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PDX1.1-dependent biosynthesis of vitamin B₆ protects roots from ammonium-induced oxidative stress

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Running title: Vitamin B₆ alleviates ammonium toxicity

Keywords: ammonium nutrition, apoplastic pH, Fe mobilization, root elongation, pyridoxine, ROS scavenging.

SHORT SUMMARY: Ammonium supply triggers a localized Fe-dependent oxidative burst that arrests primary root elongation but also induces PDX1.1-dependent vitamin B₆ production to quench ammonium-induced ROS formation.

ABSTRACT

Despite serving as a major inorganic nitrogen source for plants, ammonium causes toxicity at elevated concentrations, inhibiting root elongation early on. While previous studies have shown that ammonium-inhibited root development relates to ammonium uptake and formation of reactive oxygen species (ROS) in roots, it remained open which mechanisms are underlying the repression of root growth and how plants cope with this inhibitory effect of ammonium. Here, we demonstrate that ammonium-induced apoplastic acidification co-localizes with Fe precipitation and hydrogen
peroxide (H₂O₂) accumulation along the stele of the elongation and differentiation zone in root tips, indicating Fe-dependent ROS formation. By screening ammonium sensitivity in T-DNA insertion lines of ammonium-responsive genes, we identified PDX1.1, which is upregulated by ammonium in the root stele and catalyzes biosynthesis de novo of vitamin B₆. Root growth of pdx1.1 mutants is hypersensitive to ammonium, while chemical complementation or overexpression of PDX1.1 restores root elongation. This salvage strategy requires non-phosphorylated forms of vitamin B₆ that are able to quench reactive molecular oxygen species and rescue root growth from ammonium inhibition. We propose PDX1.1-mediated synthesis of non-phosphorylated B₆ vitamers as a primary strategy to protect roots from ammonium-dependent ROS formation.
INTRODUCTION

Nitrogen (N) is an essential mineral element for plant development and extensively applied in crop production (Xu et al., 2012). While synthetic N fertilizers greatly improve global crop yield, the input of nitrate-based N fertilizers also bears environmental risks when leached as nitrate or emitted as nitrogen oxide (Ju et al., 2009; Sutton et al., 2011). Since ammonium is less prone to leaching than nitrate, ammonium-based N fertilizers are widely used nowadays to replace nitrate in agricultural plant production. To increase fertilizer use efficiency, ammonium is often supplied in locally restricted fertilizer strips, where it is present at very high concentrations (Nkebiwe et al., 2016). Although ammonium is a preferential inorganic N source for many plant species (Gazzarrini et al., 1999), excessive ammonium causes toxicity resulting in leaf chlorosis and suppressed root growth (Britto and Kronzucker, 2002). These symptoms rely on ammonium triggering multiple physiological and morphological responses, including changes in apoplastic and cytosolic pH, gene expression, protein modification, ion transport, N metabolism, redox and phytohormone status, as well as root system architecture (Britto and Kronzucker, 2002; Li et al., 2014; Liu and von Wirén, 2017). Against this background and with the increasing use of ammonium-based N fertilizers in agricultural nutrient management, it is important to obtain a deeper understanding of how roots adapt to an ammonium-replete environment.

The most typical morphological changes in root system architecture of ammonium-exposed roots are arrested elongation of primary and lateral roots and enhanced branching of lateral roots (Li et al., 2010; Lima et al., 2010; Liu et al., 2013). Regarding lateral root branching, it has recently been shown that apoplastic acidification caused by AMMONIUM TRANSPORTER (AMT)-mediated ammonium uptake provokes pH-dependent radial auxin diffusion to stimulate locally the emergence of lateral root primordia, forming a highly branched root system (Meier et al., 2020). Given that in several auxin-related mutants root elongation is still as sensitive to ammonium as in wild-type plants, the repression of root elongation by ammonium is unlikely due to altered distribution or action of auxin in plants (Li et al., 2010; Liu et al., 2013). Thus, altered root elongation and lateral root branching under ammonium supply are most likely governed by distinct mechanisms. Neither root elongation nor lateral root branching can be triggered by glutamine, the primary assimilation product of ammonium (Lima et al., 2010; Rogato et al., 2010),
suggesting that ammonium itself or ammonium-related metabolites or signals are critical for root development (Liu and von Wirén, 2017). Indeed, excessive ammonium accumulation and hypersensitive root growth of the cipk23 mutant indicated that ammonium-induced calcium signaling is required to re-adjust internal NH₄⁺/K⁺ ratios (Shi et al., 2020). This refers to the function of CIPK23, a CBL-interacting protein kinase, in modulating AMT-, HAK- and AKT-type transporter activities (Ragel et al., 2015; Straub et al., 2017; Sanchez-Barrena et al., 2020; Dong et al., 2021) with strong impact on cytosolic ammonium accumulation and root growth. A further ammonium-related signal may be ethylene, which appears to modulate genes involved in ammonium uptake and assimilation via the transceptor function of NRT1.1 (Jian et al., 2018). Even though these signaling processes help maintaining cytosolic ammonium homeostasis, they do not explain ammonium-dependent root growth inhibition. Horizontal split agar experiments indicate that the critical region to sense the inhibitory effect of ammonium on root elongation is the root tip (Li et al., 2010). At the cellular level, both cell division and cell elongation are substantially repressed by ammonium, while the integrity of the root stem cell niche remains unaffected (Liu et al., 2013). On the search for mutants overcoming ammonium-induced root growth inhibition, the vtc1-1 mutant was isolated, which is defective in GMP-mannose pyrophosphorylase (GMPase) and thus not only in downstream ascorbate biosynthesis but also in N-glycosylation of proteins (Qin et al., 2008). However, despite its significant impact on ammonium fluxes (Li et al., 2010), GMPase activity turned out not to be the primary cause of ammonium-dependent root growth inhibition (Kempinski et al., 2011). Transcriptome studies have indicated that enhanced formation of ROS is a typical ammonium-related physiological response (Patterson et al., 2010). Even though significantly higher steady-state H₂O₂ levels in roots were not detected, antioxidative defense systems were induced that comprise ROS-degrading enzymes, including superoxide dismutase, catalase, peroxidase, guaiacol peroxidase and glutathione reductase (Patterson et al., 2010). However, in leaves ammonium toxicity increased H₂O₂ formation, which was found to be mediated by ethylene and followed by enhanced peroxidase activity probably via ABA signaling (Li et al., 2019; Sun et al., 2020). Also in the roots of rice, ammonium treatment increased H₂O₂ contents and induced the heme-heme oxygenase OsSE5 together with other ROS-detoxifying enzymes including superoxide dismutase, catalase, and ascorbate
peroxidase to relieve ammonium-supplied plants from oxidative stress (Xie et al., 2015). These studies indicate that ammonium toxicity causes ROS formation and that the capacity of enzymatic ROS detoxification is a major determinant of ammonium tolerance in plants. So far, evidence for the involvement of non-enzymatic antioxidants in ammonium-induced oxidative stress defense was not obtained, considering that in roots the pool sizes and oxidative states of NADH, NADPH, glutathione and ascorbate were not substantially altered by ammonium supply (Patterson et al., 2010).

To explore the mechanisms by which ammonium-exposed roots cope with ammonium toxicity and ammonium-dependent ROS formation, we first used a pharmacological approach unraveling localization of the ammonium-triggered H$_2$O$_2$ burst. We then screened mutants of ammonium-responsive genes in roots and identified PDX1.1, a gene involved in the biosynthesis de novo of vitamin B$_6$. Chemical complementation and genetic approaches showed that non-phosphorylated forms of vitamin B$_6$ suppressed H$_2$O$_2$ formation under ammonium supply. With PDX1.1-dependent vitamin B$_6$ formation, our study identifies a protective mechanism that spatially overlaps with ammonium-triggered H$_2$O$_2$ formation in inner root cells and thus carries potential to better adapt plant roots to ammonium-based fertilization strategies.

**RESULTS**

**Accumulation of H$_2$O$_2$ under ammonium supply inhibits primary root elongation**

When wild-type seedlings of Arabidopsis were grown on half-strength MS medium for 6 d in the presence of either 1 or 10 mM ammonium as the sole N source, primary root length became 25 or 35% shorter, respectively, than that of nitrate-grown plants (Fig S1A, B). Likewise, total and mean length of lateral roots were repressed to a similar extent as primary root length, suggesting a common mechanism underlying ammonium-induced growth inhibition in both root types (Fig S1B-D). A significant dose-dependent decrease in root elongation rate set in already after 1 d of ammonium exposure suggesting a rapid inhibitory mechanism (Fig S1E-G). Root exposure to ammonium decreased both, cortical cell length and meristem size in a dose-dependent manner and to a similar extent (Fig S1H-J). Since cytokinins determine root meristem size by controlling cell differentiation (Dello Ioio et al. 2007),
we checked the response of cytokinin-sensitive reporter line TCS:GFP in the primary root apex, which showed highest expression in the columella and lateral root cap and slight but significant repression under ammonium, revealing that cytokinin signaling was not enhanced by ammonium (Bielach et al., 2012; Fig S2A, B). The histidine kinase AHK3 acts as a cytokinin receptor to stimulate a two-component signaling pathway that transfers a phosphorelay signal to the nucleus, where transcription factors of the type-B ARABIDOPSIS RESPONSE REGULATOR gene family (ARR-B) are activated (Ferreira and Kieber, 2005). We therefore examined ammonium sensitivity via measuring primary root length and meristem size in the ahk3-3, arr1-3 and arr12-1 mutants, which however revealed no difference to the wild-type (Fig S2C-F), indicating that the AHK3/ARR1 and AHK3/ARR12 two-component cytokinin signaling pathway was not targeted by ammonium. SHY/IAA3 acts directly downstream of ARR1 and ARR12, which increases cell differentiation rate and balances root-meristem size at the transition zone (Dello Ioio et al. 2008). However, neither the SHY2 deletion line shy2-31 nor shy2-2, which expresses a stabilized proteoform of SHY2 (Tian et al., 2002), exhibited altered ammonium sensitivity of primary root elongation or meristem formation (Fig S2G-J). Taken together, these observations indicated that cytokinin signaling is not involved in ammonium-dependent inhibition of root elongation.

As ammonium has been reported to enhance ROS levels in plants (Patterson et al., 2010; Xie et al., 2015), we stained H$_2$O$_2$ in primary roots by 3,3'-diaminobenzidine (DAB). Compared with nitrate, increasing ammonium supply gradually enhanced repression of cell length and meristem size with the detection of H$_2$O$_2$ particularly in the elongation and differentiation zones of the root apex (Fig 1A-G). Since nitrate-supplied plants also showed similar DAB staining as ammonium-exposed plants in the meristematic root zone, this zone was not considered as a primary target of ammonium-induced ROS production. To better understand the source of ammonium-dependent ROS formation in primary roots, we employed a pharmacological approach and first applied potassium iodide (KI), acting as a chemical H$_2$O$_2$ scavenger (Lee et al., 2013). KI largely reverted the ammonium-dependent inhibition of the primary root elongation rate and restored cell length and meristem size to a large extent (Fig 1A-F). Recovered root elongation coincided with lower abundance of DAB-stained ROS in the elongation and differentiation zone of the primary root.
(Fig 1G), suggesting that enhanced levels of H$_2$O$_2$ are responsible for the stunted root phenotype. When ammonium-grown roots were exposed to elevated H$_2$O$_2$ levels, which were generated either directly by supply of 1 mM H$_2$O$_2$ or indirectly by supply of the peroxidase inhibitor salicylhydroxamic acid (SHAM), primary root elongation rates declined drastically due to strongly decreased cell length and meristem size, going along with enhanced DAB staining (Fig S3A-G). Notably, in nitrate-supplied roots, meristem size and DAB staining were apparently not affected by H$_2$O$_2$ or SHAM, while cortical cell length was only slightly decreased (Fig S3D-G), relating to the fact that nitrate-supplied plants possess elevated ROS scavenging capacities (Chu et al., 2021). To distinguish between the contribution of superoxide radicals (O$_2^-$) and H$_2$O$_2$, we investigated the effects of sodium diethyldithiocarbamate (DDC), an inhibitor of Cu,Zn-superoxide dismutase (SOD) that blocks H$_2$O$_2$ generation and leads to O$_2^-$ accumulation (Auh and Murphy, 1995). Both cell length and meristem size of the primary root apex were partially but significantly rescued by DDC supplementation, allowing partial recovery of primary root elongation (Fig S4). This observation indicated that O$_2^-$ radicals are not the direct cause for ammonium-mediated inhibition of root elongation, whereas SOD-catalyzed conversion of O$_2^-$ to H$_2$O$_2$ matters. Alternatively, we suspected RBOH-type NADPH oxidases produce ammonium-dependent ROS and examined a couple of rboh single or multiple knock-out lines. However, none of them showed an ammonium-dependent root phenotype (Fig S5).

Since in the absence of ammonium H$_2$O$_2$ was less effective in inhibiting root elongation (Fig S3), we assumed ammonium-facilitated formation of H$_2$O$_2$ plays a role and speculated that this process depends on the availability of Fe (Dixon and Stockwell, 2014). Indeed, lowering Fe supply from 100 µM to 10 µM Fe(III)-EDTA resulted in weaker inhibition of cell length, meristem size and primary root elongation and prevented accumulation of DAB-stained ROS in the elongation and differentiation zone of the primary root (Fig 1A-G). Higher resolution of DAB-stained cells allowed assigning H$_2$O$_2$ to the stele of the late elongation and early differentiation zone, which became shorter under increasing ammonium supply and thus appeared more apical (Fig 2A). In addition, we localized ROS-dependent fluorescence by the cell-permeant indicator 2’,7’-dichlorofluorescin diacetate (H$_2$DCFDA), revealing enhanced ROS formation in the stele, especially along the vascular strands of ammonium-exposed roots with a maximum at the transition of the
elongation to the differentiation zone (Fig 2B, C). Considering Fe dependency of ammonium-triggered root inhibition (Fig 1), Fe distribution in the root was stained by DAB-enhanced Perl's detecting both free Fe(II) and Fe(III) (Roschztardtz et al., 2009). While nitrate-grown roots accumulated some Fe in the meristematic zone, ammonium-exposed roots showed much higher Fe accumulation especially in the differentiation zone (Fig 2D). Cross sections of Perl's/DAB-stained primary roots showed that ammonium-dependent Fe precipitation was restricted by the Casparian band and thus confined to the apoplast of outer root cells (Fig S6). Since Fe availability increases with decreasing pH and ammonium nutrition is known to acidify the rhizosphere and apoplast (Römheld and Marschner, 1986; Meier et al., 2020), apoplastic pH changes were traced by using the apo-pHusion line, a ratiometric reporter of apoplastic pH (Gjetting et al., 2012). Increasing ammonium nutrition decreased apoplastic pH in particular in the elongation and differentiation zone of ammonium-exposed roots (Fig 2E, F), which co-localized with DAB- or fluorescence-stained ROS (Fig 2A-D), suggesting that the ammonium-dependent pH decrease and enhanced Fe availability caused ROS formation.

Since ammonium-induced repression of root elongation appears to be a rapid response detected even 1 d after ammonium exposure (Fig S1E-G), we investigated dynamic changes in Fe availability and ROS status in ammonium-supplied primary roots. Enhanced Fe precipitation and ROS accumulation appeared as early as 1 d after ammonium supply in the differentiation zone of the primary root and progressed in the apical direction with time and in dependence of external ammonium supply (Fig S7). As indicated by the arrows marking the boundaries of the meristematic and elongation zones, the size of the elongation zone started declining earlier than that of the meristematic zone, which went along with progression of H$_2$DCFDA-dependent fluorescence into the elongation zone towards the meristematic zone (Fig 2B; S7B). To verify an impact of ROS on cell division, we evaluated the response to ammonium of the cell cycle reporter CycB1;1::GUS (Colón-Carmona et al., 1999). Ammonium supply significantly suppressed GUS activity in the apical meristem of primary root, and this suppression was largely alleviated by lowering external Fe supply (Fig S8). Consistent with the decrease of meristem size and cell length under the same conditions (Fig 1D-F), we conclude that Fe-dependent ROS formation is a major repressor of both cell elongation and cell division in ammonium-exposed root apices. Notably, exactly the same responses of apoplastic pH, Fe availability and ROS
formation as in primary root were detected in the apex of lateral roots (Fig S9), indicating that oxidative stress acts as a common inhibitory mechanism underlying ammonium-induced growth repression in both primary and lateral roots.

To manipulate Fe availability in the presence of ammonium, medium pH was buffered at different values (Fig S10). While ammonium-inhibited root elongation was aggravated when roots were shifted from standard pH 5.7 to pH 5.0, higher pH restored root elongation leading to almost similar length of nitrate- and ammonium-grown roots at pH 7.2 (Fig 3A-C). After 6 d of ammonium exposure, Perls/DAB staining revealed decreasing Fe accumulation in the apical root zone with increasing pH, except for the columella cells (Fig 3D). In contrast, strongly enhanced amounts of Fe accumulated at pH 5.0 but only in ammonium-exposed roots. Since the pH of the agar medium was buffered by MES, we suspected that buffer strength might affect ammonium-induced pH changes and Fe availability under ammonium supply. Indeed, lowering MES concentration from 2.5 to 0.5 mM dramatically aggravated Fe precipitation and inhibition of ammonium-exposed primary root tips, while at 10 mM MES Fe precipitation decreased and ammonium-repressed root elongation was partially attenuated (Fig S11). These results indicated that the intensity of ammonium-dependent proton release and acidification of the rhizosphere is of immediate relevance for Fe mobilization and precipitation in the root apoplast and at the root surface.

The dependence of ammonium-induced root growth inhibition on Fe recalls the role of Fe in root growth inhibition under phosphate deficiency that also relies on Fe-facilitated ROS formation (Müller et al., 2015). Recently, it has been reported that this role of Fe represents a side effect of light exposure when roots are grown in transparent Petri dishes (Zheng et al., 2019). Thus, we shaded the whole root system by aluminium foil as described in Zheng et al. (2019) and noted that in all N treatments primary root length increased when roots were shielded from white light (Fig S12A, B). Even though ammonium-dependent primary root growth inhibition was slightly alleviated by shielding roots from light, ammonium supply still arrested primary root elongation to a large extent (Fig S12B, C). This observation supported that light increases the pool of redox-active Fe (Zheng et al., 2019), which in the presence of ammonium may engage in ROS formation and inhibit primary root elongation.
Previous work under phosphate deficiency showed that blue light exposure promotes the Fenton reaction converting H$_2$O$_2$ to hydroxyl radicals that are more toxic oxygen species and strongly suppress primary root elongation (Zheng et al., 2019). Here, to evaluate the involvement of hydroxyl radicals in ammonium-dependent root growth inhibition, we supplied thiourea, a chemical scavenger of hydroxyl radicals (Wasil et al., 1987), together with ammonium to the medium. However, the presence of thiourea hardly altered the sensitivity of primary roots to ammonium (Fig S13), indicating that the formation of hydroxyl radicals via the Fenton reaction is not critical for ammonium-dependent root growth inhibition.

**Hypersensitivity of root elongation to ammonium relates to vitamin B$_6$ deficiency**

To explore how plants cope with ammonium-induced H$_2$O$_2$ formation and subsequent root growth inhibition, we searched in publicly available transcriptome data for ammonium-responsive genes that are expressed in roots (Patterson et al., 2010; Ristova et al., 2016). We then collected T-DNA insertion lines of 29 ammonium-responsive genes and screened them for primary root length under ammonium versus nitrate supply (Data Set S1). In this screen, we identified a T-DNA insertion line of PDX1.1 (SALK_024245), which was hypersensitive to ammonium treatment (Fig S14A, B). In Arabidopsis, PDX1.1 (along with PDX2) catalyzes vitamin B$_6$ biosynthesis *de novo* (Tambasco-Studart et al., 2005; Fig S15), thus pdx1.1 mutants suffer from vitamin B$_6$ deficiency (Titiz et al., 2006; Wagner et al., 2006; Boycheva et al., 2015). We confirmed the hypersensitive phenotype of pdx1.1 mutants to ammonium by examining primary root growth in SALK_024245 and the transposon insertion line pdx1.1-1 (Figure S14C-E). Since SALK_024245 and pdx1.1-1 showed the same growth phenotype under ammonium supply in all our experiments, we focused in the following on SALK_024245, naming it pdx1.1-3. Given that under nitrate supply primary root growth of pdx1.1-3 was weaker (Fig 4A), we calculated the relative primary root elongation rate by normalizing it to the root growth rate under nitrate. An earlier and steeper decrease in root elongation rate of pdx1.1-3 than of the wild type confirmed hypersensitivity of pdx1.1-3 to ammonium nutrition (Fig 4B, C). The higher sensitivity of pdx1.1-3 to ammonium went along with elevated accumulation of H$_2$O$_2$ in ammonium-exposed primary root tips already 2 d after transfer to ammonium (Fig 4D).
To confirm whether the hypersensitivity of *pdx1.1-3* to ammonium is due to the defect in vitamin B₆ biosynthesis, we supplied vitamin B₆ in the form of pyridoxine to the growth medium. While external supply of pyridoxine to nitrate-grown wild-type plants had no effect, it largely alleviated ammonium-induced inhibition of primary root growth in the wild type and particularly in the *pdx1.1-3* mutant (Fig 4A). Notably, external supplementation of pyridoxine recovered completely the primary root elongation rate of *pdx1.1-3* (Fig 4B-C), cortical cell length and meristem size (Fig S16) as well as GUS activity of the *CycB1;1::GUS* reporter (Fig S17), confirming that the hypersensitive phenotype of *pdx1.1-3* to ammonium is due to defective vitamin B₆ biosynthesis. Since external supply of pyridoxine reduced also the accumulation of DAB-stained H₂O₂ in ammonium-treated primary roots of both lines (Fig 4E), we conclude that vitamin B₆ alleviates ammonium toxicity either by suppressing H₂O₂ generation or by scavenging H₂O₂ in primary root tips.

### Non-phosphorylated forms of vitamin B₆ are critical for ammonium detoxification

Vitamin B₆ is essential for all living organisms, and refers to a group of six different vitamers that contain a pyridine ring and include the non-phosphorylated forms pyridoxal (PL), pyridoxine (PN) and pyridoxamine (PM), as well as the phosphorylated forms pyridoxal 5'-phosphate (PLP), pyridoxine 5'-phosphate (PNP) and pyridoxamine 5'-phosphate (PMP) (Fitzpatrick, 2011; Fig S15). To evaluate their efficacy in mitigating root growth inhibition, we supplied different B₆ vitamers to the medium. Compared with the mock treatment, external supply of PLP slightly retarded the ceasing primary root elongation during the first 4 d after transfer to ammonium, but it hardly restored primary root length after 6 d of ammonium supply (Fig S18A-C). By contrast, exogenous application of the vitamin B₆ forms PL or PN effectively rescued primary root growth under ammonium supply, even during later stages of the treatment, indicating a superior efficacy of the non-phosphorylated B₆ vitamers. When monitoring the H₂O₂ status in root tips in parallel with root elongation, exogenous application of PN or PL but not of PLP quenched excess accumulation of H₂O₂ in ammonium-treated primary root tips (Fig S18D). These results reveal that mitigation of root growth inhibition refers to the non-phosphorylated forms of vitamin B₆ and their ability to suppress H₂O₂ accumulation.
Over-expression of PDX1.1 improves ammonium tolerance in roots

To address the question whether PDX1.1 plays a role in eliminating ammonium-triggered ROS, we inspected three independent PDX1.1 overexpression lines (Raschke et al., 2011). All three lines continued elongating their primary roots even after 6 d of ammonium exposure when root elongation of the wild-type was almost completely exhausted (Fig 5A-C). DAB staining showed that overexpression of PDX1.1 strongly reduced the level of H$_2$O$_2$ in ammonium-treated primary roots (Fig 5D). Accordingly, ammonium-induced inhibition of root cortical cell length, meristem size and activity of the CycB1;1::GUS reporter also recovered by overexpression of PDX1.1 (Fig 5E-G; Fig S19), revealing the efficacy of endogenous vitamin B$_6$ formation under ammonium nutrition. Moreover, an $^{15}$NH$_4^+$ influx experiment revealed that the ammonium uptake capacity in roots remained unaffected in the pdx1.1-3 mutant as well as in the PDX1.1 overexpression lines (Fig 5H). Taken together, elevated expression of PDX1.1 is sufficient to restore elongation in ammonium-exposed roots and acts downstream of the ammonium uptake process.

Overexpression of PDX1.1 increases the level of non-phosphorylated B$_6$ vitamers

As the PDX1-dependent step in vitamin B$_6$ biosynthesis yields first PLP (Fig S15), we hypothesized that overexpression of PDX1.1 will primarily increase phosphorylated rather than non-phosphorylated vitamers, whose generation requires an additional phosphatase reaction. Measuring vitamin B$_6$ in roots confirmed significantly lower levels of total vitamin B$_6$ in the pdx1.1-3 mutant than in the wild type, which was caused by a significant decrease in the concentrations of all five determined B$_6$ vitamers (Fig 6A, B). At first glance, this suggested a constitutive contribution of PDX1.1 to overall vitamin B$_6$ biosynthesis because vitamin B$_6$ levels decreased in pdx1.1-3 irrespective of the supplied N form. Overexpression of PDX1.1 led to constitutively higher total vitamin B$_6$ levels only in PDX1.1OE-L5, but under ammonium supply also in PDX1.1OE-L15 and to lesser extent in PDX1.1OE-L8, which coincided with elevated PDX1.1 transcript levels in these lines (Fig 6A-C). Surprisingly, the concentration of PLP was not enhanced in ammonium-supplied PDX1.1 overexpression lines, although PLP contributed > 70% to overall vitamin B$_6$ levels. In contrast, PDX1.1 overexpression lines showed significantly higher concentrations of non-phosphorylated vitamin B$_6$ forms than the wild-type, and these
differences closely reflected the relative differences in transcript levels among the
three overexpression lines. Hence, the proportion of individual B₆ vitamers shifted
towards the non-phosphorylated vitamin B₆ forms, among which PL was the most
abundant form contributing up to 25% of total vitamin B₆ (Fig 6B). PM accounted only
for approx. 1% of total vitamin B₆ in roots. In wild-type roots, 20% of total vitamin B₆
was in non-phosphorylated forms, while in pdx1.1-3 this proportion was less than
10%. In PDX1.1 overexpression lines, between 30% and 40% of total vitamin B₆ was
converted into non-phosphorylated forms and this share even increased when plants
were grown in the presence of ammonium (Fig 6A, B). We finally correlated the levels
of individual B₆ vitamers with those of PDX1.1 transcripts and revealed close
correlations for PN, PL, and PM, but not for PMP, whereas the correlation for PLP
was also significant although much weaker (Fig 6D). These results indicate that the
extent of PDX1.1 upregulation by ammonium or by ectopic expression determines
primarily the abundance of non-phosphorylated B₆ vitamers without compromising
homeostasis of phosphorylated B₆ vitamers in roots.

H₂O₂ triggers localized upregulation of PDX1.1 under ammonium supply
To investigate the transcriptional regulation of vitamin B₆ biosynthesis and
homeostasis by ammonium, the transcript levels of all known genes involved in
vitamin B₆ biosynthesis or metabolism were examined by quantitative real-time PCR
in roots grown under different N supply. Unexpectedly, PDX1.1 turned out to be the
sole vitamin B₆-related gene that was significantly upregulated by ammonium (Fig
7A). Next, the tissue-specific expression pattern of PDX1.1 was monitored in plants
expressing a PDX1.1 promoter-GUS reporter construct (Boycheva et al., 2015).
PDX1.1 reporter activity was upregulated by ammonium supply in the root
vasculature, increasing from the elongation zone basipetally through the
differentiation zone (Fig 7B). As the reporter activity became stronger in the presence
of 10 mM ammonium, upregulation of PDX1.1 expression apparently followed
ammonium supply in a dose-dependent manner (Fig 7A, B).

Since the spatial localization of PDX1.1 expression and H₂O₂ accumulation
strongly overlapped in ammonium-treated roots (Fig 2A-B, 7B), the question arose
whether ammonium itself or H₂O₂ triggered the upregulation of PDX1.1. We thus
suppressed H₂O₂ levels in ammonium-exposed roots by supply of KI or by low Fe
and observed that PDX1.1 reporter activity disappeared (Fig 7B). By contrast,
exogenous application of 1 mM H$_2$O$_2$ strongly enhanced the promoter activity of PDX1.1 in the root stele as well as in the apical root meristem, however only in the presence of ammonium (Fig 7C). Therefore, H$_2$O$_2$ serves as a signal to upregulate PDX1.1 transcription upon ammonium nutrition and determine its restricted expression in the stele.

Given that the change of pH upon ammonium exposure is the initial event to trigger the Fe-dependent oxidative burst, we hypothesized that the transcriptional response of PDX1.1 to ammonium should be influenced by medium pH. Indeed, when pPDX1.1:GUS reporter lines were shifted from pH 5.7 to 5.0, the upregulation of PDX1.1 was enhanced in the presence of ammonium. By contrast, when raising medium pH from 5.7 to 6.5 or 7.2, the induction of PDX1.1 by ammonium almost disappeared (Fig S20). These findings confirmed that ammonium- and pH-dependent H$_2$O$_2$ generation is required to induce the expression of PDX1.1 in roots.

**Elevation of PDX1.1-mediated vitamin B$_6$ biosynthesis improves tolerance to further nutrient-related stresses**

Apart from ammonium toxicity, other nutrient-related stresses also arrest root elongation by elevated ROS formation, such as phosphate deficiency or nickel toxicity (Müller et al., 2015; Zheng et al., 2019; Lesková et al., 2020). We wondered whether PDX1.1-mediated vitamin B$_6$ biosynthesis also counteracts these oxidative stresses in plants. First, we inspected published transcriptome results (Bhosale et al., 2018; Lesková et al., 2020), to assess the response of vitamin B$_6$-related genes to phosphate deficiency and nickel toxicity. Surprisingly, none of the known genes involved in vitamin B$_6$ biosynthesis or metabolism, including PDX1.1, was differentially expressed under phosphate deficiency or nickel toxicity (Fig S21A, B). Next, the arrest of primary root elongation under phosphate deficiency or nickel toxicity showed no difference between wild-type (Col-0) and pdx1.1-3 mutant plants (Fig S21C-F). These results suggested that vitamin B$_6$-dependent ROS protection did not evolve as a strategy to alleviate oxidative stress in response to phosphate deficiency or nickel toxicity in natural conditions. Nonetheless, coinciding with their enhanced ROS scavenging capacity, all three PDX1.1 overexpression lines exhibited significantly improved root elongation under phosphate deficiency or nickel toxicity (Fig S21C-F). This result suggests that enhanced PDX1.1-mediated vitamin B$_6$
biosynthesis can be applied as a practical strategy to improve the root tolerance to multiple types of oxidative stress.

**DISCUSSION**

Application of ammonium-based N fertilizers in agricultural plant production bears the risk of impaired root development when roots are exposed to ammonium-rich soil patches (Britto and Kronzucker, 2002; Watt et al., 2006). Plants have evolved several strategies to cope with the adverse effects of predominant ammonium nutrition, which comprise enhanced N assimilation in roots (Cruz et al., 2005; Guan et al., 2016; Konishi et al., 2017), ammonium compartmentalization to the apoplast or vacuole (Loqué et al., 2005; Li et al., 2010; Bai et al., 2014), and activation of enzymatic antioxidation systems to cope with ammonium-triggered ROS production (Patterson et al., 2010; Xie et al., 2015). In this study, we describe the mechanistic basis underlying ammonium-triggered ROS formation and identify with PDX1.1-dependent vitamin B₆ synthesis a metabolic defense response to counteract ammonium-induced ROS formation.

**Ammonium-dependent acidification triggers ROS formation via enhanced Fe mobilization**

Ammonium toxicity belongs to one of several mineral element disorders that associate root growth inhibition with the production of ROS (Fig 1; Xie et al., 2015). Among those disorders, localization and generation of ROS differ in an element-specific manner, pointing to different sources of ROS formation. For instance, root growth inhibition by nickel has been related to ROS formation in outer cells of the apical root zone (Leskova et al., 2020), while ROS accumulate predominantly in the meristem and elongation zone under zinc deficiency (Nakayama et al. 2020) or in the root vasculature from the meristematic through the elongation and differentiation zone of salt-stressed plants (Jiang et al., 2012). Considering that experimental visualization and localization of ROS depend not only on the site of generation but also on chemical properties of the ROS-sensitive dye or reporter, we used here DAB staining and H₂DCFDA-dependent fluorescence to localize ROS formation under ammonium nutrition primarily to the vascular cylinder or stele in the elongation and differentiation zones (Fig 2). With increasing ammonium supply and time of exposure ROS formation gradually progressed through the elongation zone towards the
meristem (Fig 2B-C; S7B), where it coincided with suppressed cell length, meristem size and cell division activity (Fig 1; S3; S8), revealing that the apical root meristem is not the primary target of ammonium-dependent ROS generation especially at early stages of ammonium exposure. Although cytokinins are crucial in balancing cell division and meristem size (Dello Ioio et al., 2008), key components of cytokinin signaling are apparently not involved in ammonium-dependent root growth inhibition (Fig S2).

Since ammonium-dependent root growth inhibition was suppressed when H$_2$O$_2$ accumulation was chemically quenched by KI (Fig 1) but enhanced when H$_2$O$_2$ accumulation was favored in the presence of SHAM or by H$_2$O$_2$ supplementation (Fig S3), H$_2$O$_2$ or a downstream product must have caused the stunted root phenotype. This is supported by the observation that blocking H$_2$O$_2$ formation by DDC in favor of O$_2^-$ accumulation prevented ammonium-dependent root growth inhibition (Fig S4).

On the other hand, high effectivity of SOD inhibition by DDC implied that DAB-stained H$_2$O$_2$ was generated via O$_2^-$, which in turn may derive from peroxisomes and the mitochondrial electron transport chain or from the activity of NADPH oxidases in the plasma membrane (Smirnoff and Arnaud, 2018). By examining a limited number of available rboh single or multiple knock-out lines, we could not find evidence for a role of RBOH-type NADPH oxidases in ammonium-triggered root growth inhibition (Fig S5), even though our assay also included the root stele-localized RBOHF, which triggers vascular ROS formation upon salinity as a prerequisite for salt tolerance (Jiang et al., 2012). Irrespective of the O$_2^-$ source, SOD-mediated dismutation of O$_2^-$ to H$_2$O$_2$ consumes protons and is favored by low apoplastic pH (Smirnoff and Arnaud, 2018) that resulted here from ammonium uptake-induced proton secretion (Fig 2; Fig 7D; Meier et al., 2020). Suppressed DAB staining and mitigation of root growth inhibition under low Fe supply (Fig 1) indicated a key role of Fe in H$_2$O$_2$ formation. Indeed, exposure of ammonium-grown roots to light, which increases the pool of redox-active Fe (Zheng et al., 2019), aggravated inhibition of root elongation (Fig S12). Furthermore, redox-active Fe(III) likely arose from apoplastic acidification and dissolution of apoplastic Fe pools (Fig S6; Zhu et al., 2018) as well as from UV-dependent photooxidation of EDTA that sets chelated ferric Fe free (Hangarter and Stasinopoulos, 1991). A similarly critical role of ferrous Fe in ROS-mediated root growth inhibition is also known for primary root growth inhibition under phosphate starvation (Müller et al., 2015; Zheng et al., 2019). In this context, exposure of P-
deficient roots to light favors reduction of ferric to ferrous Fe and subsequent Fe$^{2+}$-mediated formation of hydroxyl radicals via the Fenton reaction. DAB-stained H$_2$O$_2$ under P deficiency is confined to the meristematic root zone and further depends on malate efflux via ALMT1 (Zheng et al., 2019) to increase Fe solubility in the apoplast (Balzergue et al., 2017; Mora-Macías et al., 2017). Since the root elongation in response to ammonium was not altered in almt1 mutants (Fig S14A, B), root shading restored root elongation only in part (Fig S12) and ammonium-dependent H$_2$O$_2$ mainly localized to the root stele above the meristematic zone (Fig 2), mechanisms underlying ROS formation clearly differ between P-deficient and ammonium-exposed roots. Moreover, supplementation of the hydroxyl radical scavenger thiourea, which can restore arrested primary root elongation under P deficiency (Zheng et al., 2019), was poorly effective in the case of ammonium (Fig S13), suggesting that formation of hydroxyl radicals via the Fenton reaction is not critical for ammonium-dependent root growth inhibition. Nonetheless, a determinant role of redox-active Fe in ammonium-triggered root growth inhibition is corroborated by its gradual relief under increasing medium pH, which also decreased Fe accumulation in the root and subsequent Fe availability for ROS metabolism (Fig 3). We thus conclude that ammonium-induced acidification of the root apoplast and additionally in the rhizosphere enhance Fe solubilization as prerequisite for re-location to the stele and subsequent Fe-mediated formation of H$_2$O$_2$.

**Plant roots counteract ammonium-induced ROS formation via PDX1.1-mediated vitamin B$_6$ biosynthesis**

In plants, vitamin B$_6$ is synthesized via the “deoxyxylulose-5-phosphate (DXP)-independent pathway”, which utilizes ribose-5-phosphate, glyceraldehyde-3-phosphate and glutamine as substrates for the glutamine amidotransferase complex comprised of pyridoxine synthase (PDX1) and pyridoxine glutaminase (PDX2) (Tambasco-Studart et al., 2005). The primary product is the phosphorylated form PLP (Tambasco-Studart et al., 2005) before enzymes of the salvage pathway facilitate the interconversion among different B$_6$ vitamers (Colinas et al., 2016; Fig S15). While PLP acts as coenzyme in numerous enzymatic reactions, including those with importance for N assimilation (Percudani and Peracchi, 2003; Fitzpatrick, 2011; Colinas et al., 2016), non-phosphorylated forms of vitamin B$_6$ serve efficiently as antioxidants *in vitro* and *in vivo* (Bilski et al., 2000; Havaux et al., 2009; Mooney and
Hellmann, 2010). Our study provides several lines of evidence indicating that plants induce PDX1.1-mediated synthesis of non-phosphorylated B<sub>6</sub> vitamers as an efficient strategy to counteract ammonium-dependent oxidative stress in roots. First, as several enzymes are required for vitamin B<sub>6</sub> biosynthesis and homeostasis, it was surprising that only one of them, PDX1.1, responded to ammonium with enhanced expression (Fig 7A). Indeed, overexpression of PDX1.1 suppressed ammonium-dependent H<sub>2</sub>O<sub>2</sub> formation and associated root length inhibition, whereas pdx1.1 mutant lines were hypersensitive to ammonium (Fig 4 and 5, Fig S14). As in these lines total vitamin B<sub>6</sub> levels closely followed PDX1.1 transcript levels in roots, transcriptional regulation of PDX1.1-dependent PLP synthesis alone provides sufficient plasticity to counteract the adverse growth effect of ammonium. Among the three paralogs of PDX1 in Arabidopsis, only PDX1.1 and PDX1.3 show catalytic activities (Tambasco-Studart et al., 2005), while PDX1.2 encodes a pseudoenzyme that can boost vitamin B<sub>6</sub> biosynthesis via heteromerization with its paralogs in response to singlet oxygen or heat stress (Moccand et al., 2014). Although spatial and temporal expression patterns of PDX1.1 and PDX1.3 largely overlap and only disruption of both genes causes embryo lethality, PDX1.3 has been found to be more abundant and requisite for stress tolerance than PDX1.1 (Titiz et al., 2006). Nonetheless, enhanced expression of PDX1.3 at the protein level appears to require PDX1.2 (Dell’Aglio et al., 2017), whereas overexpression of PDX1.1 can be achieved with the protein alone to substantially increase vitamin B<sub>6</sub> production (Raschke et al., 2011; Fig 6). This regulatory versatility of PDX1.1 may provide an advantage when plants need to respond instantly to oxidative stresses. Second, external supply of vitamin B<sub>6</sub> to pdx1.1-3 mutant or wild-type plants completely prevented ammonium-induced inhibition of root elongation (Fig 4). Although the overall vitamin B<sub>6</sub> level in the pdx1.1-3 mutant line was only 20-30% lower than in the wild type (Fig 6A), this difference as well as a further 50-100% increase in the overexpressing lines gradually improved root growth, indicating a strong dose dependence of beneficial vitamin B<sub>6</sub> action. This dose-dependent effect went back to the abundance of the non-phosphorylated B<sub>6</sub> vitamers (Fig 6A), because only PL and PN restored root length while the primary biosynthesis product PLP remained ineffective (Fig S18). Superior functionality of non-phosphorylated vitamers as ROS scavengers is most likely determined by substituents of the pyridoxine core that modulate electron density in the ring and thus the interaction with...
singlet molecular oxygen \((^1\text{O}_2; \text{Bilski et al., 2000})\). Upon quenching of \(^1\text{O}_2\) the pyridine ring is degraded (Bilski et al., 2000), explaining the dose-dependent rather than catalytic effect of vitamin B\(_6\) as observed here (Fig 5, 6). Since in overexpression lines root concentrations of the three non-phosphorylated vitamers PM, PN and PL only, correlated closely with PDX1.1 transcript levels (Fig 6D), transcriptional upregulation of PDX1.1 is sufficient to confer a B\(_6\) vitamer-specific defense response to stress.

Third, PDX1.1-mediated biosynthesis de novo of vitamin B\(_6\) spatially and temporally coincides with ammonium-induced ROS formation in roots. Within a time frame of a few days, there was a robust temporal coincidence between root elongation rates and H\(_2\)O\(_2\) accumulation, even when vitamin B\(_6\) was provided externally or PDX1.1 expression levels were modulated (Fig 4, 5). Also at the tissue level, spatial patterns of PDX1.1 transcript levels and H\(_2\)O\(_2\) accumulation strongly overlapped in ammonium-treated roots (Fig 2, 7), supporting the notion that vitamin B\(_6\) biosynthesis is targeted to those tissues and root developmental zones that are most severely affected by ROS accumulation. In plants, biosynthesis de novo of vitamin B\(_6\) relies on PDX2 using glutamine as a substrate to produce PLP (Fig S15; Tambasco-Studart et al., 2005; Fitzpatrick et al., 2007; Boycheva et al., 2015). Glutamine is also the most abundant product of ammonium assimilation in roots (Xu et al., 2012; Liu and von Wirén, 2017), which is mediated by cytosolic glutamine synthetase. Interestingly, GLN1;2-mediated glutamine synthesis preferentially localizes in the root vasculature and GLN1;2 represents the most strongly upregulated GLN isoform in ammonium-supplied roots (Ishiyama et al., 2004; Lothier et al., 2011; Guan et al., 2016). Alternatively, ammonium may also be used directly by PDX1.1, independently of PDX2 action, as has been demonstrated in vitro (Raschle et al., 2007). In this context, elevated availability not only of glutamine but also of ammonium as one educt for vitamin B\(_6\) synthesis in those cells that suffer most from ROS production may be a factor why plants favored vitamin B\(_6\) as preferential ROS scavenger during evolutionary adaptation to elevated external ammonium levels. Given that ammonium-induced ROS formation occurs in all the cell types of the root differentiation zone and meristem (Fig 2A-C; S7B) while the enhancement of PDX1.1-mediated vitamin B\(_6\) biosynthesis under ammonium supply is confined to the vasculature (Fig 7B; S20), the question arises how root cells that do not produce vitamin B\(_6\) are protected from oxidative stress. Likely there is a radial transport
pathway bringing vitamin B\textsubscript{6} from the vasculature to the outer root cells, driven either by diffusion of non-phosphorylated B\textsubscript{6} vitamers that are considered able to permeate membranes (Stolz and Vielreicher, 2003), or by radial transport via membrane proteins, such as purine permeases (PUPs). In particular PUP1 has been shown to transport non-phosphorylated B\textsubscript{6} vitamers in Arabidopsis as well as after heterologous expression in yeast (Szydlowski et al., 2013).

Vitamin B\textsubscript{6} has proven effective in ROS detoxification in a variety of systems and conditions. In different human cell cultures, the supplementation of non-phosphorylated forms of vitamin B\textsubscript{6} alleviates superoxide-induced damage and lipid peroxidation (Jain and Lim, 2001; Kannan and Jain, 2004; Mahfouz et al., 2009). In Arabidopsis protoplasts, PN supplementation can reduce oxidative damage generated upon high illumination (Danon et al., 2005). When leaf discs were exposed directly to \textsuperscript{1}O\textsubscript{2}, lipid peroxidation in the \textit{pdx1.3} background was higher than in the wild type (Havaux et al., 2009). Since this effect was not observed with O\textsubscript{2}\textsuperscript{-} or H\textsubscript{2}O\textsubscript{2}, and since \textsuperscript{1}O\textsubscript{2} levels rise during illumination more drastically in the \textit{pdx1.3} mutant than in the wild type, vitamin B\textsubscript{6} has been proposed to act as \textsuperscript{1}O\textsubscript{2} quencher (Danon et al., 2005; Havaux et al., 2009). Moreover, \textsuperscript{1}O\textsubscript{2} can easily convert to O\textsubscript{2}\textsuperscript{-} by electron transfer (Khan and Kasha, 1994) and is further converted to H\textsubscript{2}O\textsubscript{2} by SOD, allowing reactive Fe to take in a key role in ammonium-induced ROS processing (Fig 7D). Our study could not identify the source of produced \textsuperscript{1}O\textsubscript{2} or O\textsubscript{2}\textsuperscript{-} species, as analysis of a few tested \textit{rboh} mutants did not provide sufficient evidence to rule out NADPH oxidases as a source for O\textsubscript{2}\textsuperscript{-} formation (Fig S5). Alternatively, \textsuperscript{1}O\textsubscript{2} or O\textsubscript{2}\textsuperscript{-} species may derive from electron transport processes in root plastids or mitochondria (Smirnoff and Arnaud, 2018). Irrespective of their origin, their accumulation during repression of SOD by DDC was ineffective in inhibiting root elongation (Fig S4), indicating that the conversion of O\textsubscript{2}\textsuperscript{-} to H\textsubscript{2}O\textsubscript{2} by SOD was required to inhibit root elongation (Fig 7D). Actually, these two ROS species greatly influence root development by regulating the balance between cell proliferation and cell differentiation in the root tips (Tsukagoshi, 2016). Specifically, O\textsubscript{2}\textsuperscript{-} localized in the meristematic zone maintains cell division, while H\textsubscript{2}O\textsubscript{2} prevailing in the elongation zone stimulates cell differentiation (Tsukagoshi et al., 2010). However, excessive accumulation of H\textsubscript{2}O\textsubscript{2} in the elongation zone leads to a repression of root growth reflected by inhibited cell elongation and smaller meristem size (Tsukagoshi et al.,
This may explain how H$_2$O$_2$ accumulation inhibited root elongation in ammonium-grown plants (Fig 7D).

Based on our study, we propose a working model of the processes underlying primary root growth inhibition under ammonium nutrition (Fig 7D). Ammonium uptake, which is particularly high in the elongation zone (Duan et al., 2018), provokes apoplastic acidification (Meier et al., 2020) that increases Fe solubilization and re-precipitation in inner root cells (Fig 2, 7D). It is not yet completely clear why Fe preferentially accumulates along the stele and whether the required change in Fe binding forms for xylem loading is involved, but previous studies have confirmed enhanced Fe precipitation in the pericycle and xylem (Green and Rogers, 2004; Roschztattardtz et al., 2013). There, elevated Fe availability and acidic pH favor O$_2$- dismutation and H$_2$O$_2$ formation (Smirnoff an Arnaud, 2018, Fig 7D). Considering that ammonium uptake-dependent acidification stimulates the generation of H$_2$O$_2$, H$_2$O$_2$ or a downstream product induces PDX1.1 expression at the site of Fe and ROS localization (Fig 7B, D). Predominant biosynthesis of non-phosphorylated B$_6$ vitamers (Fig 6) can subsequently quench reactive molecular oxygen (‘O$_2$/O$_2$·’) that serves as source for H$_2$O$_2$ formation (Fig S4) and thereby counteract ammonium-induced H$_2$O$_2$ formation to restore root growth. We find that this PDX1.1-mediated biosynthesis de novo of vitamin B$_6$ is not only essential for protecting roots against ammonium toxicity, as occurring in ammonium-enriched fertilizer bands in agricultural plant production, but also effective against other adverse growth conditions that involve Fe-dependent ROS formation such as P deficiency and nickel toxicity.

MATERIALS AND METHODS

Plant materials and growth conditions

*Arabidopsis thaliana* accession Col-0 and Ler served as wild type. The following mutants and transgenic lines were used: *ahkan*3-3 (Dello loio et al., 2007), *arr1*-3 (Dello loio et al., 2007), *arr12*-1 (Dello loio et al., 2007), *shy2*-31 (Dello loio et al., 2008), *shy2*-2 (Dello loio et al., 2008), *TCS:GFP* (Biela et al., 2012), *pdx1.1*-1 (Titiz et al., 2006), *pdx1.1*-3 (SALK 024245), *apo-pHusion* (Gjetting et al., 2012), *PDX1.1OE-L5* (Raschke et al., 2011), *PDX1.1OE-L8* (Raschke et al., 2011), *PDX1.1OE-L15* (Raschke et al., 2011), *pPDX1.1:GUS* (Boycheva et al., 2015), *CycB1;1::GUS* (Colón-Carmona et al., 1999). The cell cycle reporter *CycB1;1::GUS* (Col-0
CycB1;1::GUS (pdx1.1-3) and CycB1;1::GUS (PDX1.1OE-L5) lines. Complete information of T-DNA insertion lines of 29 ammonium-responsive genes used in the mutagenesis screen are listed in Data Set S1. Arabidopsis seeds were surface sterilized by 70% ethanol with 0.05% (v/v) Triton X-100, and cultured on modified half-strength Murashige and Skoog (MS) medium containing 100 µM Fe(III)-EDTA, 0.5% sucrose, 1% Duchefa Phyto agar (Duchefa Biochemie), 2.5 mM MES pH 5.7, and N sources were added to different final concentrations as described in the figure legends. Seedlings grown in Petri dishes (12 × 12 cm) were cultured vertically in a growth chamber under a 22°C/18°C and 10/14 h (light/dark) regime at the light intensity of 120 µmol photons m⁻² s⁻¹. For root phenotyping experiments, plants were pre-cultured on half-strength MS medium containing 1 mM KNO₃ for 6 d, and then transferred to treatment plates supplemented with half-strength MS medium containing 1 mM KNO₃, 1 mM NH₄Cl or 10 mM NH₄Cl respectively. 0.5 mM K₂SO₄ was added to balance K⁺ concentration in the ammonium treatment. Root phenotypes were measured 6 d after transfer. To generate P deficiency, 6 days-old plants were transferred to half-strength MS medium containing 625 µM KH₂PO₄ (+P) or 5 µM KH₂PO₄ (-P), KCl was added to -P medium to balance K⁺ concentrations. For Ni toxicity, 6 days-old plants were transferred to half-strength MS medium in the absence (-Ni) or presence (+Ni) of 75 µM NiSO₄. Root phenotypes were assessed 6 d after transfer. When supplementing H₂O₂, fresh 30% H₂O₂ (Roth) solution (9.79 mol L⁻¹) was diluted to a concentration of 1 M and supplied to the medium after autoclaving. In the root shading experiment, root-containing segments of Petri dishes were covered with aluminium foil as described in Zheng et al. (2019). For vitamin B₆ supplementation experiments, different B₆ vitamers were dissolved in water and supplied to agar medium at a final concentration of 5 µM. Key chemical information is given in Table S1.

**Root growth measurements**

During root phenotyping experiments, the position of primary roots was labeled every day after treatment in order to calculate primary root elongation rate. Root images were taken by an Epson 10000XL scanner at a resolution of 300 dpi. Primary root growth parameters were measured by WinRHIZO Pro 2007 software (Regents Instruments Canada). Differential interference contrast (DIC) microscopy images of primary root tips were taken to assess the size of primary root meristem and the
length of mature cortical cell as described in Dello Ioio et al. (2007). All the experiments were performed at least twice with similar results.

**ROS detection**

H$_2$O$_2$ in primary root tips were detected by 3,3′-Diaminobenzidine (DAB) staining as described by Thordal-Christensen et al. (1997). Briefly, primary roots were incubated in 1 mg ml$^{-1}$ DAB solution for 8 h, and then imaged by DIC microscopy. H$_2$O$_2$ is visualized as a reddish-brown coloration. A cell-permeant fluorogenic dye 2′,7′-dichlorodihydrofluorescein diacetate (H$_2$DCFDA) was also used to measure ROS activity in primary roots. Seedlings were stained for 20 min in a solution of 50 µM H$_2$DCFDA in 50 mM potassium phosphate buffers (pH 7.0). DCF fluorescence was excited at 488 nm and detected at 517-527 nm. Virtual color images were generated by a rainbow color code to indicate the fluorescence intensity of DCF in roots.

**Apoplastic and rhizosphere pH measurements**

Apoplastic pH changes were measured by the ratiometric pH reporter apo-pHusion (Gjetting et al., 2012). The fluorescence intensity ratio between pH-sensitive GFP and pH-insensitive RFP was calculated to indicate the changes of apoplastic pH, and virtual ratio images were generated by ImageJ v1.53 software. The changes of rhizosphere pH under different N supply were monitored by the pH indicator bromocresol purple (BCP) that changes color from yellow at pH 5.0 to violet at pH 7.2 (Meier et al., 2020). Six days after growth on different N sources, agar medium containing six seedlings per plate were stained overnight (18 h) with BCP solution at a final concentration of 0.1 mg ml$^{-1}$.

**Histological staining**

Promoter-driven β-glucuronidase (GUS) activity was determined by GUS staining as described previously (Li, 2011). Roots were rinsed once with staining buffer containing 50 mM NaHPO$_4$ buffer (pH 7.2), 2 mM potassium ferrocyanide, and 2 mM potassium ferricyanide, and then incubated at 37°C in staining solution containing 2 mM X-Gluc. After staining, roots were cleared by HCG solution (chloral hydrate:water:glycerol = 8:3:1) and imaged by DIC microscopy. To assess the size of the primary root meristem and the length of mature cortical cell, non-staining primary root tips were cleared by HCG solution and imaged by DIC microscopy. To stain Fe accumulation in roots, a Perls staining and DAB/H$_2$O$_2$ intensification was performed according to Roschztattardtz et al. (2009). Roots were rinsed three times with 10 mM EDTA, and then incubated for 5 min in a freshly prepared Perls staining solution.
Afterward, roots were incubated in a methanol solution containing 0.01 M NaN₃ and 0.3% (v/v) H₂O₂ for 1 h at room temperature. After washing three times with 0.1 M phosphate buffer (pH 7.4), roots were finally incubated for 5 min in an intensification solution containing 0.025% (w/v) DAB and 0.005% (v/v) H₂O₂ in 0.1 M phosphate buffer (pH 7.4). Roots were mounted with HCG solution before imaging by light microscopy.

**Microscopy analyses**

Fluorescent images were taken by laser scanning confocal microscopy Zeiss LSM 780. Root samples were stained with 10 µg ml⁻¹ propidium iodide (PI) for 5 min to visualize cell walls. GFP was excited at 488 nm and detected at 505-535 nm; RFP was excited at 561 nm and detected at 580-630 nm; PI was excited at 561 nm and detected at 600-700 nm. DCF fluorescence was excited at 488 nm and detected at 517-527 nm. The same microscope settings were kept to measure all confocal sections across samples. Fluorescence quantification of apo-pHusion and H₂DCFDA were conducted by Zeiss ZEN microscope software (version 2.6). Virtual ratio images were generated by Image J 1.53 software. DAB staining, Perls staining, GUS staining and DIC images were taken by Zeiss Axio Imager 2 system. For light microscopy of in situ-localized Fe, root cuttings of 5 mm length of 5 seedlings from each growth condition, were dissected approximately 1 cm above the root tip and subjected to aldehyde fixation, dehydration and resin embedding as described in Table S2. Semi-thin sections of 2.5 µm thickness were cut with a Leica UCT microtome (Leica Microsystems, Wetzlar, Germany), and mounted on slides in rapid mounting medium Entelan (Sigma-Aldrich, Darmstadt, Germany). Sections were recorded with a 40x lense at fixed exposure time using a Zeiss Axio Imager M2 (Carl Zeiss Microscopy GmbH, Oberkochen, Germany).

**Vitamin B₆ quantification analysis**

The abundance of all individual B₆ vitamers in roots were determined by HPLC. Vitamin B₆ quantification analyses were performed as described previously (Colinas et al., 2016) with the following changes: two separate extractions were performed with 15 volumes and 8 volumes of 50 mM ammonium acetate (pH 4.0), respectively, and a 50 µL injection volume was used for a single run per extract.

**Real-time quantitative PCR**
Total RNA was extracted from 10-20 mg frozen root samples by RNeasy plant mini kit (Qiagen) following the manufacturer’s protocol. Template cDNA was synthesized from 1 µg total RNA using SuperScript II Reverse Transcriptase (Thermo Fisher Scientific) and Oligo d(T)12-18 primer. Real-time qPCR was performed by CFX384 Touch Real-Time PCR Detection System (Bio-Rad) using iQ SYBR Green Supermix (Bio-Rad). According to the multiple internal control method (Vandesompele et al., 2002), relative transcript levels of target genes were calculated by the geNorm algorithm (https://genorm.cmgg.be), using UBQ10 (AT4G05320) and ACTIN2 (AT3G18780) as the multiple internal control genes in this study. Gene specific primers for qPCR are listed in Table S3.

**Statistical analysis**

Data were collected and analyzed by Microsoft Excel 2016. Statistical analyses were conducted by Graphpad Prism 8 (version 8.3.0). Two-tailed Student’s t-test, Dunnett’s multiple test or Tukey’s HSD test was performed to test the statistical significance, and the P-value of each statistical analysis is described in the figure legends. Graphs were plotted by Graphpad Prism 8 (version 8.3.0), and edited by Adobe Illustrator 2020 (version 24.2.1).

**Accession numbers**

Sequence data in this study can be found in the Arabidopsis Information Resource (TAIR) according to the following accession numbers: PDX1.1 (AT2G38230), PDX1.2 (AT3G16050), PDX1.3 (AT5G01410), PDX2 (AT5G60540), PDX3 (AT5G49970), SOS4 (AT5G37850), PLR1 (AT5G53580), UBQ10 (AT4G05320) and ACTIN2 (AT3G18780).

**AUTHOR CONTRIBUTIONS**

Y.L. and N. v. W. conceived the project and designed the experiments. Y.L., R.F.H.G., R.A.M., M.M., P.S. and T.B.F. performed the experiments and analyzed the data. G.K. provided the transcriptome data. Y.L. and N.v.W. wrote the manuscript with support of R.F.H.G. and T.B.F.

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FIGURE LEGENDS

Figure 1. Involvement of H$_2$O$_2$ and Fe in ammonium-dependent root growth inhibition.  
(A) Root phenotype of wild-type plants 6 d after transfer to different N supply, which contained additionally either 0.5 mM potassium iodide (KI) or 10 µM Fe(III)-EDTA (low Fe) instead of 100 µM Fe(III)-EDTA as in control plates. Horizontal marks along the root axis indicate daily positions of primary root tips. Root images were scanned 6 d after transfer. Scale bar = 1 cm. (B) - (C) Relative primary root elongation rate under 1 mM ammonium supply (B) or under 10 mM ammonium supply (C), normalized to the growth rate of plants treated with 1 mM nitrate. Symbols represent means ± SE, n = 20 plants per treatment. Asterisks denote significant differences between control and indicated treatments at each time point as * P<0.05 ** P<0.01 *** P<0.001 according to Dunnett's multiple test. (D) Cell length and meristem size of primary roots. The length of mature cortical cells and the size of the root apical meristem are indicated by yellow arrowheads. DIC images of primary root tips were taken at 6 d after transfer. Scale bar = 100 µm. (E) - (F) Quantitative readout of mature cortical cell length (E), and primary root meristem size (F). Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 16 plants. Different letters represent significant differences between treatments according to two-way ANOVA followed by Tukey's HSD test, P<0.05. (G) DAB staining of H$_2$O$_2$ in primary root tips at 6 d after treatment. The reddish-brown coloration indicates H$_2$O$_2$. Representative images from 10 plants per treatment are shown. Scale bar = 200 µm.

Figure 2. Ammonium-dependent histochemical changes in the primary root apex.  
(A) DAB staining of H$_2$O$_2$ in primary root tips at 6 d after treatment. The reddish-brown coloration indicates H$_2$O$_2$. Representative images from 10 plants per treatment are shown. Scale bar = 100 µm. (B) Staining of ROS (oxidant levels) by H$_2$DCFDA
staining. Rainbow color code (black to white) indicates DCF fluorescence intensity (low to high). Scale bars: 100 µm. (C) Quantitative readout of the fluorescence intensity of H₂DCFDA staining in different zones of the primary root. The boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values. n = 10 independent plants. Different letters represent significant differences within each individual root zone at P<0.05 according to Tukey’s HSD test. (D) Perls/DAB staining of Fe as indicated by a reddish-brown color. Scale bars: 100 µm. (E) Activity of the apoplastic pH sensor apo-pHusion. Color code (black to white) indicates fluorescence intensity ratio of eGFP/mRFP1 (low to high) and thus apoplastic pH. Scale bars: 100 µm. (F) Quantitative readout of the intensity ratio of eGFP/mRFP in different developmental zones of the primary root. Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values; n = 10 independent plants. Different letters represent significant differences within each individual root zone at P<0.05 according to Tukey’s HSD test.

After a pre-culture of 6 d, wild-type or apo-pHusion plants were transferred to the treatment medium supplied with differing N forms. Histological staining and fluorescent imaging were performed 6 d after treatment. Representative images from 10 plants per treatment are shown. Yellow arrowheads in (A) and (D), or white arrowheads in (B) and (E), indicate the boundaries of the meristematic zone, elongation zone and differentiation zone along the primary root.

Figure 3. Influence of medium pH on primary root growth and root tissue Fe. (A) Root phenotype of wild-type plants 6 d after transfer to different N supply buffered at different pH. Medium pH of 5.0, 5.7 and 6.5 was buffered by 2.5 mM MES, while pH 7.2 was buffered by 2.5 mM MOPS. Horizontal marks along the root axis indicate daily positions of primary root tips. Scale bar = 1 cm. (B) - (C) Relative primary root elongation rate under 1 mM ammonium supply (B) or under 10 mM ammonium supply (C), normalized to the growth rate of plants treated with 1 mM nitrate. Symbols represent means ± SE, n = 20 plants per treatment. Asterisks denote significant differences between control (pH 5.7) and indicated treatments at each time point as * P<0.05 ** P<0.01 *** P<0.001 according to Dunnett’s multiple test. (D) Perls/DAB staining of Fe as indicated by a reddish-brown color. Perls/DAB staining was conducted at 6 d after transfer. Representative images from 10 plants per treatment are shown. Scale bars: 100 µm.
Figure 4. Exogenous application of vitamin B₆ alleviates ammonium toxicity in roots.

(A) Root phenotype of Col-0 and pdx1.1-3 mutant plants 6 d after transfer to different N supply in the absence or presence of 5 µM vitamin B₆ (pyridoxine). Horizontal marks along the root axis indicate daily positions of primary root tips. Scale bar = 1 cm. (B) - (C) Relative primary root elongation rate under 1 mM ammonium supply (B) or under 10 mM ammonium supply (C), normalized to the growth rate of plants treated with 1 mM nitrate. Data represent means ± SE, n = 20 plants per treatment. Different letters represent significant differences at each time point at P<0.05 according to Tukey’s HSD test. (D) H₂O₂ accumulation in the primary root as visualized by DAB staining in wild-type and pdx1.1-3 mutant plants under differing N supply. DAB staining was conducted 2 d after treatment (2DAT) or 6 d after treatment (6DAT). Representative images from 10 seedlings per treatment are shown. Scale bar = 200 µm. (E) DAB staining of H₂O₂ in primary root tips 6 d after treatment. The reddish-brown coloration indicates H₂O₂. Representative images from 10 seedlings per treatment are shown. Scale bar = 200 µm.

Figure 5. Over-expression of PDX1.1 enhances ammonium tolerance in roots.

(A) Root phenotype of wild-type, pdx1.1-3 mutant and three independent PDX1.1 overexpression lines subjected to different N supply. Horizontal marks along the root axis indicate daily positions of primary root tips. After pre-culture of 6 d, plants were transferred to media containing different N forms. Root images were taken at 6 days after transfer. Scale bar = 1 cm. (B) - (C) Relative primary root elongation rate under 1 mM ammonium supply (B) or under 10 mM ammonium supply (C), normalized to the growth rate of plants treated with 1 mM nitrate. Symbols represent means ± SE, n = 20 plants per treatment. Asterisks denote significant differences between wild type and indicated lines at each time point as * P<0.05 ** P<0.01 *** P<0.001 by Dunnett’s multiple test. (D) DAB staining of H₂O₂ in primary root tips of wild type, pdx1.1-3 and PDX1.1OE-L5 at 6 d after treatment. The reddish-brown coloration indicates H₂O₂. Representative images from 10 plants per treatment are shown. Scale bar = 200 µm. (E) Cell length and meristem size of primary roots of wild type, pdx1.1-3 and PDX1.1OE-L5. The length of mature cortical cells and the size of the root apical meristem are indicated by yellow arrowheads. DIC images of primary root tips were taken 6 d after treatment. Scale bar = 100 µm. (F) - (G) Quantitative readout of mature cortical cell length (F), and primary root meristem size (G). Boxes show the
first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 16 plants. Different letters represent significant differences between lines and treatments according to two-way ANOVA followed by Tukey’s HSD test, P<0.05. (H) Influx of $^{15}$NH$_4^+$ into roots of wild type, pdx1.1-3 mutant and PDX1.1 overexpression lines. Plants were pre-cultured hydroponically for 5 weeks and then subjected to N starvation for 4 d before being transferred to 200 µM $^{15}$N-labeled NH$_4^+$ for 6 min. Bars represent means ± SD, n = 5 biological replicates. Asterisks denote significant differences between wild type and indicated lines at * P<0.05 by Dunnett’s multiple test; ns = not significant.

Figure 6. Vitamin B$_6$ profiling of pdx1.1 mutant and PDX1.1-overexpressing plants under different N regimes.
(A) Root concentrations of vitamin B$_6$, pyridoxamine 5’-phosphate (PMP), pyridoxal 5’-phosphate (PLP), pyridoxamine (PM), pyridoxine (PN) and pyridoxal (PL) in wild type (Col-0), pdx1.1-3 mutant and three PDX1.1 overexpression lines 6 d after transfer to different N supply. The amount of pyridoxine 5’-phosphate (PNP) was too low to be detected in this study. Bars represent means ± SD, n = 4 independent biological replicates. Asterisks denote significant differences between wild type and indicated lines at * P<0.05 ** P<0.01 *** P<0.001, by Dunnett’s multiple test; ND = not detected. (B) Proportion of individual B$_6$ vitamers in the roots and (C) relative transcript abundance of PDX1.1 in the roots of wild type, pdx1.1-3 and PDX1.1 overexpression lines 6 d after transfer to different N supplies. The relative transcript level of PDX1.1 was determined by quantitative real-time PCR, and normalized by using ACTIN2 and UBQ10 as internal controls. Bars represent means ± SD, n = 4 independent biological replicates. (D) Correlation between the transcript abundance of PDX1.1 and the concentration of individual B$_6$ vitamers in roots of wild type, pdx1.1-3 and PDX1.1 overexpression lines. Pearson’s correlation coefficients are shown as R values.

Figure 7. Transcriptional regulation of PDX1.1 and other vitamin B$_6$-related genes in response to ammonium.
(A) Relative transcript abundance of PDX1.1, PDX1.2, PDX1.3, PDX2, PDX3, SOS4 and PLR1 in roots of wild-type plants 6 d after transfer to different N supplies. Relative transcript levels were determined by quantitative real-time PCR and normalized by using both ACTIN2 and UBQ10 as multiple internal controls. Bars represent means ± SD, n = 3 biological replicates. Different letters represent
significant differences among means at $P<0.05$ according to Tukey’s HSD test, ns = not significant. (B) - (C) Promoter activity of the $pPDX1.1$:GUS reporter in primary roots 6 d after transfer to different N treatments in the absence or presence of 0.5 mM KI or under supply of 10 $\mu$M Fe(III)-EDTA (low Fe) (B) or in the presence of 1 mM H$_2$O$_2$ (C). Representative images from 10 plants per treatment are shown. Scale bar = 100 $\mu$m. (D) Working model for the role of ammonium-triggered proton release in Fe-dependent ROS metabolism and subsequent vitamin B$_6$ formation. Ammonium uptake provokes proton secretion and apoplastic acidification which increases Fe solubilization in the root apoplast and the rhizosphere along the elongation and differentiation zones of the root. Secreted protons i) mobilize Fe from ETDA or the apoplastic Fe pool (Fe(OH)$_3$), ii) enable superoxide dismutase (SOD)-mediated conversion of superoxide (O$_2^-$) to H$_2$O$_2$. H$_2$O$_2$ upregulates $PDX1.1$ expression to enhance accumulation of non-phosphorylated B$_6$ vitamers, in particular pyridoxine (PN) and pyridoxal (PL), which serve as antioxidants quenching the reactive molecular oxygen species $^1$O$_2$ or O$_2^{}$•. The root images on the right and left side refer to the primary root tips of DAB-stained seedlings and $pPDX1.1$:GUS reporter lines under ammonium supply, respectively.
SUPPLEMENTAL FIGURE LEGENDS

Figure S1. Ammonium suppresses root elongation in Arabidopsis.

(A) Visual appearance of seedlings subjected to different N supplies. Wild-type plants were pre-cultured on half-strength MS medium containing 1 mM NO$_3^-$ for 6 d before being transferred to 1 mM NO$_3^-$, 1 mM NH$_4^+$ or 10 mM NH$_4^+$ as sole N source. During plant growth, the positions of primary root tips were labeled every day to calculate primary root elongation rates. Images were taken 6 d after transfer. Scale bar = 1 cm.

(B) - (D), Primary root length (B), total lateral root length (C), and mean lateral root length (D) of wild-type plants 6 d after transfer to different N treatments. Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 20 independent plants. Different letters represent significant differences at $P<0.05$ according to Tukey’s HSD test.

(E) - (G), Primary root length (E), primary root elongation rate (F), and relative primary root elongation rate (G) of wild-type plants under differing N supplies. Symbols represent means ± SE, n = 20 plants per treatment. Different letters in (E) and (F) represent significant differences within each time point at $P<0.05$ according to Tukey’s HSD test. Different letters in (G) denote significant differences among time points under 1 mM or 10 mM NH$_4^+$ respectively, according to Tukey’s HSD test.

(H) Cell length and meristem size of primary roots. Primary roots were stained by propidium iodide (PI), and confocal images were taken 6 d after transfer. The length of mature cortical cells and the size of the root apical meristem are indicated by red arrowheads. Scale bar = 100 µm.

(I) - (J), Quantitative readout of primary root meristem size (I), and mature cortical cell length (J). Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 20 plants. Different letters represent significant differences at $P<0.05$ according to Tukey’s HSD test.

Figure S2. Cytokinin signaling is not involved in ammonium-dependent inhibition of root growth.

(A) Expression pattern of the cytokinin reporter TCS:GFP in primary root tips under different N supply. Images were taken 6 d after transfer. (B) Quantitative readout of TCS:GFP fluorescence intensity in primary roots under different N supply. Boxes show the first quartile, median and third quartile; the whiskers show minimum and maximum values, n = 10 plants. Asterisks denote significant differences between wild type and indicated lines at * $P<0.05$ by two-tailed Student’s $t$-test.

(C) and (G), Root phenotype of wild-type (Col-0), ahk3-3, arr1-3, arr12-1 mutants (C), and wild-type...
(Ler), shy2-31, shy2-2 mutants (G) subjected to different N supply. Roots were scanned at 6 d after transfer. Scale bar = 1 cm. The position of primary root tips at the day of transfer is labeled by black arrowheads. (D) and (H), Primary root length after transfer of Col-0, ahk3-3, arr1-3, arr12-1 (D), and Ler, shy2-31, shy2-2 (H) to different N supply. Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 10 plants in (D), n = 12 plants in (H). Different letters represent significant differences between treatments and lines according to two-way ANOVA followed by Tukey’s HSD test at P<0.05. (E) and (I), Primary root meristems of Col-0, ahk3-3, arr1-3, arr12-1 (E), and Ler, shy2-31, shy2-2 (I) under different N supply. The size of the primary root meristem is indicated by yellow arrowheads in the images. Root images were taken at 6 d after transfer to indicated N sources. Scale bar = 100 µm. (F) and (J), Quantitative readout of the primary root meristem size of Col-0, ahk3-3, arr1-3, arr12-1 (F), and Ler, shy2-31, shy2-2 (J). Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 10 plants in (F), n = 12 plants in (J). Different letters represent significant differences between treatments and lines according to two-way ANOVA followed by Tukey’s HSD test at P<0.05.

**Figure S3. H2O2 aggravates root growth inhibition in the presence of ammonium.**

(A) Root phenotype of wild-type plants 6 d after transfer to different N supply, in the absence or presence of either 1 mM H2O2 or 20 µM SHAM. Horizontal marks along the root axis indicate daily positions of primary root tips. Root images were scanned 6 d after transfer. Scale bar = 1 cm. (B) - (C) Relative primary root elongation rate under 1 mM ammonium supply (B) and under 10 mM ammonium supply (C), normalized to the growth rate of plants treated with 1 mM nitrate. Symbols represent means ± SE, n = 20 plants per treatment. Asterisks denote significant differences between control and indicated treatments at each time point as * P<0.05 ** P<0.01 *** P<0.001 according to Dunnett’s multiple test. (D) Cell length and meristem size of primary roots. The length of mature cortical cells and the size of the root apical meristem are indicated by yellow arrowheads. DIC images of primary root tips were taken at 6 d after transfer. Scale bar = 100 µm. (E) - (F) Quantitative readout of cortical cell length (E), and meristem size (F). Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 16 plants. Different letters represent significant differences between samples according
to two-way ANOVA followed by Tukey’s HSD test at $P<0.05$. (G) DAB staining of 
H$_2$O$_2$ in primary root tips at 6 d after treatment. The reddish-brown coloration 
indicates H$_2$O$_2$. Representative images from 10 plants per treatment are shown. 
Scale bar = 200 µm.

**Figure S4. Superoxide is not causal for ammonium-dependent root growth 
inhibition.**

(A) Schematic diagram of ROS metabolism and inhibitor action. Sodium 
diethyldithiocarbamate (DDC) acts as an inhibitor of superoxide dismutase (SOD), 
potassium iodide (KI) acts as H$_2$O$_2$ scavenger, and salicylhydroxamic acid (SHAM) is 
a peroxidase inhibitor (Lee et al., 2013). (B) Root phenotype of wild-type plants 6 d 
after transfer to different N supply in the absence or presence of 100 µM DDC. 
Horizontal marks along the root axis indicate daily positions of primary root tips. 
Scale bar = 1 cm. (C) - (D) Relative primary root elongation rate under 1 mM 
ammonium supply (C) and under 10 mM ammonium supply (D), normalized to the 
growth rate of plants treated with 1 mM nitrate. Symbols represent means ± SE, n = 
20 plants per treatment. Asterisks denote significant differences between control and 
DDC treatment at each time point according to two-tailed Student’s t-test at *** $P<0.001$. (E) Cell length and meristem size of primary roots. Primary roots were 
stained by propidium iodide, and confocal images were taken 6 d after transfer. The 
length of mature cortical cells and the size of the root apical meristem are indicated 
by red arrowheads. Scale bar = 100 µm. (F) - (G) Quantitative readout of cortical cell 
length (F), and meristem size (G). Boxes show the first quartile, median and third 
quartile; the whiskers show the minimum and maximum values, n = 20 plants. 
Different letters represent significant differences between samples according to two-
way ANOVA followed by Tukey’s HSD test at $P<0.05$.

**Figure S5. Screening of rboh mutants under different N supply.**

(A) Root phenotypes of wild-type plants and single or multiple rboh mutants in 
response to different N supply. After a pre-culture of 6 d, seedlings were transferred 
to medium containing either 1 mM nitrate or 1 mM ammonium as sole N source. 
Horizontal marks along the root axis indicate daily positions of primary root tips. 
Images were taken 6 d after treatment. Scale bar = 1 cm. (B) Primary root length 
after transfer. Boxes show the first quartile, median and third quartile; the whiskers 
show the minimum and maximum values, n = 18 plants. Different letters represent
significant differences between samples according to two-way ANOVA followed by
Tukey’s HSD test at $P<0.05$.

**Figure S6. Fe precipitation in the primary root under nitrate or ammonium supply.**

After a pre-culture of 6 d, wild-type plants were transferred to the medium supplied with 1 mM nitrate, 1 mM ammonium or 10 mM ammonium. Perls/DAB stained roots were embedded in Spurr resin. Root cross-sections were obtained from the differentiation zone of primary roots. Fe precipitation in primary roots is indicated by a reddish-brown colour. Scale bars: 50 µm.

**Figure S7. Dynamic changes of the Fe and ROS status in primary root tips under different N supply.**

(A) Perls/DAB staining of Fe as indicated by a reddish-brown color. After a pre-culture of 6 d, wild type (Col-0) plants were transferred to the treatment medium supplied with differing N forms. Histological staining was performed at the indicated days after transfer (DAT). Scale bars: 100 µm. (B) ROS status (oxidant levels) in primary root tips as monitored by H$_2$DCFDA staining. DCF fluorescence was detected by confocal microscopy at the indicated time points. Rainbow color code (black to white) indicates DCF fluorescence intensity (low to high). Scale bars: 100 µm. (C) Quantitative readout of the fluorescence intensity of H$_2$DCFDA staining. The boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 12 plants. Different letters represent significant differences between samples at $P<0.05$ according to Tukey’s HSD test. Orange arrowheads in (A), or white arrowheads in (B), indicate the boundaries of the meristematic zone, elongation zone and differentiation zone along the primary root.

**Figure S8. Fe aggravates ammonium-repressed cell division in roots.**

(A) GUS activity in primary root tips of a CycB1;1::GUS reporter line. After a pre-culture of 6 d, CycB1;1::GUS reporter plants were transferred to treatment medium supplied with different N forms. The treatment medium contained 100 µM Fe(III)-EDTA (control) or 10 µM Fe(III)-EDTA (low Fe). Histological staining was performed 6 d after transfer. Scale bar = 100 µm. (B) Quantitative readout of the dividing cell number in primary root tips of CycB1;1::GUS reporter line. Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 12 plants. Different letters represent significant differences between samples according to two-way ANOVA followed by Tukey’s HSD test at $P<0.05$. 
**Figure S9. Ammonium-dependent histochemical changes in the lateral root apex.**

(A) Cell length and meristem size of lateral root. After a pre-culture of 6 d, wild type (Col-0) seedlings were transferred to the treatment medium supplied with different N forms. Six days after transfer, lateral roots were stained by propidium iodide and imaged by confocal microscopy. The length of mature cortical cells and the size of the meristem are indicated by red arrowheads. Scale bar = 100 µm. (B) - (C), Quantitative readout of lateral root meristem size (B), and lateral root cortical cell length (C). Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 20 plants. Different letters represent significant differences at P<0.05 according to Tukey’s HSD test. (D) DAB staining of H$_2$O$_2$ in lateral root tips at 6 d after treatment. The reddish-brown coloration indicates H$_2$O$_2$. Representative images from 10 plants per treatment are shown. Scale bar = 100 µm. (E) Detection of ROS (oxidant levels) by H$_2$DCFDA staining. Rainbow color code (black to white) indicates DCF fluorescence intensity (low to high). Scale bars: 100 µm. (F) Quantitative readout of the fluorescence intensity of H$_2$DCFDA staining in lateral roots. The boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values. n = 12 plants. Different letters represent significant differences at P<0.05 according to Tukey’s HSD test. (G) Perls/DAB staining of Fe in lateral root tips as indicated by a reddish-brown color. Representative images from 10 plants per treatment are shown. Scale bars: 100 µm. (H) Activity of the apoplastic pH sensor apo-pHusion in lateral roots. Color code (black to white) indicates fluorescence intensity ratio of eGFP/mRFP1 (low to high) and thus apoplastic pH. Scale bars: 100 µm. (I) Quantitative readout of the intensity ratio of eGFP/mRFP in lateral root. Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values; n = 12 plants. Different letters represent significant differences at P<0.05 according to Tukey’s HSD test. Yellow arrowheads in (D) and (G), or white arrowheads in (E) and (H), indicate the boundaries of the meristematic and elongation zone along the lateral root.

**Figure S10. Changes in rhizosphere pH after supply of different N forms.**

After a pre-culture of 6 d, wild-type (Col-0) seedlings were transferred to the treatment medium supplied with different N forms. Medium pH of 5.0, 5.7 and 6.5 was buffered by 2.5 mM MES, while pH 7.2 was buffered by 2.5 mM MOPS. Changes of medium pH through plants are monitored by the pH indicator
bromocresol purple (BCP) that changes color from yellow at pH 5.0 to violet at pH 7.2, while the left side of the plates without plants serves as a blank.

**Figure S11. Influence of buffer strength on root growth and root Fe status.**

(A) Root phenotype of wild-type (Col-0) plants 6 d after transfer to treatment medium containing MES buffer at different strength. Horizontal marks along the root axis indicate daily positions of primary root tips. Scale bar = 1 cm. (B) - (C) Relative primary root elongation rate under supply of 1 mM ammonium (B) or 10 mM ammonium (C), normalized to the growth rate of plants treated with 1 mM nitrate. Symbols represent means ± SE, n = 20 plants per treatment. Asterisks denote significant differences between 2.5 mM MES and indicated treatments at each time point as * P<0.05 ** P<0.01 *** P<0.001 according to Dunnett’s multiple test. (D) Perls/DAB staining of Fe in primary root tips as indicated by a reddish-brown color. Perls/DAB staining was conducted at 6 d after transfer. Representative images from 10 plants per treatment are shown. Scale bars: 100 µm.

**Figure S12. Light is not essential for ammonium-dependent inhibition of primary root elongation.**

(A) Experimental setup of the root shading treatment. Petri dish-grown plants were exposed to white light in the control treatment (light), while the whole root zone was covered by aluminium foil in the shading treatment (shading), as described in Zheng et al. (2019). (B) Appearance of plants grown on the indicated nitrogen supply. After a pre-culture of 6 d, wild-type (Col-0) plants were transferred to the treatment plates and either exposed to light or shaded. Images were taken 6 d after transfer. Scale bar = 1 cm. (C) Primary root length after transfer of wild-type plants to light or shading. Boxes show the first quartile, median and third quartile; the whiskers show minimum and maximum values; n = 20 plants. Different letters represent significant differences at P<0.05 according to Tukey’s HSD test.

**Figure S13. The formation of hydroxyl radicals is not critical for ammonium-inhibited root elongation.**

(A) Root phenotype of wild-type (Col-0) plants 6 d after transfer to different N supplies in the absence (control) or presence of 0.5 mM or 1.0 mM thiourea. Horizontal marks along the root axis indicate daily positions of primary root tips. Scale bar = 1 cm. (B) - (C) Relative primary root elongation rate under supply of 1 mM ammonium (B) or 10 mM ammonium (C), normalized to the growth rate of plants treated with 1 mM nitrate. Symbols represent means ± SE, n = 20 plants per treatment. Asterisks denote significant differences between control and thiourea.
treatments at each time point as * $P<0.05$ ** $P<0.01$ *** $P<0.001$ according to Dunnett’s multiple test.

**Figure S14.** *pdx1.1* mutants are hypersensitive to ammonium.

(A) Phenotypic screening of T-DNA insertion lines of ammonium-responsive genes. Primary root length under ammonium supply was taken as a read-out for ammonium sensitivity. Candidate lines were pre-cultured on half-strength MS medium containing 1 mM nitrate for 6 d before being transferred to medium containing either 1 mM nitrate or 1 mM ammonium as sole N source. Horizontal marks along the root axis indicate daily positions of primary root tips. Images were taken 6 d after transfer. Scale bar = 1 cm.

(B) Relative primary root length after transfer to 1 mM ammonium, normalized to the root length of plants treated with 1 mM nitrate. Asterisks denote significant differences between wild-type (Col-0) and indicated lines as *** $P<0.001$ according to by Dunnett's multiple test; ns = not significant.

(C) Schematic diagram of the T-DNA insertions in the *PDX1.1* gene. Positioning of the T-DNA insertion in *pdx1.1*-3 (*SALK* _024245) and the transposon insertion in *pdx1.1*-1 (*SM_3_22664*) (Titiz *et al.*, 2006). (D) Root phenotype of wild-type (Col-0), *pdx1.1*-1 and *pdx1.1*-3 plants subjected to different N supplies. Horizontal marks along the root axis indicate daily positions of primary root tips. Images were taken 6 d after transfer. Scale bar = 1 cm.

(E) Primary root length of wild-type (Col-0) and *pdx1.1* mutant plants after transfer to different N forms. Boxes show the first quartile, median and third quartile; the whiskers show minimum and maximum values; $n = 18$ plants. Different letters represent significant differences between samples according to two-way ANOVA followed by Tukey’s HSD test, $P<0.05$.

**Figure S15. Vitamin B₆ metabolism in Arabidopsis.**

Vitamin B₆ comprises six different vitamers. PLP, as a phosphorylated B₆ vitamer is a bioactive form participating as coenzyme in numerous enzymatic reactions. Non-phosphorylated forms of vitamin B₆ are PN, PL, and PM. In plants, PLP is produced by biosynthesis *de novo* (blue) or by the salvage pathway (brown) via the conversion among different B₆ vitamers. This schematic is modified from Colinas *et al.* (2016). Gln, glutamine; Glu, glutamate; R5P, ribose 5-phosphate; G3P, glyceraldehyde 3-phosphate; PNP, pyridoxine 5’-phosphate; PLP, pyridoxal 5’-phosphate; PMP, pyridoxamine 5’-phosphate; PN, pyridoxine; PL, pyridoxal; PM, pyridoxamine; P-ase, phosphatase.
Figure S16. Exogenous application of vitamin B₆ rescues cell length and meristem size under ammonium supply.
(A) - (B) Cell length and meristem size of primary roots of wild-type (Col-0) (A) or pdx1.1-3 mutant (B). Plants grown under different N supply and in the absence (- VB₆) or presence (+ VB₆) of 5 µM pyridoxine. DIC images of primary roots were taken 6 d after transfer. The length of mature cortical cells and the size of the root apical meristem are indicated by yellow arrowheads. Scale bar = 100 µm. (C) - (D) Quantitative readout of cortical cell length (C), and meristem size (D). Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 16 plants. Different letters represent significant differences between samples according to three-way ANOVA followed by Tukey’s HSD test, P<0.05.

Figure S17. Exogenous application of vitamin B₆ restores cell division under ammonium supply.
(A) GUS activity of the CycB1;1::GUS reporter in primary roots of wild-type (Col-0) and pdx1.1-3 mutant plants. After a pre-culture of 6 d, CycB1;1::GUS reporter lines were transferred to different N forms in the absence (- VB₆) or presence (+ VB₆) of 5 µM pyridoxine. Histological staining were performed 6 d after treatment. Scale bar = 100 µm. (B) Quantitative readout of the number of dividing cells. Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 12 plants. Different letters represent significant differences among treatments according to three-way ANOVA followed by Tukey’s HSD test at P<0.05.

Figure S18. Non-phosphorylated forms of vitamin B₆ are critical for ammonium detoxification.
(A) Root phenotype of wild-type (Col-0) plants 6 d after transfer to different N supply in the absence or presence of 5 µM vitamin B₆ provided either as pyridoxal 5’-phosphate (PLP), pyridoxal (PL), or pyridoxine (PN). Horizontal marks along the root axis indicate daily positions of primary root tips. Seedlings subjected to N supplies with exogenous supplementation of different Bs vitamers. Scale bar = 1 cm. (B) - (C) Relative primary root elongation rate under 1 mM ammonium supply (B) or under 10 mM ammonium supply (C), normalized to the growth rate of plants treated with 1 mM nitrate. Symbols represent means ± SE, n = 20 plants per treatment. Asterisks denote significant differences between control and indicated treatments at each time point as * P<0.05 ** P<0.01 *** P<0.001 according to Dunnett’s multiple test. (D) DAB staining of H₂O₂ in primary root tips 6 d after treatment. The reddish-brown coloration
indicates H$_2$O$_2$. Representative images from 10 seedlings per treatment are shown. Scale bar = 200 µm.

**Figure S19. Over-expression of PDX1.1 rescues cell division under ammonium supply.**

(A) GUS activity of the CycB1;1::GUS reporter in primary root tips of wild-type (Col-0), pdx1.1-3 and PDX1.1OE-L5 plants. After a pre-culture of 6 d, CycB1;1::GUS reporter lines were transferred to the treatment medium supplied with different N forms. Histological stainings were performed 6 d after transfer. Scale bar = 100 µm. (B) Quantitative readout of the number of dividing cells in primary root tips of CycB1;1::GUS reporter lines. Boxes show the first quartile, median and third quartile; the whiskers show the minimum and maximum values, n = 12 plants. Different letters represent significant differences between treatments and lines according to two-way ANOVA followed by Tukey’s HSD test at P<0.05.

**Figure S20. Influence of medium pH on the expression of PDX1.1 upon ammonium supply.**

Promoter activities of the pPDX1.1:GUS reporter in primary roots subjected to different N forms and buffered at different pH were monitored by GUS staining 6 d after transfer. Medium pH of 5.0, 5.7 and 6.5 was buffered by 2.5 mM MES, while pH 7.2 was buffered by 2.5 mM MOPS. Representative images from 10 plants per treatment are shown. Scale bar = 100 µm.

**Figure S21. Over-expression of PDX1.1 improves tolerance to P deficiency and Ni toxicity.**

(A) - (B) Heat map displaying the relative expression level of vitamin B$_6$-related genes (highlighted in boxes) in response to phosphorus deficiency (A) or nickel toxicity (B). Data were retrieved from published microarray experiments (Bhosale et al., 2018; Lesková et al., 2020). Selected genes previously shown to be up- or down-regulated by phosphorus deficiency or nickel toxicity were included as controls. -P, 5 µM phosphate; +P, 625 µM phosphate; ++Ni, 100 µM nickel; control, no nickel added to growth medium. (C) Appearance of plants grown on different phosphate (P) supplies. After a pre-culture of 6 d, the seedlings of Col-0, pdx1.1-3, PDX1.1OE-L5, PDX1.1OE-L15 and PDX1.1OE-L8 were transferred to treatment medium containing 625 µM P (+P) or 5 µM P (-P). Images were taken 6 d after transfer. Scale bar = 1 cm. (D) Primary root length after transfer. Root phenotyping of indicated lines was conducted 6 d after transfer. Boxes show the first quartile, median and third quartile;
the whiskers show minimum and maximum values; n = 20 plants. Different letters represent significant differences between treatments and lines according to two-way ANOVA followed by Tukey's HSD test at $P<0.05$. (E) Appearance of plants subjected to nickel (Ni) toxicity. After a pre-culture of 6 d, the seedlings of wild-type (Col-0), pdx1.1-3, PDX1.1OE-L5, PDX1.1OE-L15 and PDX1.1OE-L8 were transferred to half-strength MS medium in the absence (- Ni) or presence (+ Ni) of 75 µM Ni. Images were taken 6 d after transfer. Scale bar = 1 cm. (F) Primary root length after transfer. Root phenotypes were recorded 6 d after transfer. Boxes show the first quartile, median and third quartile; the whiskers show minimum and maximum values; n = 20 plants. Different letters represent significant differences between samples according to two-way ANOVA followed by Tukey's HSD test at $P<0.05$.

**Table S1. Chemical Information.**

**Table S2. Protocol for fixation, dehydration and embedding of root tissue.**

**Table S3. Primers used in this study.**

**Data Set S1. Information on ammonium-responsive genes used for the mutant screening in this study.**
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