional regulation of PSR genes.

Common and Distinct Pi, Signaling Pathways Mediated
by PHO2 and CAX

Because both pho2 and cax1/cax3 mutants displayed
an increased level of shoot Pi, it is tempting to know
whether PHO2 and CAX1/CAX3 act in the same Pi
signaling pathway. Down-regulation of PHO2 in the
roots results in increased uptake and root-to-shoot
translocation of Pi; however, the role of PHO2 in the
shoots is unclear (Lin et al., 2008). Here, we observed
that the PHO2 transcript level was greatly increased in
the shoot of cax1/cax3 under +Pi and −Pi conditions,
whereas the level in the root of cax1/cax3 under +Pi
conditions was slightly reduced but still higher than
that in the −Pi root of wild-type plants (Fig. 4C). Given
that, under Pi deficiency, the level of PHO2 transcript
in the root of cax1/cax3 was as low as that in the root of
wild-type plants (Fig. 4C), we argue that PHO2 is not
involved in enhancing the Pi uptake activity in cax1/
cax3 under −Pi conditions. Although both CAX1/CAX3
and PHO2 inhibited the root Pi uptake activity, they
seem to mediate different pathways leading to Pi
accumulation. First, in contrast to pho2, which exhibits
a dramatically increased Pi uptake activity only when
Pi remains adequate, cax1/cax3 displayed an enhanced
Pi uptake activity under both +Pi and −Pi conditions
(Fig. 2, A and B). Notably, although the increase of Pi
uptake activity of cax1/cax3 under −Pi conditions was
striking, the underlying mechanism is not clear at
present. Second, unlike the pho2 mutant, the cax1/cax3
mutant had a similar shoot-to-root ratio of Pi distribu-
tion as the wild type under +Pi conditions (Fig. 2C).
Third, up-regulation of miR399, which acts upstream
of the PHO2-dependent Pi signaling pathway, was not
involved in the moderate reduction of PHO2 mRNA
level in the root of cax1/cax3 under +Pi conditions
(Fig. 4, C and D). These lines of evidence suggest that
CAX1/CAX3 and miR399-mediated PHO2 Pi signal-
ning pathways are two distinct pathways attributed to
the consequences of Pi uptake and accumulation.

In the pho2 mutant, the expression levels of PHT1;8
and PHT1;9 were shown to be increased under +Pi
conditions and assumed to contribute to the establish-
ment of high Pi in the shoot of pho2 (Aung et al., 2006;
Bari et al., 2006). However, in the root of cax1/cax3
under both +Pi and −Pi conditions, no up-regulation
of PHT1 genes at the transcript level was observed
(Fig. 6D). Although the protein level of PHT1;1/2/3
was not increased (Fig. 7B), our results from grafting
demonstrate that PHT1;1 is partially responsible for Pi
accumulation in the cax1/cax3 scion (Fig. 8C), indicat-
ing that a posttranslational regulation of PHT1;1 may
be involved. Several consensus sites for N-linked gly-
cosylation and phosphorylation have been predicted
in PHT1;1 (Muchhal et al., 1996). Indeed, PHT1;1/2

\[ \text{Ca}^{2+}_{\text{cyt}} \]

\[ \text{Ca}^{2+}_{\text{compartmentation}} \]
have been identified in phosphoproteomics of the Arabidopsis plasma membrane, and a phosphorylation site was detected in its C-terminal peptide (Nühse et al., 2004; Hem et al., 2007). Furthermore, as we did not examine the subcellular localization of PHT1;1 in cax1/cax3, changes in the membrane distribution of PHT1;1 cannot be excluded.

CAX1/CAX3-Mediated Shoot-to-Root Pi Signaling

A paradigm for systemic regulation of Pi homeostasis has been recently established. The miR399 generated in shoots after the onset of Pi starvation serves as a long-distance signal to activate P1 transport systems by suppressing PHO2 expression in roots (Lin et al., 2008; Pant et al., 2008). Our results obtained in split-root experiments indicate that CAX1 and CAX3 do not mediate the generation and movement of systemic suppressors from the +P root half to the −P root half. By contrast, the results of reciprocal grafting experiments clearly suggest the involvement of CAX1 and CAX3 in shoot-to-root Pi signaling. We hypothesize that abrogation of CAX1 and CAX3 in the cax1/cax3 mutant may relieve the repression of shoot PSR genes through alteration of \([Ca^{2+}]_{\text{cyt}}\) and/or misregulated compartmentation of Ca2+, thereby triggering a systemic signal that moves from shoots to roots to activate the PHT1 P1 transporters or other unidentified Pi transport systems (Fig. 9). Because mature miR399 was not observed in the shoot and root of cax1/cax3 under +P conditions, it is clear that miR399 is not up-regulated to serve as a systemic signal traveling to roots in the CAX1/CAX3-mediated Pi signaling pathway. Several molecules, including hormones, sugars, nutrients themselves or their metabolites, and small RNAs, have been suggested as systemic signals in the long-distance signaling of nutrient status (Liu et al., 2009). In the future, it will be interesting to identify such a shoot-derived signal and the molecular components involved in the up-regulation of the P1 transport system and to establish how those PSR genes misregulated in cax1/cax3 can modulate the P1 transport activity.

MATERIALS AND METHODS

Plant Materials and Growth Conditions

Seeds of Arabidopsis (Arabidopsis thaliana) cax1, cax3, and cax1/cax3 were kindly provided as gifts by Drs. Kendal Hirschi and Ning-Hui Cheng (Baylor College of Medicine). Seeds of the wild type (Col-0) and the ph1;1 T-DNA line (SALK_088586) were obtained from the Arabidopsis Biological Resource Center. The +P and −P media were supplemented with 250 μM KH2PO4 and 0 μM KH2PO4, respectively, unless specified otherwise. For hydroponic growth, 9-d-old seedlings grown on agar plates with half-modified Hoagland nutrient solution containing 250 μM KH2PO4 and 1% Suc solidified with 0.8% agar were transferred to the same nutrient solution containing 250 μM KH2PO4 without Suc for 8 to 10 d. A 5-d treatment of P1 starvation was initiated by replacing 250 μM KH2PO4 with P1-free medium. For plants grown in the medium supplemented with Ca2+ and Mg2+, 9-d-old seedlings grown on agar plates were transferred to hydroponic medium supplemented with various concentrations of Ca2+ and Mg2+ as indicated and grown for another 12 d. All plants were grown under a 16-h-light/8-h-dark cycle.

Grafting of Arabidopsis Plants

Hypocotyl reciprocal grafting was performed as described previously with minor modifications (Lin et al., 2008). Briefly, micrografting was conducted with 8-d-old seedlings, which were then incubated vertically in the dark for 1 d before being transferred to the culture room under dim light for another 2 d. Two weeks after micrografting, plants were transferred to hydroponic culture and grown for another 2 weeks before sample collection. Lack of contamination of adventitious roots in grafted plants was confirmed by genotyping using PCR.

Affymetrix ATH1 Array Hybridization and Data Analysis

Transcriptomic analyses of plants were conducted using Affymetrix ATH1 arrays. Wild-type and cax1/cax3 plants grown in hydroponic cultures under +P and −P conditions (see above) were harvested for RNA isolation. Two independent biological replicates were performed. Ten micrograms of total RNA (see below) was used for cDNA synthesis, labeled by in vitro transcription, and followed by fragmentation according to the manufacturer’s recommendations (GeneChip Expression Analysis Technical Manual, Revision 5; Affymetrix). The labeled samples were hybridized to the ATH1 array at 45°C for 16.5 h. Washing and staining were done on a Fluidics Station-450, and the ATH1 array was scanned using the Affymetrix GeneChip Scanner 7G. The results were quantified and analyzed using MicroArray Suite 5.0 software (Affymetrix). The obtained data were normalized using Robust Multichip Average (Irizarry et al., 2003), and the statistical significance of differential expression was determined by Limma analysis (Smyth, 2004).

P1 Concentration and P1 Uptake Analysis

P1 concentration and uptake activity were determined as described (Chiou et al., 2006). To assay the P1 uptake, 4-week-old plants grown under +P or −P conditions were transferred to medium containing 250 μM KH2PO4 (+P) or 10 μM KH2PO4 (−P) for the measurement of \([^{32}P]P1\) uptake.

RNA Isolation, RT-PCR, and qRT-PCR

Total RNA from hydroponic samples was isolated by the use of TRIzol reagent (Invitrogen) and treated with DNase I (Ambion) before qRT-PCR to eliminate genomic DNA contamination. cDNA was synthesized from 1 to 1 μg of total RNA by use of Moloney murine leukemia virus reverse transcriptase (Promega) with oligo(dT) primer. RT-PCR conditions and sequences of primers used in our study were identical to those listed in supplemental table S1 of Aung et al. (2006). Sequences of additional primers are listed in Supplemental Table S3. qRT-PCR was performed using the Power SYBR Green PCR Master Mix kit (Applied Biosystems) on a 7300 Real-Time PCR system (Applied Biosystems) according to the manufacturer’s instructions. Relative expression levels were normalized to that of an internal control, UBQ10.

Immunoblot Analyses

For extraction of total protein, 10-d-old seedlings of wild-type and ph1;1 plants with or without 5-d treatment of P1 deficiency were ground in liquid nitrogen and dissolved in protein lysis buffer (2% SDS, 60 mM Tris-HCl [pH 8.5], 2.5% glycerol, 0.13 mM EDTA, and 1× complete protease inhibitor [Roche]). Twenty micrograms of total protein was loaded onto the SDS-PAGE apparatus for each sample. Hydroponically grown cax1/cax3 and wild-type plants were harvested for total membrane protein extraction. One milligram of root tissues was ground with an ice-cold mortar and pestle and dissolved in 3 ml of membrane extraction buffer (330 mM Suc, 50 mM Tris [pH 7.5], 10 mM KCl, 5 mM EDTA, 5 mM dithiothreitol, 1 mM phenylmethylsulfonyl fluoride, and 1× complete protease inhibitor [Roche]). The extracts were then collected and centrifuged at 2,000g for 10 min at 4°C. Supernatants were collected and centrifuged at 400,000g for another 40 min at 4°C. Pellets were dissolved in membrane extraction buffer and collected as total membrane proteins. Twenty micrograms of total membrane protein was loaded onto the SDS-PAGE apparatus for each sample. Polyclonal rabbit antibodies were raised and affinity purified.
against an internal fragment of PHT1;1 corresponding to amino acid residues 266 to 285 (ELEEEVDDKVPDQRNYGLF). The final concentration of 20 to 100 ng mL−1 affinity-purified antibodies was used for immunoblot analysis.

Sequence data from this article can be found in the GenBank/EMBL data libraries under the following accession numbers: UBQ10 (At4g05320), PHT1;1 (At5g33390), PHT1;2 (At5g33370), PHT1;3 (At5g33360), PHT1;4 (At2g38940), PHT1;5 (At2g32830), PHT1;7 (At3g54700), PHT1;8 (At1g28060), PHT1;9 (At1g76320), PHO2 (At2g37770), CAX1 (At2g38170), CAX3 (At3g1860), At4 (At5g03545), SPX1 (At2g19150), and SPX3 (At2g45130).

Supplemental Data

The following materials are available in the online version of this article.

Supplemental Figure S1. Comparison of 163 and 455 differentially expressed genes (more than 2-fold change) in wild-type shoots and roots under –Pi conditions, respectively, with those reported by Morcuende et al. (2007).

Supplemental Figure S2. qRT-PCR analysis of SPX1 (A and B), PHT1;4 (C and D) and At4 (E and F) expression in the shoot (A, C, and E) and root (B, D, and F) of split-root wild-type and cax1/cax3 plants.

Supplemental Figure S3. qRT-PCR analysis of several PSR genes in the shoot of wild-type and cax1/cax3 plants under –Pi, or –P conditions.

Supplemental Table S1. Genes significantly differentially expressed (P ≤ 0.01, more than 2-fold change) in data sets derived from microarray analysis.

Supplemental Table S2. Ca2+–related genes that are differentially expressed in the shoot and root of cax1/cax3 under P-sufficient conditions.

Supplemental Table S3. Sequences of qRT-PCR primers used in this study.

ACKNOWLEDGMENTS

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LITERATURE CITED

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