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Natural variation at *XND1* impacts root hydraulics and trade-off for stress responses in *Arabidopsis*

Ning Tang ¹, Zaigham Shahzad^{1,4}, Fabien Lonjon², Olivier Loudet ³, Fabienne Vaillau² & Christophe Maurel ¹

Soil water uptake by roots is a key component of plant performance and adaptation to adverse environments. Here, we use a genome-wide association analysis to identify the XYLEM NAC DOMAIN 1 (*XND1*) transcription factor as a negative regulator of *Arabidopsis* root hydraulic conductivity (L_p). The distinct functionalities of a series of natural *XND1* variants and a single nucleotide polymorphism that determines *XND1* translation efficiency demonstrate the significance of *XND1* natural variation at species-wide level. Phenotyping of *xnd1* mutants and natural *XND1* variants show that *XND1* modulates L_p through action on xylem formation and potential indirect effects on aquaporin function and that it diminishes drought stress tolerance. *XND1* also mediates the inhibition of xylem formation by the bacterial elicitor flagellin and counteracts plant infection by the root pathogen *Ralstonia solanacearum*. Thus, genetic variation at *XND1*, and xylem differentiation contribute to resolving the major trade-off between abiotic and biotic stress resistance in *Arabidopsis*.

¹BPMP, CNRS, INRA, Montpellier SupAgro, Université de Montpellier, 34060 Montpellier, France. ²LIPM, Université de Toulouse, INRA, CNRS, 31326 Castanet-Tolosan, France. ³Institut Jean-Pierre Bourgin, INRA, AgroParisTech, CNRS, Université Paris-Saclay, 78000 Versailles, France. ⁴Present address: Institute of Molecular, Cell and Systems Biology, College of Medical, Veterinary and Life Sciences, University of Glasgow, Bower Building, Glasgow G12 8QQ, UK. Correspondence and requests for materials should be addressed to C.M. (email: christophe.maurel@cnrs.fr)

The growth performance and survival of terrestrial plants, whether in favorable or adverse environments, crucially depend on a proper uptake and management of water. Most plant species forage the soil for water through continuous growth and development of roots into a ramified architecture. The intrinsic water transport properties of root tissues (i.e., their root hydraulic conductivity, L_{p_r}) are also important for efficient uptake and transfer of water towards the shoots. L_{p_r} shows a high environmental plasticity, with typical regulations depending on the availability of water, mineral nutrients or oxygen in the soil¹. The same stimuli act on root growth and development, thereby altering root system architecture (RSA)². Overall, growth and water transport properties of roots, which combine into the so-called root hydraulic architecture, determine the plant's capacity to capture soil water under changing or heterogeneous soil conditions. Plants also display remarkable intraspecific genetic variations in RSA and hydraulics^{3–7}, with possible impacts on abiotic stress responses. Thus, the ultimate question is to understand how combined genetic and physiological adjustments of RSA and root water permeability contribute to plant adaptation to specific habitats or climatic scenarios.

Root water transport per se relies on several fundamental processes, often presented as sequential. Radial water flow, from the soil to the vasculature in the stele, is mediated through cell walls (apoplastic path) or from cell-to-cell. The latter path combines transcellular (across cell membranes and aquaporins) and symplastic (across plasmodesmata) transport. Water is then axially transported to the aerial parts through xylem vessels. The contexts in which root xylem can be hydraulically limiting are still debated⁸. Based on Poiseuille's law of laminar flow, it was calculated that under water sufficient conditions this tissue is supposedly not limiting with respect to root structures mediating radial transport⁹. This may not be true in root tips, whereby vessels are not fully differentiated. Xylem cavitation under drought can also dramatically reduce plant hydraulic conductance and confer high plant vulnerability¹⁰. Conversely, drought impacts xylem differentiation further supporting a crucial role of vascular transport in these conditions^{5,8}. However, this view may not apply to certain species or under mild water stress since intraspecific variation of xylem size in major crops such as rice was not associated to any growth advantage, especially under water deficit¹¹. Thus, the links that xylem vessel differentiation establishes between root growth and development, hydraulics, and stress responses are not fully established⁸. In contrast, detailed studies have revealed how positioning of xylem axes contributes to early vascular pattern formation and how subsequent differentiation of xylem tracheary elements occurs through cell clearance by programmed cell death and deposition of lignin in secondary cell walls¹². These developmental processes are controlled by regulatory networks involving NAC (NAM, ATAF1,2, and CUC2)¹³ and MYB (myeloblastosis)-type transcription factors¹⁴. The former proteins have played a key evolutionary role in water-conducting structures from moss to vascular plants¹⁵.

While root hydraulics is prone to fine biophysical analyses, genetic dissection of this trait has been somewhat lagging due in part to technical difficulties in defining proper hydraulic phenotypes. Reverse genetic analyses of candidate genes have uncovered the limiting role of aquaporins¹ or endodermal barriers^{16,17}. In contrast, other components which determine root anatomy or environmental signaling and which can potentially interfere with root hydraulic properties have been poorly explored¹⁸. Quantitative genetics approaches based on intraspecific variations of root hydraulics^{3,4} could help uncover such molecular components. In line with these ideas, quantitative trait locus (QTL) analysis of L_{p_r} in a biparental recombinant population

(Bur-0 × Col-0) of *Arabidopsis* led to the cloning of *hydraulic conductivity of root 1*, a Raf-like MAPKKK gene that acts as a negative regulator of L_{p_r} ¹⁹. Here, we perform a genome-wide association analysis as another approach to identify genes controlling root hydraulics in *Arabidopsis*. We identify *XYLEM NAC DOMAIN 1 (XND1)* as a key negative regulator of *Arabidopsis* root hydraulics at the species-wide level. Our study also reveals how genetic variation at *XND1* may contribute to the trade-off between abiotic stress tolerance and biotic defense in *Arabidopsis*.

Results

A GWA study uncovers two novel genes controlling L_{p_r} . A set of 143 *Arabidopsis* accessions from the RegMap panel²⁰ was phenotyped for root hydraulics (Supplementary Data 1). A fourfold variation of L_{p_r} was observed among accessions (Supplementary Figure 1), with a coefficient of variation of 0.245 and a broad-sense heritability $h^2 = 0.36$. Conditional genome wide association (GWA) mapping using 250k single nucleotide polymorphisms (SNP) data²⁰, and an accelerated mixed-model algorithm method with four markers as cofactors²¹ revealed two SNPs that were significantly associated with L_{p_r} variation (Bonferroni multiple testing correction at $\alpha = 0.05$) and contributed to 18.3% and 7.3% of the genetic variance, respectively (Fig. 1a). One was located on chromosome (Chr) 1 (position 13,612,169), while the other was on Chr 5 (position 25,787,448). Considering 20-kb genomic regions surrounding these two association SNPs (Fig. 1b, c), we identified eight candidate genes for L_{p_r} variation. None of these genes showed SNPs that were in strong linkage disequilibrium (LD) with the corresponding GWA studies (GWAS) peak SNP (Supplementary Figure 2). They were therefore further evaluated using a total of 15 Col-0-derived T-DNA insertion lines. As for the Chr1 region, two allelic insertion lines for At1g36240 showed, with respect to Col-0, an increase in L_{p_r} by 11 or 17%, while mutant lines for three neighboring genes did not show any significant L_{p_r} phenotype (Supplementary Table 1). When considering the four genes located in the Chr 5 region, we observed significantly altered L_{p_r} specifically for three T-DNA insertion lines of At5g64530 (Supplementary Table 1; Fig. 1d). The data reveal the power of GWA mapping for identifying genetic determinants of specific traits like root hydraulics. At1g36240 encodes a putative ribosomal protein which will be investigated in another work. At5g64530 encodes Xylem NAC Domain 1 (XND1), a NAC transcription factors that antagonizes xylem differentiation, by negatively regulating secondary cell wall synthesis and programmed cell death²². Its putative function in axial water transport led us to examine in closer details its contribution to root hydraulics, which has not been described previously.

XND1 negatively regulates root hydraulics. Quantitative reverse transcription polymerase chain reaction (qRT-PCR) analyses revealed contrasting *XND1* mRNA abundance in the three *xnd1* allelic mutants (Fig. 1d). *xnd1-3* and *xnd1-4*, which both exhibit a T-DNA insertion in the *XND1* promoter region appeared as knock-down and activation lines, respectively. Consistent with a T-DNA insertion in the second intron, *xnd1-5* can rather be considered as a knock-out allele. When considering the root dry weight (DW), primary and total root length, or lateral root density in plants grown in hydroponics, all three *xnd1* genotypes had a root architecture similar to that of Col-0 (Supplementary Figure 3). In contrast, and by reference to Col-0, L_{p_r} was increased in hypofunctional mutants (*xnd1-3*, *xnd1-5*) by up to $29.7 \pm 7.6\%$ and decreased by $18.6 \pm 4.7\%$ in the *xnd1-4* over-expression mutant. These results indicate that XND1 acts as a true negative regulator of root hydraulics. Consistent with this,

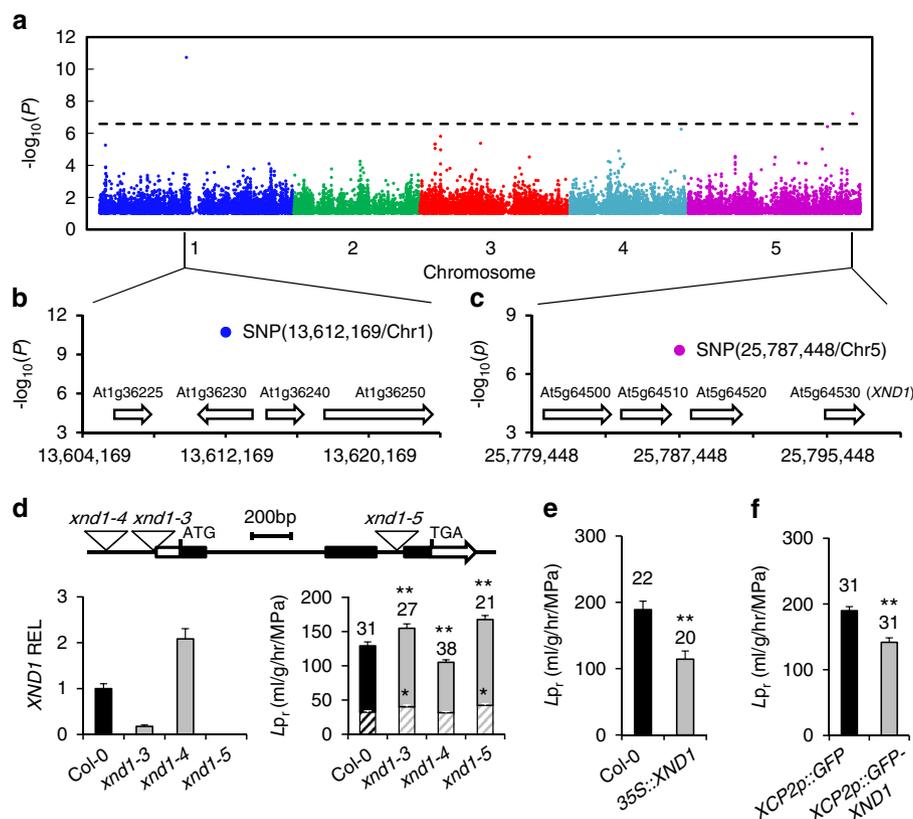


Fig. 1 Identification of *XND1* as a genetic determinant of root hydraulic conductivity. **a** Manhattan plot of GWA for root hydraulic conductivity (L_p_r) data based on a conditioned accelerated mixed-model. The chromosomes are depicted in different colors and the x-axis represents the chromosome number and position. A Bonferroni corrected significance threshold at $\alpha = 0.05$ is indicated by the horizontal dashed line. **b, c** Gene models of 20-kb genomic regions surrounding the two most significantly associated SNPs. The blue and purple dots represent these SNPs, and numbers on x-axes indicate chromosomal positions in base pairs. **d** Molecular and phenotypic characterization of three *xnd1* T-DNA insertion lines. The upper diagram shows a schematic representation of the *XND1* genomic region with positions of the three T-DNA insertions. The bar graph on the left shows *XND1* transcript abundance relative to Col-0 (relative expression level, REL) in the *xnd1* lines (means \pm SE, based on two biological repeats). The bar graph on the right shows L_p_r phenotypes before (whole bars) or after (hatched bars) root treatment for 30 min with the aquaporin blocker NaN_3 . **e, f** L_p_r phenotype of Col-0 transgenic lines with *XND1* constructs under control of a *CaMV35S* (**e**) or a xylem specific *XCP2Pro* (*XCP2p*) (**f**) promoter. In **f**, plants expressing GFP alone under the control of *XCP2Pro* were used as negative controls. In all panels, L_p_r data (means \pm SE) were based on the indicated number of plants in two to three independent experiments. Asterisks indicate significant differences with respect to control lines (Student's *t* test; * $P < 0.05$, ** $P < 0.01$)

overexpression of *XND1* under the control of a *CaMV35S* promoter (Supplementary Figure 4A, B) resulted in a dramatic ($-39.5 \pm 7.7\%$) reduction in L_p_r (Fig. 1e). *XND1* is expressed in root xylem, preferentially in association with differentiating tracheary elements^{22,23}. Transgenic expression of a GFP-*XND1* fusion protein under the control of xylem specific promoter *XCP2Pro* (Supplementary Figure 4C, D) reduced L_p_r by $25.3 \pm 4.3\%$ (Fig. 1f), implying that *XND1* acts in vascular tissues to exert hydraulic effects.

Allelic diversity at *XND1* validates GWA analyses. The L_p_r associated SNP identified at position 25,787,448/Chr 5 during GWA analysis is located 7.9 kb apart from the *XND1* coding region, whereas SNPs that are closer did not show any significant association. This hints to possible allelic heterogeneity and multiple mutations in *XND1* that would contribute to L_p_r variation and are linked to the associated SNP. Newly released genomic sequences²⁴ were used to evaluate in closer details natural variation at *XND1* among 112 accessions, of which 85 belong to the initially investigated panel while the remaining corresponds to accessions for which phenotypic data were generated later (Supplementary Data 2). Considering a genomic region from 2 kb

upstream to 0.3 kb downstream of the *XND1* coding region, we used a generalized linear model to test the association with L_p_r at 27 polymorphic sites (minor allele frequency (MAF) > 0.05), including SNPs and INDELs (Fig. 2a, b). One single SNP located at position 25,795,349 in the 5'-UTR of *XND1* surpassed the significance threshold and was thereafter named SNP_{UTR}. Several SNPs also pointed to possible associations in the promoter region, consistent with putative allelic heterogeneity. Further, we used the eight polymorphisms showing the lowest *P* values ($P < 0.075$) in this association analysis to define six haplogroups, each containing 7–38 accessions (Fig. 2c; Supplementary Data 2). These haplogroups fall into four phenotypic classes with significant differences in mean L_p_r values (ANOVA; $P < 0.05$) (Fig. 2c) suggesting that genetic diversity at *XND1* truly contributes to shaping L_p_r variation at the species level. To establish this point further, we selected four accessions (Bur-0, Ty-0, Col-0, and Fei-0) as representative members of haplogroups with contrasting mean L_p_r values (Fig. 2c). Corresponding genomic fragments encompassing *XND1* were introduced into *xnd1-5* and tested for their capacity to complement the mutant L_p_r phenotype (Fig. 2d). In these experiments, we used a set of three to five independent transgenic lines per haplotype, which provide mean *XND1* expression levels similar to that in native Col-0 (Supplementary

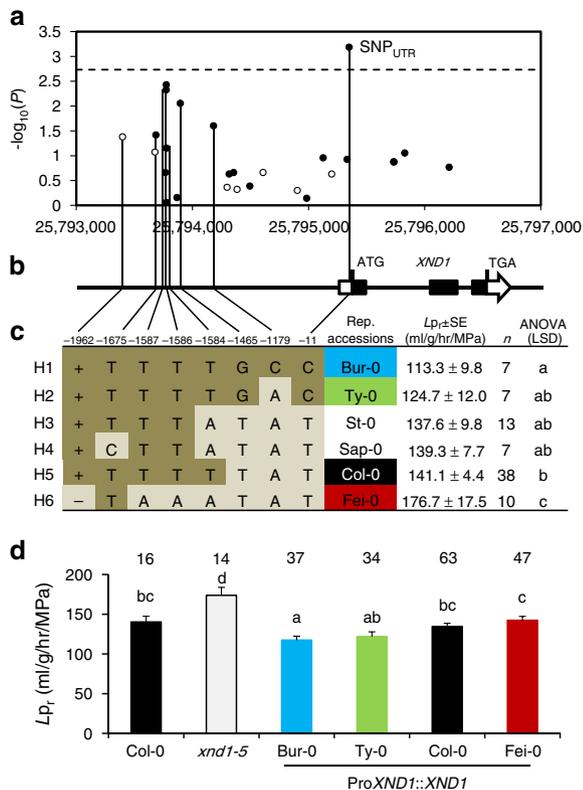


Fig. 2 A *XND1*-based association analysis allows refining the natural allelic variations of *XND1*. **a** The association with L_{p_r} of 27 polymorphic sites (MAF > 0.05) in the indicated *XND1* genomic region was investigated in a set of 112 accessions. The x-axis shows the nucleotide position of each variant, with empty and filled circles indicating INDELs and SNPs, respectively. The y-axis shows the $-\log_{10}(P)$ for the association tests, with the significance threshold at $\alpha = 0.05$ indicated with a dashed line. **b** The eight polymorphisms selected for further analysis are projected onto a schematic representation of *XND1* gene structure. For position -1962 , + and $-$ represent an insertion and deletion, respectively. The boxes represent exons, with solid and empty boxes showing translated and untranslated regions, respectively. The SNP at Chr 5-P25,795,349 that surpassed the significance threshold in **a** is located in the 5'-UTR of *XND1* and indicated as SNP_{UTR} . **c** The eight selected nucleotide polymorphisms (with their distance from translation start site shown on the top) define six haplogroups (H1–H6). Representative accessions and mean $L_{p_r} \pm SE$ within each haplogroup (n , accessions number) are shown. One-way ANOVA (Fisher's LSD, $P < 0.05$) was used to test the significance of the L_{p_r} data. **d** Transgenic complementation of *xnd1-5* with different allelic forms of *XND1*. The L_{p_r} of Col-0, *xnd1-5*, and *xnd1-5* plants expressing *XND1* genomic fragments from Bur-0, Ty-0, Col-0, or Fei-0, was tested using 3–5 independent transgenic lines per *XND1* allele (see Supplementary Figure 5). Mean values $\pm SE$ ($n = 14$ –63 plants) are shown with sample size indicated on the top. One-way ANOVA (Fisher's LSD, $P < 0.05$) was used to test the significance of the data

Figure 5A,B). All four complemented series exhibited lower L_{p_r} than *xnd1-5*, indicating that each of *XND1* allelic forms was functional in a Col-0 background (Fig. 2d; Supplementary Figure 5C; Supplementary Data 3). In addition, transformation with the Bur-0 or Ty-0 alleles yielded lower L_{p_r} values than when using the Col-0 or Fei-0 allele, or the Fei-0 allele, respectively. Thus, with respect to Col-0 and Fei-0, the first two accessions harbor somewhat hyperfunctional alleles.

To further investigate the significance of variation at *XND1*, we revisited previous phenotyping data from a Bur-0 \times Col-0 cross¹⁹. Although there was no sign of a purely additive QTL at the

bottom of Chr 5 in this cross¹⁹, we detected an epistatic QTL controlling L_{p_r} variation (Supplementary Figure 6), possibly including *XND1* effect. The phenotypic consequence of segregation at *XND1* region is conditioned by the allele present at a locus on Chr 2, suggesting that phenotypic expression of *XND1* variation may depend on the genetic background. We note that, consistent with the effects of introducing a Bur-0 *XND1* allele in Col-0 (Fig. 2d), the RILs fixed for Bur-0 at *XND1* region and Col-0 at the Chr2 locus exhibited a reduced L_{p_r} (Supplementary Figure 6).

The overall genetic data validate the identification of *XND1* by GWA mapping, and support the idea that allelic differences at *XND1* contribute to the natural variation of L_{p_r} in *Arabidopsis*.

A SNP in *XND1* can account for natural variation of L_{p_r} .

Among the 112 accessions analyzed above, the SNP_{UTR} , with C vs. T variation, identifies two haplogroups with significantly different L_{p_r} (SNP_{UTR-C} , $L_{p_r} = 114.8 \pm 7.6$, $n = 16$; SNP_{UTR-T} , $L_{p_r} = 145.5 \pm 3.3$, $n = 96$; $P < 0.01$). Bur-0 and Col-0 belong to the first and second haplogroups, respectively (Fig. 2c). We addressed the functional significance of SNP_{UTR} by transgenic complementation of *xnd1-5* mutant with Bur-0 or Col-0 allelic forms of *XND1*, either native or with a converting point mutation at SNP_{UTR} (Fig. 3a, d). While mutation of the Col-0 allele did not influence L_{p_r} (Fig. 3b, c; Supplementary Figure 7A, C; Supplementary Data 4), the conversely mutated Bur-0 allele conferred a higher L_{p_r} than its native counterpart in *xnd1-5* complemented lines (Fig. 3e, f; Supplementary Figure 7B, D; Supplementary Data 4). These data establish that SNP_{UTR} contributes to natural variation of *XND1* function. Since L_{p_r} phenotypic differences induced by these point mutations were in either case smaller than when comparing the two wild-type alleles (Fig. 2d; Fig. 3c, f), we hypothesize that other polymorphisms act, independently or in interaction with SNP_{UTR} , to induce the full allelic effects. Effects of these polymorphisms would be more pronounced in the Col-0 allele than in the Bur-0 allele.

Next, we addressed the general molecular mechanisms that can account for natural variation of *XND1* function. Knowing that the *XND1* allelic variation pointed by GWA and validated by transgenic complementation does not reside in the *XND1* coding sequence, we investigated natural variation in *XND1* expression and its potential functional significance. The abundance of *XND1* transcripts was measured in the roots of 49 randomly selected accessions representative of the six previously defined H1–H6 haplogroups. No correlation of L_{p_r} with *XND1* transcript abundance was observed when considering all individual accessions (Fig. 4a), or their grouping in the H1–H6 (Fig. 4b) or SNP_{UTR-C} vs. SNP_{UTR-T} (Fig. 4c) haplogroups. Thus, the SNP_{UTR} and other putative SNP(s) causing L_{p_r} variations have marginal effects, if any, on mRNA abundance at this scale. To investigate putative effects of SNP_{UTR} on *XND1* protein translation, a *XND1*-GFP fusion was placed under the control of a *CaMV35S* promoter and a *XND1* 5'-UTR, in its SNP_{UTR-C} or SNP_{UTR-T} form, and expressed in transgenic *xnd1-5* (Supplementary Figure 8). Based on 9–10 lines per genotype with overall similar transcript abundance, we used GFP fluorescence as a reporter of *XND1*-GFP accumulation (Supplementary Figure 9A–D). For each line, a relative translation efficiency was deduced from the ratio of *XND1*-GFP protein and mRNA abundances (Supplementary Figure 9E). Combined data (Fig. 4d) revealed that relative translation efficiency of the SNP_{UTR-C} form was 63% higher than that of its SNP_{UTR-T} counterpart. The positive effect of SNP_{UTR-C} on *XND1* translation and accumulation is consistent with lower L_{p_r} in the corresponding haplogroup.

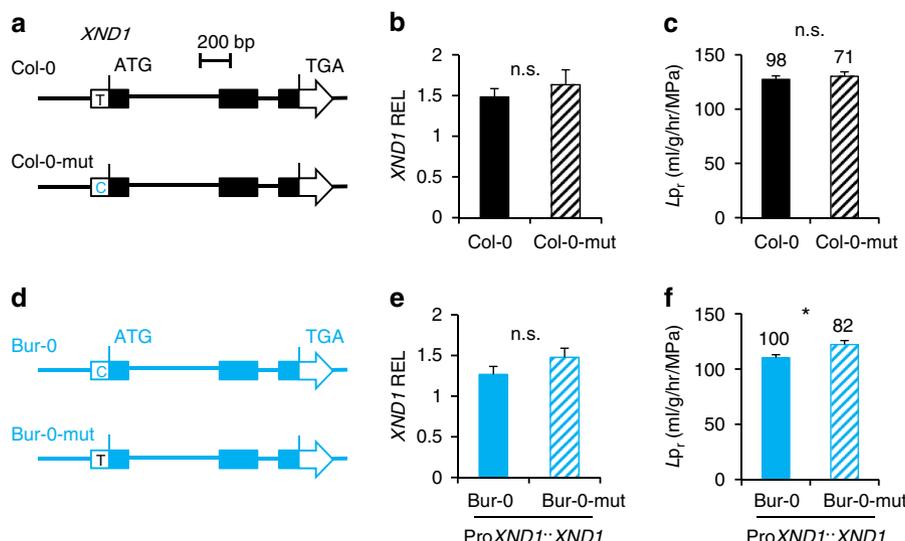


Fig. 3 SNP_{UTR} of *XND1* contributes to *L_p* variation. Transgenic complementation of *xnd1-5* with Col-0 or Bur-0 allelic forms of *XND1*, either wild-type or with a site-directed mutation at SNP_{UTR}. **a, d** Schematic representation of investigated *XND1* allelic forms. Col-0 and Bur-0 indicate the wild-type forms while Col-0-mut and Bur-0-mut indicate the corresponding point mutated forms. **b, e** Transcript abundance of *XND1* in *xnd1-5* homozygous transgenic lines expressing the Col-0, Col-0-mut, Bur-0, or Bur-0-mut forms of *XND1*. Three to five independent transgenic lines (two biological replicates for each) were studied for each form. Mean values ± SE (*n* = 18–30) were normalized to transcript abundance in Col-0. **c, f** *L_p* of the transgenic lines described above. Three to five independent transgenic lines were phenotyped for each form. Mean values ± SE are shown with total number of plants indicated on the top. Student’s *t* test (*P* < 0.05) was used to assess the statistical significance of the data (n.s. = not significant). *XND1* expression and *L_p* data in individual transgenic lines are shown in Supplementary Figure 7

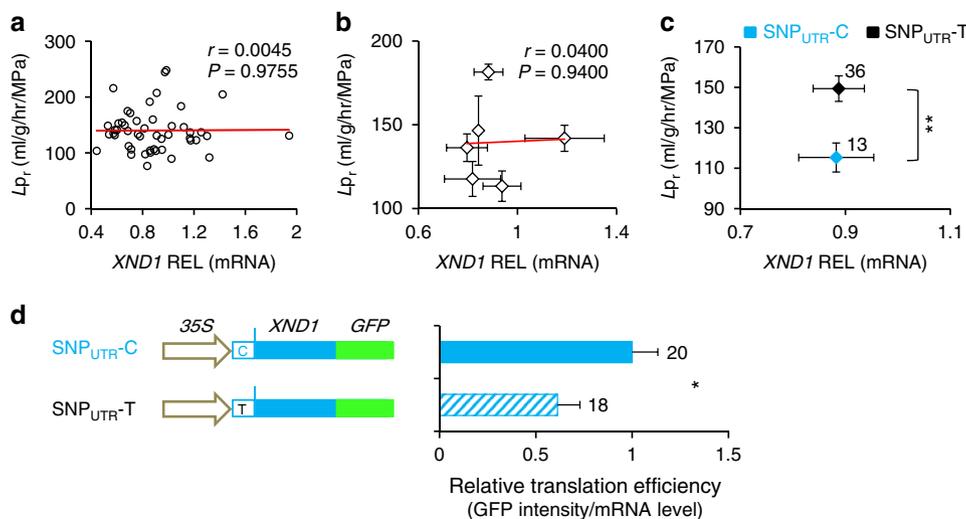


Fig. 4 *XND1*-dependent natural variation of *L_p* is contributed by effects of SNP_{UTR} on *XND1* translation. **a–c** *XND1* transcript abundance was measured in the roots of 49 accessions using 6–10 plants and two independent experiments per accession. Data were analyzed considering each individual accession (**a**) or their grouping in the H1–H6 (**b**) or SNP_{UTR}-C vs. SNP_{UTR}-T haplogroups (**c**). Pearson correlation coefficient (*r*) and *P* value (*P*) between *XND1* transcript level and *L_p* are shown in **a** and **b**. Error bars in **b** and **c** indicate SE. Accession number per haplogroup was *n* = 3–18 in **b** and as indicated in **c** (*P* < 0.01 in Student’s *t* test). **d** A *XND1*–GFP fusion construct was placed downstream of a *CaMV35S* promoter and a *XND1* 5’-UTR, with either C or T at SNP_{UTR}. Note that this SNP is the only variation between Bur-0 and Col-0 *XND1* alleles present in the construct sequence. Transcript abundance and fluorescence intensity of *XND1*–GFP was quantified in the roots of 9–10 transgenic lines per genotype, with two biological replicates. Relative translation efficiency of *XND1*–GFP was calculated in each individual line from the ratio of fluorescence to mRNA abundance. Mean values ± SE based on total number of lines and repeats are indicated on the right and were normalized to the data for SNP_{UTR}-C. Student’s *t* test (*P* < 0.05) was used to assess the statistical significance of the data

***XND1* affects both xylem formation and aquaporin activity.** To understand further the causes of *XND1*-mediated variation of *L_p*, we used sodium azide (NaN₃), a common blocker of plant plasma membrane aquaporins^{3,19}. NaN₃ dramatically reduced *L_p* in Col-0 and three T-DNA insertion *xnd1* genotypes, yielding a residual *L_p*, which was higher in loss-of-function alleles (*xnd1-3*, *xnd1-5*) than in wild type (Col-0) or gain-of-function (*xnd1-4*)

plants (Fig. 1d). NaN₃-resistant *L_p* accounts for aquaporin-independent water transport pathways, whether radial (apoplasm) or axial (xylem vessels). Here, facilitation of NaN₃-resistant pathways in *xnd1-3* and *xnd1-5* is consistent with *XND1* acting as a negative regulator of xylem differentiation. To assess this point, we probed xylem morphology along the root axis by sampling tissues in 2-cm-long consecutive segments from the

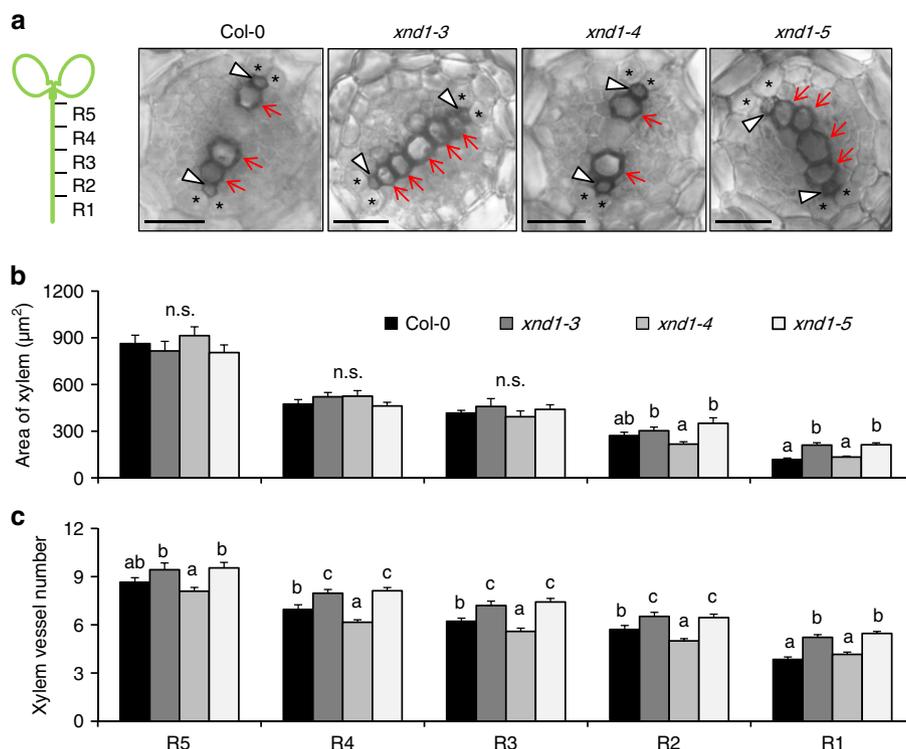


Fig. 5 XND1 negatively regulates root xylem formation. **a** The figure shows schematic representation of 2-cm-long segments (R1–R5) used for characterization of xylem anatomy in the indicated genotypes and representative cross-sections within the R2 segment. Arrowheads and arrows indicate protoxylem and metaxylem, respectively. Asterisks indicate pericycle cells adjacent to xylem poles. Scale bars represent 20 μm . **b, c** Xylem area (**b**) and vessel number (**c**) (mean values \pm SE) were measured in the indicated root segment and genotype, in 20–61 sections from 10–15 plants and 2–3 independent experiments. One-way ANOVA (Fisher's LSD, $P < 0.05$) was used to test the significance of the data (n.s. = not significant)

root tip (Fig. 5). By comparison to Col-0, formation of the xylem (especially metaxylem) was advanced in *xnd1-3* and *xnd1-5*, while delayed in *xnd1-4* (Fig. 5a). Specifically, overall xylem area was enhanced in *xnd1-3* and *xnd1-5* in the most apical segment (R1) (Fig. 5b). Xylem vessel number was consistently increased in several segments (R1–R4) of loss-of-function *xnd1* alleles whereas it was reduced in the R2–R4 segments of *xnd1-4* (Fig. 5c).

We also observed that the absolute difference in L_{p_r} between Col-0 and *xnd1* mutants was much higher in the absence than in the presence of NaN_3 (Fig. 1d), suggesting that, with respect to Col-0, loss and gain-of-function *xnd1* mutants show higher and lower plasma membrane aquaporin activity, respectively. Because XND1 serves as a transcription factor, we investigated whether it may alter, either directly or indirectly, the abundance of the corresponding aquaporin transcripts in roots. Thus, we considered all the 13 *Plasma membrane Intrinsic Protein (PIP)* genes and investigated the three T-DNA insertion *xnd1* lines and XND1 overexpressors described above (*35S::XND1*, *XCP2p::GFP-XND1*), taking Col-0 and *XCP2p::GFP* plants as controls. None of the *PIP* genes showed differences in expression that would correlate to L_{p_r} (Supplementary Figure 10), suggesting that effects of XND1 on aquaporin function are not mediated through *PIP* transcript abundance. The overall data indicate that XND1 acts as a negative regulator of L_{p_r} by repressing both xylem development and aquaporin activity. The latter effects are not yet understood and may well be indirect.

XND1 negatively acts on plant drought stress tolerance. By determining the efficiency of water supply to the plant's aerial parts, root hydraulics can play a key role in maintaining the overall plant water status, especially in conditions of strong water demand or water deprivation. To investigate the general role of

XND1 in plant water relations, we first compared the wild type (Col-0) and three *xnd1* genotypes grown under various water regimes. Under sufficient water supply, all genotypes showed comparable growth performance, although *xnd1-4* showed a slight deficit in shoot water content (Supplementary Figure 11A, B). When seedlings were grown for three weeks in these favorable conditions and thereafter subjected to a long-term (24 days) water deprivation, all plant genotypes showed signs of severe stress and growth retardation, which, however, were the most pronounced in *xnd1-4* (Fig. 6a). Plant resilience to these extreme conditions was investigated at 5 days after rewatering. With respect to Col-0 plants, loss-of-function *xnd1* mutants showed a higher shoot fresh weight (FW) and a tendency to higher shoot DW (Fig. 6b and Supplementary Figure 11C). The gain-of-function *xnd1* plants (*xnd1-4*) showed a mirror phenotype with reduced shoot growth (lower FW and DW) and reduced shoot water content (Fig. 6b and Supplementary Figure 11C). The overall data indicate that XND1 negatively acts on plant tolerance to water deficit. To address the significance of XND1 natural variation in plant drought tolerance, we investigated *xnd1-5* lines complemented with the Bur-0, Col-0, or Bur-0 SNP_{UTR}-mutated allelic forms of XND1. Under water sufficient condition, all complemented lines showed comparable growth performance with respect to Col-0 and *xnd1-5*, with a slight reduction in shoot water content in the Bur-0 allele complemented lines (Supplementary Figure 12). After successive drought and recovery, all complemented lines showed lower shoot FW than *xnd1-5* indicating the functionality of the three XND1 allelic forms in these conditions (Fig. 6c and Supplementary Figure 13). Interestingly, complementation with the Bur-0 allele resulted in lower shoot FW than with the Col-0 or Bur-0 SNP_{UTR}-mutated allele (Fig. 6c and Supplementary Figure 13). In addition, the shoot DW and

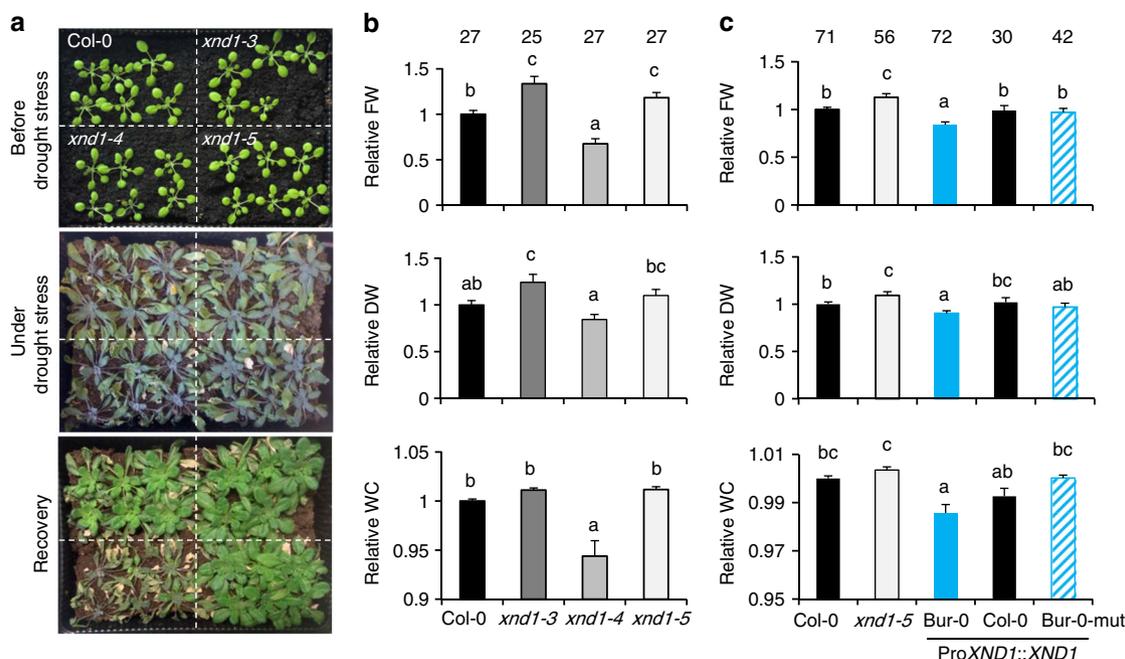


Fig. 6 XND1 negatively regulates plant drought stress tolerance. **a** Shoot phenotypes of Col-0 and *xnd1* plants at 22 days postgermination (upper panel), after water deprivation for 24 additional days (middle panel), and at 5 days after rewatering (lower panel). **b** Relative fresh weight (FW), dry weight (DW) and water content (WC) in shoots of indicated genotypes at 5 days after rewatering (see **a**). All data were normalized to values measured in Col-0 plants from the same pot. The figure shows mean values \pm SE from the indicated number of plants and four biological replicates. One-way ANOVA (Fisher's LSD, $P < 0.05$) was used to test the significance of the data. **c** Same parameters as in **b** but for Col-0, *xnd1-5*, and *xnd1-5* plants complemented with the Bur-0, Col-0, or Bur-0-mut *XND1* alleles. Three to five independent transgenic lines were phenotyped for each allele. Mean values \pm SE from the indicated number of plants are shown. One-way ANOVA (Fisher's LSD, $P < 0.05$) was used to test the significance of the data

water content of the Bur-0 allele complemented lines was lower than in lines containing the Col-0 or Bur-0 SNP_{UTR}-mutated allele, respectively (Fig. 6c and Supplementary Figure 13). These differences establish that natural variation at *XND1*, at the SNP_{UTR} in particular, confers variations in integrated plant responses to drought.

XND1 contributes to protection against a vascular pathogen.

Plant vasculature serves as an infection route for pathogens and its development is targeted by multiple biotic factors. Along these lines, abundance of *XND1* transcripts in *Arabidopsis* is enhanced up to fivefold by bacterial infiltration (e.g., *Pseudomonas syringae* pv. *phaseolicola*) and exposure to pathogen-associated molecular patterns (e.g., flg22) (see *Arabidopsis* eFP Browser database at http://bar.utoronto.ca/efp_arabidopsis/cgi-bin/efpWeb.cgi).

These data led us to examine in closer details the effects of flg22 on xylem formation. A treatment with 0.25 μ M flg22 for 4 days reduced the overall xylem area in R2 and R3 segments of Col-0, but not in *xnd1-3* or *xnd1-5* (Fig. 7a). In R2 root segments of Col-0, xylem vessel number was reduced by flg22 whereas it was unresponsive to the peptide in corresponding root segments of the three *xnd1* insertion mutants (Fig. 7b). Thus, XND1 mediates in part adjustments of vasculature formation under biotic constraints. The significance of these effects was further tested by exposing roots of various *xnd1* genotypes to a soil borne pathogenic bacterium (*Ralstonia solanacearum*) and scoring plant infection symptoms for up to 9 days (Fig. 7c, Supplementary Figure 14A). Based on plant survival rate, the *xnd1-3* knock-down line was indistinguishable from Col-0. In contrast, *xnd1-5* and *xnd1-4* showed, with respect to Col-0, a higher and lower susceptibility to the bacterial strain, respectively. Measurements of bacterial growth in rosettes confirmed this ranking, indicating that at 3 days

postinoculation, *xnd1-5* and *xnd1-4* showed *in planta* bacterial multiplication that was higher ($P = 0.007$) and lower ($P = 0.004$), respectively, than in Col-0 (Fig. 7d). At a later stage (4 days postinoculation), all genotypes exhibited similar bacterial contents (Supplementary Figure 14B). Transformation of *xnd1-5* with the Bur-0, Col-0, or Bur-0 SNP_{UTR}-mutated allelic forms of *XND1* abolished the higher susceptibility of *xnd1-5* to *R. solanacearum*, with bacterial wilt phenotypes of the complemented lines similar to that of Col-0 (Supplementary Figure 14C). The overall data indicate that XND1 limits bacterial proliferation in the plant and subsequent bacterial wilt appearance, supporting the idea that reduced xylem formation contributes to plant protection against bacterial infection.

Discussion

Recent work from our group¹⁹ has shown that, although technically challenging and under strong environmental and developmental dependency, root hydraulics (L_{p_r}) is amenable to quantitative genetics analyses. With respect to the recombinant population mapping approach used in our previous work, GWA studies could be more straightforward for identifying candidate genes quantitatively acting on a trait of interest. In any case, it is described that these two approaches are certainly complementary to reveal the genetic architecture of a trait²⁵. In the present study, a one shot phenotyping of 143 accessions allowed identification of two novel genes, *At1g36240* and *At5g64530*, which proved to act as negative regulators of L_{p_r} .

While additional work is required to assess the role of *At1g36240*, several complementary lines of evidence indicate that *At5g64530* (*XND1*) corresponds to the gene spotted by the GWA signal on Chr 5. First, loss- and gain-of-function of *XND1* in Col-0 insertion mutants resulted in opposite L_{p_r} phenotypes.

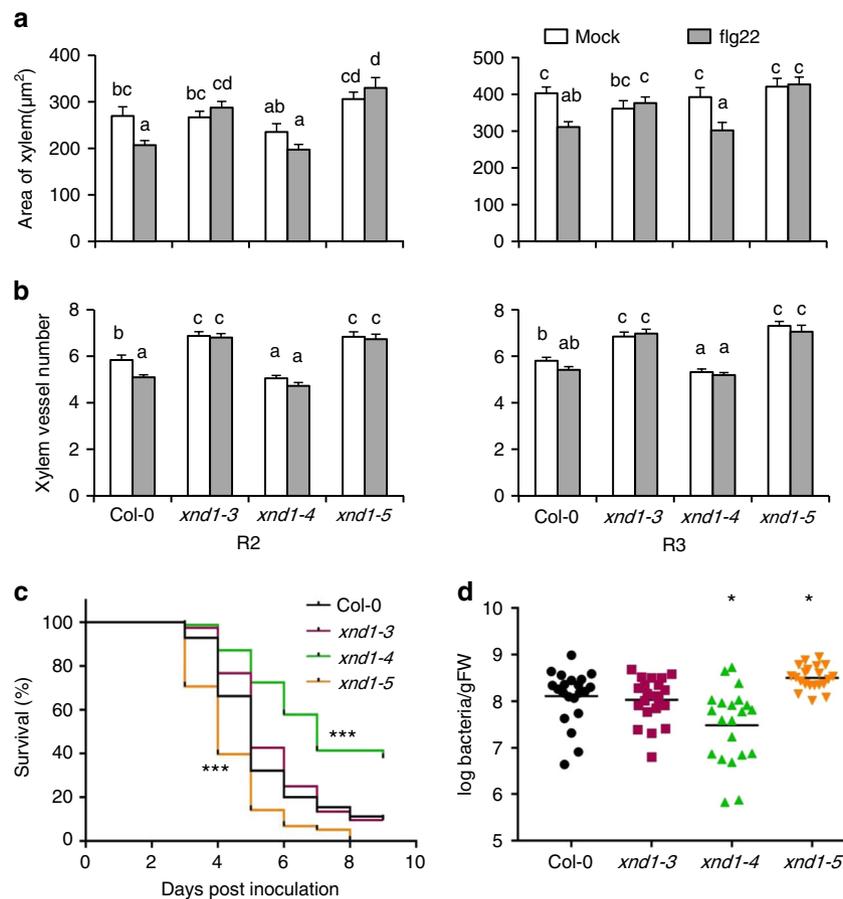


Fig. 7 XND1 mediates the inhibition of xylem formation by flg22 and resistance to *R. solanacearum*. **a, b** Plants of the indicated genotype were grown for 4 days in the absence (mock) or in the presence of 0.25 µM flg22 (flg22). R2 and R3 root segments (see Fig. 5a) were cross-sectioned and xylem area (**a**) and xylem vessel number (**b**) were measured. Data are means ± SE from 18–50 sections in 6–10 plants and two independent experiments. One-way ANOVA (Fisher’s LSD, $P < 0.05$) was used to test the significance of the data. **c** The indicated genotypes were inoculated with *R. solanacearum* through root dipping and plant survival (in %) was scored at the indicated days postinoculation. Cumulated data from three independent biological experiments, each with three technical repeats comprising at least 24–32 plants. Stars indicate significant differences from Col-0 based on a Gehan-Breslow-Wilcoxon test ($P < 0.0001$). The Col-0 and *xnd1-3* curves were not significantly different ($P = 0.1304$). **d** *in planta* growth measurement of *R. solanacearum* at 3 days after inoculation of the indicated genotypes. Each dot represents a replicate of two plants and the black line indicates the mean of all the replicates. Cumulated data from four independent biological repeats. Asterisks indicate a significant difference with Col-0 (Mann-Whitney test; $P < 0.05$)

Although very encouraging to confirm our candidate gene’s association²⁶, this is not sufficient per se. Thus, we also showed that four natural *XND1* allelic forms have distinct abilities to complement the enhanced Lp_r phenotype of *xnd1-5* knock-out plants. Remarkably, the corresponding haplogroups, each containing 7–38 natural accessions, showed consistent quantitative differences in Lp_r further supporting the idea that an allelic series at *XND1* significantly modulates root hydraulics at the species-wide level. Additionally, an epistatic Lp_r QTL corresponding to the *XND1* region was mapped in a Bur-0 × Col-0 RIL population, with effects on Lp_r consistent with those of corresponding *XND1* alleles. Because of confounding genetic relatedness or allelic heterogeneities, the statistics of GWAS detection can lead to complex patterns where causal polymorphisms in the gene of interest are away from and do not show LD with the most significantly associated SNPs²⁷. The present work may typically represent such case since the Chr 5 GWAS peak SNP (SNP_{peak}) was not in strong LD with any single SNP of neighboring genes in a 48 kb region (Supplementary Figure 2). Yet, a gene-based association study with a different accession set and an extended polymorphism data, pointed to a SNP present in the 5’-UTR of

XND1 (SNP_{UTR}) as two allelic forms, and located at a 7901 bp distance from the Chr 5 GWAS peak SNP. The fact that SNP_{UTR} and SNP_{peak} do not appear in strong LD ($r^2 = 0.23$, Supplementary Figure 2), could be explained in part by a bias due to the relatively low minor allele frequency (14%) at SNP_{UTR}. Again, we used transgenic complementation to assess the functional significance of this variation: the contribution of this SNP_{UTR} to Lp_r was established by site-directed allelic conversion of a Bur-0 allele and functional expression in a Col-0 background. Thus, the hyperfunctionality of the Bur-0 vs. Col-0 allele is due in part to a T-to-C substitution at this position. The failure of a symmetrical mutation in the Col-0 allele to modify Lp_r indicates that SNP_{UTR} functionally interacts with other intragenic components to yield the *XND1*-dependent variation of Lp_r . In particular, SNPs in the 3’UTR were not considered and could contribute to natural phenotypic variation. In summary, the overall work provides a thorough dissection of natural variations at *XND1* and demonstrates their contribution to root hydraulics.

Causal polymorphisms can be located in coding regions, in regulatory non-coding regions (this work) or can correspond to genomic structural variations²⁸. Their identification usually

provides highly relevant insights into the function and regulation of the gene or gene product of interest. In the case of *XND1*, two protein interaction motifs present in the C-terminal domain are essential for its function as they allow binding to the general cell cycle and differentiation regulator RETINOBLASTOMA-RELATED²⁹. Yet, these motifs were highly conserved in the accession panel of this study and seem not to underlie *XND1*-based natural variation of *L_p*.

Here, we found that *L_p* natural variation was not correlated to *XND1* transcript abundance when natural accessions were assembled in functionally discriminating haplogroups (Fig. 4b, c; Supplementary Figure 15). Knowing that the 5'-UTR of plant genes can harbor regulatory elements that affect RNA structure, RNA-protein interactions and/or ribosome recruitment^{30,31}, we investigated whether allelic variation at SNP_{UTR} interferes with *XND1* translation efficiency. Because *XND1* is a transcription factor that is lowly abundant and primarily confined to the nucleus of xylem cells, we monitored fluorescence of a *XND1*-GFP construct in a specific root tip region as it was the most suitable to proper quantification. Although a definitive analysis such as ribosome profiling remains to be done, our results based on ratio of protein and mRNA abundances in cells ectopically expressing *XND1* suggest that SNP_{UTR} potentially affects translational efficiency. As a result, we propose that variation at SNP_{UTR} leads to distinct *XND1* protein abundance, xylem differentiation and *L_p*. We also stress that, as indicated by the local association study, SNP_{UTR} is likely not the only causal SNP, and other variants, for instance specific to haplotype H6 (Fig. 2c), may also be functional.

The identification of *XND1* in our GWA study was strikingly in line with the notion that NAC transcription factors control water-conducting structures from primitive terrestrial plants to most recently evolved vascular plants^{15,22,32}. Further, *XND1* is an angiosperm-specific NAC that contributes to specific vessel differentiation features in these plants²⁹. Consistent with earlier studies²², we validated that *XND1* negatively acts on xylem differentiation and, based on Hagen-Poiseuille law³³, we propose that its developmental function results in an axial hydraulic limitation in root tips. A genetic link between xylem function and root hydraulics has previously been pointed by Lefebvre et al.¹⁸, based on the dramatic drop in *L_p* displayed by *esk1* loss-of-function mutants. With respect to the general collapse of xylem vessels and pleiotropic stress phenotypes shown by these mutants, *xnd1* genotypes showed milder alterations of xylem tissues, specifically in roots tips. Thus, a slightly altered abundance and overall size of the xylem vessels seem to be sufficient to perturb *L_p*. A more precise analysis indicated, however, that *XND1* may also act on aquaporin function to reduce *L_p*. This could be due to indirect effects on aquaporin expression and/or function due to locally altered vascular tissue differentiation. While these conclusions rely on analysis of *xnd1* transgenic lines, the precise mechanisms that underlie effects of natural *XND1* variants on *L_p* remain to be explored.

Beyond root hydraulics, our study underscores a general role of xylem in plant response to environmental stresses. In particular, we showed that *XND1* negatively acts on drought tolerance. Based on the phenotype of *XND1* overexpressors, we propose that reduced xylem differentiation and water transport in root tips hampers water extraction in deep, newly colonized soil strata. A reduced leaf water content was also observed in well-watered conditions, suggesting a general hydraulic limitation in these plants. These findings contribute to a longstanding debate on the role of xylem in drought tolerance. Genetic selection programs in wheat and rice have targeted xylem morphology using root thickness as a proxy^{7,34}. However, the significance of these variations with respect to drought adaptation has been controversial

(discussed by Lynch et al.⁵). In wheat, reduced xylem diameter was linked to better growth in dry environments whereas, in rice, genotypic variation in xylem vessel diameter was not linked to the plant's capacity to regulate leaf water status under drought¹¹. The genetic variants of *XND1* investigated in the present work provide more precise insights into these questions, at least in the case of *Arabidopsis*. We also realize that, besides efficiency for conducting the transpiration stream, xylem anatomy contributes to cavitation resistance and thereby to tolerance to extreme drought events. However, this critical trait, which shows high phenotypic and genotypic plasticity in trees³⁵, could not be addressed in our work since no natural cavitation has ever been described in *Arabidopsis*.

Xylem differentiation is sensitive to many more environmental factors than water availability^{36,37}. For instance, a link between PAMP-triggered immunity and xylem differentiation pathways involving plant glycogen synthase kinase 3 proteins is emerging^{38,39}. In the present work, we investigated the significance of *XND1* induction by flg22, and found this gene to mediate the inhibiting effects of the elicitor on vascular formation. A general role for *XND1* in plant defense was further investigated using *R. solanacearum* a prototypal root vascular pathogen that causes bacterial wilt in many plant species⁴⁰. As for drought assays, series of hypo- and hyperfunctional, artificial and natural alleles were used to address *XND1* function. Consistent with its induction upon infection, enhanced expression of *XND1* in *xnd1-4* led to plants more tolerant to *R. solanacearum* than Col-0, whereas a loss-of-function mutation (*xnd1-5*) caused a mirror phenotype that could be complemented using either Col-0 or Bur-0 alleles. Several modes of actions may explain the role of *XND1* in the present pathosystem. In agreement with *in planta* bacterial growth measurements, we propose that restriction or delayed differentiation of the vasculature mediated by *XND1* may antagonize microbe propagation, and/or the establishment of an aqueous living space for pathogens⁴¹, thereby contributing to plant defense. Alternatively, and as shown for other NACs⁴², *XND1*-mediated transcriptional reprogramming could enhance plant immunity.

Because of their general role in differentiation of water-conducting tissues, NACs have played a central role during plant evolution, and contributed to their adaptation to aerial environments¹⁵. Our study shows that natural selection is still at work in higher plants, and refines the role of a crucial NAC member in diverse vascular functions. By negatively acting on the differentiation of xylem vessels in root tips, *XND1* sensitizes the plant to extreme drought events while directly or indirectly providing a partial protection against systemic pathogen attack. Thus, *XND1* mediates a trade-off between plant responses to biotic and abiotic stresses. While environmental stresses can occur independently, sequentially or in combination, this work exclusively reports on effects of individual stresses. Since the role of *XND1* could conceivably be more complex when stresses occur in concert than when applied individually, additional dimensions of the role of this gene in plant stress responses remain to be explored. Nevertheless, our work indicates that natural variation at *XND1* may contribute to the extensive geographic distribution of species such as *Arabidopsis*⁴³. Interestingly, the minor allelic form (14% of the 112 accessions) carrying SNP_{UTR}-C was associated with hyperfunctional alleles, lower *L_p*, and lower drought tolerance. This indicates that the selective balance of drought over pathogen attacks may have been predominant to maintain SNP_{UTR}-T as the major allele. While our study emphasizes the adaptive significance of *XND1* variations in natural plant populations, it will also be crucial to investigate the genetic variation of this gene and of other NAC homologs in trees³² and herbaceous crops, for

applied purposes such as wood quality, productivity under drought conditions or disease control.

Methods

Plant materials. Seeds for natural accessions of *Arabidopsis thaliana* (Supplementary Data 1) or T-DNA insertion lines (Supplementary Table 1) were obtained from the Nottingham Arabidopsis Stock Centre (NASC) except for seeds of FLAG_153A06 which came from the Versailles Arabidopsis Stock Center. When required, T-DNA insertions were confirmed by PCR using primers listed in Supplementary Table 2. *XND1* overexpression lines under control of a *CaMV35S* or xylem specific *XCP2* promoter⁴⁴ were as described previously^{22,29}.

Hydroponic growth conditions. Seeds were surface sterilized and sown on petri plates (12 × 12 cm, Gosselin, BP124-04) with 0.5 × Murashige and Skoog (MS) solid medium [2.2 g L⁻¹ MS (Sigma, M5519), 10 g L⁻¹ sucrose (Euromedex, 200–301 B), 0.5 g L⁻¹ MES (Euromedex, EU0033), and 0.7% agar (Sigma, A4675), pH 5.7 adjusted using KOH]. Following stratification for 2 days in the dark at 4 °C, plates were incubated vertically for 9 days in a growth chamber at 20 °C and 70% relative humidity, with cycles of 16 h of light (250 μmol photons m⁻² s⁻¹) and 8 h of night. Plants were then transferred in hydroponic culture under the same growth conditions. The hydroponic medium (0.75 mM MgSO₄, 0.5 mM KH₂PO₄, 1.25 mM KNO₃, 1.5 mM Ca(NO₃)₂, 50 μM Fe-EDTA, 100 μM Na₂SiO₃, 50 μM H₃BO₃, 12 μM MnSO₄, 1 μM ZnSO₄, 0.7 μM CuSO₄, 0.24 μM Na₂MoO₄) was replaced weekly.

Root hydraulic conductivity (*L_p*) measurements. Measurements were performed essentially as described³ on 21- to 23-day-old plants cultured in hydroponics. The intact root system of a freshly de-topped plant was inserted into a pressure chamber filled with hydroponic solution (pH 6.5 adjusted using KOH), and sealed with a silicon dental paste (PRESIDENT Light Body, Coltene, Switzerland). The root system was subjected to a pressure pretreatment at 350 kPa for 10 min to attain equilibration of sap flow exuded from the hypocotyl section, and to successive treatments at 320, 160, and 240 kPa, each for 2 min. High-accuracy flow meters (Bronkhorst, France) in combination with a LabVIEW-derived application were used to record the rate of pressure (*P*)-induced sap flow (*J_s*). DW_r was determined after desiccation at 80 °C for at least 24 h. *L_p* (in ml g⁻¹ h⁻¹ MPa⁻¹) was calculated as: $L_p = J_s / (DW_r \cdot P)$. In azide (NaN₃) inhibition experiments, *J_s* was measured at continuous 320 kPa and its percentage inhibition was measured at 30 min following the addition of 1 mM NaN₃.

GWA, candidate-based association and haplotype analysis. For GWA analysis, *L_p* was measured on 143 natural accessions, each with 6 individual plants randomized over a measuring period of four weeks (Supplementary Data 1). For the calculation of broad-sense heritability, average intra-accession variance was taken as the environmental variance, and data for all individuals were used for calculating phenotypic variance. A conditional GWA analysis was conducted on mean *L_p* values using the GWAPP web interface (<https://gwas.gmi.oeaw.ac.at/>) and the accelerated mixed-model (AMM) algorithm method with cofactors²¹. Genotype data for 206,087 SNPs from 250k SNP chip²⁰ were used to carry out GWA analysis, and only SNPs with MAF > 0.05 (equivalent to a minor allele count (MAC) ≥ 8) were considered. For conditional analysis, four successive steps were performed following an initial calculation with AMM. At each step, one among the nine most highly associated SNPs identified in the previous calculation was arbitrarily selected as a cofactor. In practice, four SNPs (Chr1-P13,612,169; Chr 5-P25,787,448; Chr 5-P21,846,701, and Chr4-P17,518,747) were stepwise included as cofactors. To correct for multiple testing, a Bonferroni correction with a nominal significance threshold (α) of 0.05 was applied, corresponding to an uncorrected *P* value of 2.61 × 10⁻⁷. The proportion of genetic variance explained by SNP (13,612,169/Chr1) and SNP (25,787,448/Chr 5) was determined using coefficients of determination from simple linear regressions. For *XND1*-based local association analysis, we extracted from Salk Arabidopsis 1001 Genomes database (<http://signal.salk.edu/atg1001/index.php>) genomic sequences of *XND1* (encompassing a region from 2 kb upstream to 300 bp downstream of *XND1* coding sequence) from 112 accessions. Of these, 85 belong to the initially investigated panel for GWA mapping (Supplementary Data 2). Association analysis between polymorphic sites (including INDELs and SNPs with MAF > 0.05) and *L_p* were performed with TASSEL version 5 using a generalized linear model⁴⁵. The significance threshold was set at a *P* value of 0.05 per marker number. Haplotypes were classified based on eight polymorphic sites showing the lowest *P* values according to *XND1*-based local association analysis. The haplogroups containing at least five accessions were used for further comparative analysis.

Genetic complementation of *xnd1*. A 7088 bp genomic region harboring *XND1* was amplified from Col-0, Bur-0, Ty-0, and Fei-0 genomic DNA using a iProof High-Fidelity PCR Kit (Bio-Rad) and cloned into a pGreen0179 vector⁴⁶. A QuikChange Site-Directed Mutagenesis Kit (Agilent stratagene) was used to mutate the SNP_{UTR} (Chr5_P25,795,349) of the cloned Col-0 or Bur-0 *XND1* fragments. All primer sequences used are listed in Supplementary Table 2. The constructs were

confirmed by sequencing and transferred into *xnd1-5* mutant plants using the floral dip method⁴⁷. For each construct, three to five independent homozygous transgenic lines were selected in T3 generation on 30 mg L⁻¹ hygromycin B (Sigma), checked for *XND1* expression (see below) and phenotyped for *L_p*.

Quantitative gene expression. *XND1* and *PIP* mRNA abundance was characterized in transgenic lines and/or natural accessions using qRT-PCR. Total RNA was extracted from *Arabidopsis* roots and reverse-transcribed using a SV Total RNA Isolation System (Promega) and M-MLV reverse transcriptase (Promega), respectively. PCR was performed on an optical 384-well plate with a LightCycler[®] 480 system (Roche) using SYBR Green I Master (Roche) or SYBR Premix Ex Taq (TaKaRa), according to the manufacturer's instructions. *TIP41-like protein* (At4g34270), *PP2A3* (At1g13320), and *SAND family protein* (At2g28390) were selected as reference genes⁴⁸, based on their expression stability among accessions evaluated using a NormFinder software⁴⁹. All primer sequences used are listed in Supplementary Table 2. Relative expression levels were determined using the 2^{(-ΔΔC(T))} method⁵⁰, and calibrated with respect to transcript abundance in the wild-type control, unless otherwise stated.

Characterization of RSA. Roots of hydroponically grown 22-day-old plants were harvested and preserved in 20% ethanol solution. The whole root systems were immersed in water and positioned on a square petri dish (24 cm × 24 cm), so as to avoid root overlapping, and then imaged using an Epson V850 Pro scanner at 600 dpi. Primary and total root lengths, and lateral root density were analyzed with an OPTIMAS software (version 6.1). The lateral root density was determined on the primary root, in a 12 cm region starting from the tip. Root DW was determined after desiccation at 80 °C for at least 24 h.

Root histological analyses. Roots of 21- to 23-day-old plants were cut in 2-cm-long segments starting from 0.3 cm of the tip. Root segments were embedded in 4% low-melting point agarose (Euromedex, 1670-B) and cross-sectioned (~100 μm thickness) using a Micro-Cut H1200 Vibratome (Bio-Rad) according to the manufacturer's instruction. Xylem morphology was observed under a BH-2 Bright field Microscope (Olympus) and quantified using ImageJ. Xylem vessels were identified from their thickened cell wall. Xylem size and abundance were assessed from the area of all vessels and vessel number, respectively. For testing the effect of flg22 on xylem formation, 19-day-old plants were exposed for 4 days to a hydroponic solution containing 0.25 μM flg22 or 0.025% DMSO as mock treatment, and xylem morphology was analyzed as described above.

Expression of fluorescent *XND1* fusion proteins. A 1465 bp region harboring *XND1* 5'-UTR and coding sequence, without stop codon and in its wild-type or SNP_{UTR}-mutated Bur-0 form, was amplified from above mentioned *XND1*-pGreen0179 clones. The fragment was cloned using the Gateway Technology (Invitrogen), downstream of a 35S*CaMV* promoter and in fusion with *GFP* in a pGWB505 vector⁵¹. The primers used are listed in Supplementary Table 2. The constructs were confirmed by sequencing and transferred into *xnd1-5* mutant plants. For each construct, nine to ten independent transgenic lines were selected in T2 generation on 30 mg L⁻¹ hygromycin B (Sigma), cultured in hydroponics and checked for *XND1-GFP* transcript abundance and GFP fluorescence intensity in roots. The primers used for qRT-PCR are listed in Supplementary Table 2. For quantification of the GFP fluorescence intensity, roots of 21- to 23-day-old plants were observed using a fluorescence microscope (Zeiss Axio Observer 7) and mean gray values in 400 μm root tips were quantified using an ImageJ program (NIH, USA). Data were normalized to the corresponding value of the line C9-3, which possessed the lowest transcript abundance and fluorescence intensity.

Drought stress treatments. Col-0 plants, *xnd1* T-DNA insertion and complementation lines were grown in trays filled with peat soil (Neuhaus Humin Substrat N2, Klasmann-Deilmann). Four to six trays were used per experiment. Each tray was divided in four quarters, each containing six plants of a specific genotype (quarter-split manner). The dimensions (length × width × depth) of the trays were 18 cm × 13 cm × 5.5 cm. Plants were maintained in a growth chamber at 20 °C and 65% relative humidity, with cycles of 8 h of light (250 μmol photons m⁻² s⁻¹) and 16 h of night, and sufficient watering. After 22 days, plants were subjected to drought by withholding water for 24 days and re-irrigated for 5 days. Gravimetric soil water content was around 30% at the end of the water deficit period. The control treatment was conducted in the same conditions, but with continuous watering. Shoot fresh weight (FW) was determined immediately after harvest whereas shoot DW was measured after further desiccation for at least 4 days at 60 °C. Shoot water content was calculated as the FW-to-DW difference. All data were normalized to the corresponding mean value of Col-0 plants in the same tray.

Bacterial inoculations. Seeds of Col-0 plants and *xnd1* T-DNA insertion lines were surface sterilized for 20 min with a 12% sodium hypochlorite solution, washed five times with sterile water and sown on a MS solid medium. After 8 days at 20 °C in a growth chamber, plantlets were transferred to Jiffy pots (Jiffy France, Lyon, France) and grown for 3 weeks under short days conditions at 22 °C and 70%

relative humidity with 9 h of light ($250 \mu\text{mol m}^{-2} \text{s}^{-1}$). Exposed roots of the plants were immersed for 20 min in a suspension containing 10^8 bacteria/mL of *R. solanacearum* GMI1000 strain⁵². Inoculated plants were then transferred to a new tray on a firm surface of potting soil, and incubated in a growth chamber at 75% relative humidity with cycles of 12 h of light ($100 \mu\text{mol m}^{-2} \text{s}^{-1}$) at 27 °C and 12 h of night at 26 °C. Plant position was randomized prior to inoculation. Symptom appearance was scored daily and independently for each plant, using a macroscopic scale describing the observed wilting: 0, no wilting; 1, 25% of leaves wilted; 2, 50%; 3, 75%; 4, complete wilting. For subsequent analysis, the data were transformed into a binary index: 0, < 50% leaves wilted; 1, \geq 50% wilted leaves in order to construct survival curves. We then applied the Kaplan–Meier survival analysis⁵³ with the Gehan–Breslow–Wilcoxon method to compute the *P* value and test the null hypothesis of identical survival experience of the tested mutant. A *P* value lower than 0.05 was considered to be significant. The survival curves represent a pool of three technical replicates, each with 24–32 plants, and 3 independent biological replicates corresponding to 240 plants for Col-0 and for each of the *xnd1* T-DNA insertion lines. For bacterial internal growth measurements, a *R. solanacearum* GMI1000 derivative strain carrying a gentamycin resistance cassette⁵⁴ was used. Rosettes of three to six pairs of plants were harvested at 3 and 4 days post-inoculation, sterilized in 70% ethanol and rinsed three times in sterile water. The rosettes were then weighted, grinded and re-suspended in sterile water. Bacterial concentrations were determined by plating dilutions on B medium. Four biological replicates were done. Comparison of *in planta* bacterial multiplication in *xnd1* genotypes with that in Col-0 was performed through a Mann–Whitney test. All statistical analyses were performed with a Prism version 5.0 software (GraphPad Software, San Diego, CA, USA).

Statistical analyses. Unless otherwise indicated, statistical significance of the data was assessed using either a Student's *t* test ($^*P < 0.05$) or one-way ANOVA (lowercase letters: $P < 0.05$). Student's *t* tests were performed using EXCEL whereas a STATISTICA software was used for ANOVA and multiple comparison tests (Fisher's least significant difference).

Data availability

The data supporting the findings of the study are available as Supplementary data or from the corresponding author on reasonable request.

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Author contributions

N.T. and C.M. conceived and designed the experiments with input from Z.S., O.L., and F.V. N.T. and Z.S. performed GWAS and phenotype screening of T-DNA insertion lines. F.L. and F.V. performed the *Ralstonia* infection assays. N.T. conducted all other experiments. N.T. and C.M. analyzed the data and wrote the paper which was checked by all authors.

Additional information

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