A mechanosensitive Ca\(^{2+}\) channel activity is dependent on the developmental regulator DEK1

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Responses of cells to mechanical stress are thought to be critical in coordinating growth and development. Consistent with this idea, mechanically activated channels play important roles in animal development. For example, the PIEZO1 channel controls cell division and epithelial-layer integrity and is necessary for vascular development in mammals. In plants, the actual contribution of mechanoperception to development remains questionable because very few putative mechanosensors have been identified and the phenotypes of the corresponding mutants are rather mild. Here, we show that the Arabidopsis Defective Kernel 1 (DEK1) protein, which is essential for development beyond early embryogenesis, is associated with a mechanically activated Ca\(^{2+}\) current in planta, suggesting that perception of mechanical stress plays a critical role in plant development.
Multicellular development is dependent upon cell–cell communication. Recent research has highlighted the fact that, in addition to chemical signals, the perception of mechanical stress, at both the cell and tissue level, is a key factor underlying growth coordination and morphogenesis (e.g., ref. 1). However, while in animal systems mechanosensors with important roles in development have been identified8–9, the molecular basis for mechanoperception in plants remains enigmatic. As in animals, plants respond to mechanical stimuli by an elevation in cytoplasmic Ca2+10–12. Based on the knowledge of plasma-membrane mechanosensins in animal systems, the most probable trigger for Ca2+ release from internal cellular compartments (Ca2+-induced calcium release) is either the opening of plasma membrane-localized mechanosensitive Ca2+ permeable channels, or the opening of voltage-dependent Ca2+ channels in response to changes in membrane potential caused by mechanosensitive channels permeable to other ions13. Recent research in Arabidopsis has led to the identification of several plasma membrane-localized mechanosensitive ion channels. These include proteins similar to the bacterial mechanosensitive channel of small conductance (the MSL family13–16), the MCA1 protein that rescues the yeast Ca2+ channel mutant mid1, and its homolog MCA217–20. In addition, the membrane protein OSCA1 forms a hyperosmolarity-gated Ca2+ permeable channel required for osmosensing in Arabidopsis21. However, although the MSL8 protein has recently been shown to be required for pollen grains to survive rapid rehydration during fertilization14, the very mild developmental phenotypes in single and multiple mutants of genes encoding the channels described above, suggests that they are unlikely to play a major role in mechanosensing during development. Furthermore, these proteins have not conclusively been shown to be responsible for any of the mechanosensitive Ca2+ currents which have been detected and extensively described by electrophysiologists over the past few decades in planta20, 22–26. Since genes encoding voltage-sensitive Ca2+ channels similar to those identified in animal systems have not been found in plant genomes, it has been proposed that plants may have evolved novel systems for mediating mechanosensitive Ca2+ fluxes at the plasma membrane to control development (reviewed in ref. 12).

The Defective Kernel 1 (DEK1) protein is encoded by a highly conserved uni-gene found in all multicellular plant genomes sequenced so far, and it is absolutely required for both embryonic and post-embryonic development in angiosperms27–31. Null dek1 mutant embryos do not develop beyond the early globular stage. When plants with reduced DEK1 activity can be obtained, they show major developmental defects, notably in epidermal differentiation and adhesion30, 32, 33.

The DEK1 protein contains multiple predicted transmembrane (TM) spans interrupted by a loop, and an intracellular tail including a linker domain and a C-terminal domain showing similarity to animal calpains, a class of Ca2+-dependent cysteine

**Fig. 1** A mechanically activated current permeable to Ca2+ is present at the plasma membrane. **a** Representative membrane patches from Col-0 (left) and an msl quintuple mutant msl4;msl5;msl6;msl9;msl10 (msl15, right), exposed to increased positive pressure steps in an outside-out patch configuration show a rapidly activated, rapidly inactivated current in ionic conditions favorable for Ca2+ current recording. We have named the channel responsible for this activity the RMA channel (see text). Time constants are means ± SE (n = 6); **b** Under the same conditions, single channel I/V curves show similar RMA channel conductance in Col-0 (solid square) and in the msl15 mutant (open square). Values are means ± SE (n = 6); **c** Representative single channel recordings showing that the RMA current is reversibly eliminated by exchanging Ca2+ ions with non-permeant TEA+ ions. **d** Open probability (for n channels, (nP(o))) is severely decreased in TEA+ bath solution. A paired t-test was used to compare means (**P < 0.01). Values are means ± SE. For all experiments, the membrane potential was held at −196 mV. Ionic conditions are described in the Methods section.

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proteases. DEK1 localizes to the plasma membrane and to internal compartments. In their original model of DEK1, Lid et al. predicted the presence of 21 TM domains with an extracellular localization for the loop and cytoplasmic localization (subsequently confirmed in planta) for the C-terminus. However, in a more recent analysis, Kumar et al. proposed a consensus model with 23 TM domains and with a cytoplasmic localization for the loop. Structure-function studies have shown that the cytosolic CALPAIN domain of the protein can, alone, complement the embryo lethality of *dek1* mutants suggesting that this domain, which is removed from the rest of the protein by an autolytic cleavage event, represents an active form of the DEK1 protein. Consistent with this scenario, and with a role for DEK1 in maintaining epidermal integrity, overexpression of the CALPAIN domain of DEK1 leads to thickening of the outer epidermal cell wall in leaves, and increased deposition of pectins, which are important for cell adhesion (reviewed in ref.37). The epidermis is thought to be under tension during much of plant growth. One plausible function of DEK1 could, therefore, be to perceive and respond to this tension to coordinate epidermal development and maintain epidermal integrity. As the epidermis plays a critical role in organ growth in plants (e.g., refs. 39–42), this, in turn, has major implications for plant development as a whole.

The activity of the CALPAIN domain has been shown to be Ca2+-dependent in vitro. These findings are in line with results in animal systems where the activation of cytoplasmic calpains is associated with the Ca2+-dependent autolytic-removal of an N-terminal extension, which in some cases can inhibit calpain activity. Because calpains have been shown to act downstream of mechanosensitive channels such as PIEZO in animals, we tested the hypothesis, and demonstrated, that the TM domain...
of DEK1 is associated with Ca\textsuperscript{2+} dependent mechanoperception at the plasma membrane in plants.

**Results**

**A rapidly mechanically activated plasma membrane Ca\textsuperscript{2+} channel.** Combining genetic and electrophysiological approaches, the *Arabidopsis* MSL9 and MSL10 proteins have been shown to mediate a mechanically activated channel activity in protoplasts derived from wild-type plant root cells\textsuperscript{13}. Selectivity characterization indicated that anions permeate preferentially through the wild-type MSL channels\textsuperscript{13}. Because in this work we used excised membrane patches from callus protoplasts, rather than whole root cell-derived protoplasts, we first investigated whether mechanically activated ion currents due to MSLs and other unidentified channels could also be detected in this material.

Reverse transcription quantitative-PCR (RT-qPCR) analysis confirmed the expression of MSL9 and MSL10 in wild-type (Columbia-0) callus tissue (Supplementary Fig. 1a). When Cl\textsuperscript{–} is provided at the cytosolic face of plasma membrane patches, a mechanically activated channel with a high conductance (46 ± 1.4 pS; ± indicates standard deviation, n = 6) is elicited upon application of positive pressure using a high-speed pressure clamp, as previously described in root protoplasts (Supplementary Fig. 1b)\textsuperscript{13}. This activity, which is mainly due to MSL9 and MSL10, was completely abolished in protoplasts obtained from quintuple *msl4 msl5 msl6 msl9 msl10* (*msl\textsuperscript{Δ}5*) mutant callus (Supplementary Fig. 1c). In addition, to MSL activity, we observed a rapidly activated-inactivated current with a smaller conductance, elicited immediately after the pressure increase (Supplementary Fig. 1b). Unlike the situation for the MSL current, removing Cl\textsuperscript{–} from the cytosolic face of the membrane did not modify this rapid current (Fig. 1a, Supplementary Fig. 1b). The rapid current was still present in protoplasts derived from *msl\textsuperscript{Δ}5* mutant callus with or without Cl\textsuperscript{–} ions at the cytosolic face (Fig. 1a, Supplementary Fig. 1c). In addition this
Because the dek1-2 CALPAIN-OE background cannot produce the intact N-terminal region of DEK1, we next tested the potential effect of DEK1 TM domain disruption on the RMA channel activity. In the majority of responsive patches obtained from dek1-2 CALPAIN-OE calli we observed a complete absence of the RMA current upon membrane stretching (Fig. 3a, Supplementary Fig. 7). Only a residual current was detectable in most patches of this background. In patches from dek1-3 CALPAIN-OE calli the RMA channel activity was still detactable, but showed 6–8 times more rapid inactivation kinetics (Fig. 3a, b, Supplementary Fig. 7) compared to those observed in callus from Col-0 plants.

The production of viable dek1 mutant plants, as mentioned before, requires the expression of at least the CALPAIN domain of DEK1. In our lines, the CALPAIN domain is overexpressed in the callus system at the RNA level (Supplementary Fig. 4). To confirm that the loss of the RMA channel activity in the dek1-2 CALPAIN-OE background is not a consequence of CALPAIN overexpression, we generated calli from Col-0 plants overexpressing the CALPAIN domain (Supplementary Figs. 4d and 5) 18. This material was submitted to electrophysiological analysis. The RMA current was still present in patches from this material, and behaved similarly to the current in Col-0 patches. (Fig. 3a, b, Supplementary Fig. 7). Therefore, we conclude that the CALPAIN domain overexpression per se is not responsible for the loss of RMA channel activity in the dek1-2 CALPAIN-OE background.

Loss of the DEK1 TM domains does not affect MSL activity. Next, we investigated whether the removal of the DEK1 TM domains could cause a perturbation of other mechanoresponsive channel activities at the plasma membrane by testing whether MSL channels can be activated in the dek1-2 CALPAIN-OE line. We found that the MSL activity in dek1-2 CALPAIN-OE protoplasts was still detectable upon membrane stretching, and its characteristics were comparable to that of Col-0 (Supplementary Fig. 8). Thus, loss of the DEK1 TM domains leads to a specific perturbation of the mechanosensitive RMA channel activity in our system.

DEK1 function influences root growth inhibition by Gd³⁺ ions. Based on the mild developmental defects observed in the dek1 CALPAIN-OE lines 32, 34, the CALPAIN domain of DEK1 is likely the “active” domain of the protein. The DEK1 CALPAIN domain is autolytically cleaved from the full length DEK1 protein 32, and its activity has been shown to be Ca²⁺ dependent in vitro 43. Because the release of the CALPAIN domain upon DEK1 cleavage could be triggered by Ca²⁺ influx via the RMA channel, blocking the RMA channel might affect development in wild-type plants, but would have a reduced impact in the dek1-2 CALPAIN-OE line, where the expression of a “pre-cleaved” version of the DEK1 CALPAIN would partially bypass the need for the RMA channel activity. Considering the sensitivity of the RMA current to Gd³⁺, and the absence of this current in CALPAIN complemented dek1-2 mutants we tested whether CALPAIN complemented dek1-2 mutants might show such reduced sensitivity to Gd³⁺. Gd³⁺ dramatically inhibits Arabidopsis root growth 57. We generated a dose sensitivity curve of wild-type Col-0 seedlings to concentrations of Gd³⁺ between 0 and 200 μM (Supplementary Fig. 9). We found strong growth inhibition (40–50%) in wild-type seedlings even at the lowest concentrations tested (30 and 60 μM) (Supplementary Fig. 9). This inhibition was alleviated in CALPAIN complemented dek1-2 plants (Supplementary Fig. 10), formally relating DEK1 function to Gd³⁺ sensitivity in planta.
Reintroducing full-length DEK1 restores RMA current. We hypothesize that the loss of RMA current observed in dek1-2 CALPAIN-OE callus is caused by the absence of an intact DEK1 transmembrane region. We propose that in the dek1-3 CALPAIN-OE plants a truncated DEK1 protein, which contains the DEK1 transmembrane span but which cannot confer normal inactivation kinetics on the RMA current is produced (Supplementary Fig. 2c). To test both these hypotheses we investigated the RMA channel activity in both the previously published and characterized dek1-3 line complemented with the full length GFP-tagged DEK1 protein expressed under the RPSSA promoter (dek1-3 DEK1) 30, and in a dek1-2 mutant background into which the same, complementing full length DEK1-encoding construct had been introduced by crossing (dek1-2 DEK1). In the majority of patches obtained from these lines, we recorded an RMA channel activity with inactivation kinetics and current amplitude resembling those of wild-type plants (Fig. 3a, b; Supplementary Fig. 7). The properties of this restored current were tested in the complemented dek1-2 DEK1 mutant callus and found to be strikingly similar to those of the RMA current (Supplementary Fig. 11). Our results show that reintroducing the full length DEK1 into dek1 mutant backgrounds leads to the restoration of an RMA current similar to that observed in wild-type callus.

Discussion

In summary, we show that the mechanosensitive activity of a Ca2+-permeable channel present in the plasma membrane of Arabidopsis callus-derived protoplast requires the TM domains of the DEK1 protein. It has previously been demonstrated that the CALPAIN domain of DEK1 is released from the plasma membrane by an autolytic cleavage event 32, and that the CALPAIN activity is enhanced by Ca2+ ions 43. We therefore propose a model in which DEK1 activity leads to transient elevation of cytoplasmic Ca2+ concentration during mechanical stimulation, which is locally transduced by autolytic cleavage (and thus activation) of the CALPAIN domain 32. Based on the phenotype of lines with reduced DEK1 activity, this mechanotransduction pathway is likely required for the maintenance of cell–cell contacts and epidermis integrity 33, consistent with embryo lethality in loss of function alleles. In turn, DEK1-dependent epidermis integrity is required for the propagation of mechanical signals between neighboring cells, in a feedback loop. Given the absolute requirement for DEK1 in both coherent embryogenesis and post-embryonic growth 29–31, 33 (summarized in Supplementary Fig. 12), our work suggests a key contribution of mechanical perception to plant development. The absence of cell migration in plant tissues may explain why the perception of tension between adjacent cells plays such an essential role in development in this kingdom.

This study, together with previously published results from both our work and the work of others 32, 50 supports the idea that the CALPAIN domain of DEK1 is the effector component of the protein in terms of mechanotransduction. Whether the transmembrane portion of DEK1 forms a mechanically activated Ca2+ channel per se, or it is the mechano-sensor associated with an independent Ca2+ channel remains to be determined. However, the published Ca2+ dependence of CALPAIN catalytic activity, combined with our electrophysiological results and the fact that CALPAIN complemented dek1 null mutants develop relatively normally, suggests that once activated by cleavage, the CALPAIN domain of DEK1 can respond to changes in cytoplasmic Ca2+ levels mediated by other Ca2+ channels. These could include proteins such as MCA1 and MCA2, or potentially the plant PIEZO protein, which has yet to be functionally characterized. The presence of such channels is consistent with our finding that CALPAIN complemented dek1-2 plants retain some Gd3+ sensitivity in planta.

Interestingly, intracellularly localized calpains have been proposed to act downstream of mechanosensitive ion channels in animals, to regulate a variety of cellular processes, including cell-to-cell adhesion 58–61. In this context, the association of a calpain protease with a domain influencing a mechanosensitive Ca2+ channel activity in the membranes of multicellular plants may reflect evolutive convergence between these two kingdoms.

In animals, the characterization of mechanosensitive channels Piezo1 and Piezo2 has revealed novel roles of mechanoperception in development and physiology. For instance, respiration in lungs relies on Piezo2-expressing sensory neurons, which use mechanical signals to sense airway stretching 62. Axon growth and trajectory was also shown to depend on Piezo1-dependent perception by the mechanical environment of neurons 53. The association of DEK1, a key developmental regulator, with a mechanosensitive channel activity paves the way to the identification of new roles of mechanical signals in plant development.

Methods

Plant material. Arabidopsis thaliana (Arabidopsis italiana) Columbia-0 (Col-0) wild-type seeds were obtained from Nottingham Arabidopsis Stock Centre (NASC, School of Biosciences, University of Nottingham, United Kingdom). Plants expressing the prpSSA:CALPAIN-HIS in a wild-type background (CALPAIN-OE), prpSSA: CALPAIN-GFP, or prpSSA:DEK1 in the dek1-3 mutant background and prpSSA: CALPAIN-GFP in the dek1-2 background have been previously described 18, 30. The prpSSA:DEK1 construct was transferred from the dek1-3 to the dek1-2 background by crossing, and plants were genotyped as described in Supplementary Fig. 3.

Plant and callus growth conditions. For in vitro cultures, seeds were surface-sterilized with chlorogenic acid, sown on square plates, and stratified for 2 or 3 days in the dark at 4 °C. After stratification, seeds were germinated in a growth chamber under a 16-h light/8-h dark cycle at 21 °C.

Callus generation. Surface-sterilized seeds were sown on “initiation medium” containing 4.3 g/L Murashige and Skoog salts (MS, Sigma-Aldrich), 2% sucrose, 10 mg/L myo-inositol, 100 µg/L nicotinic acid, 1 mg/L thiamine-HCl, 100 µg/L pyridoxine-HCl, 400 µg/L glycine, 0.23 µM kinetin, 4.5 mM 2,4-D, 1% Phytagel (pH 5.7). For callus generation, seeds were cultured in a growth chamber for 15 days. Cali were then transferred onto “maintenance medium” containing 4.3 g/L MS salts (Sigma-Aldrich), 2% sucrose, 10 mg/L myo-inositol, 100 µg/L nicotinic acid, 1 mg/L thiamine-HCl, 100 µg/L pyridoxine-HCl, 400 µg/L glycine, 0.46 µM kinetin, 2.25 µM 2,4-D, 1% Phytagel (pH 5.7), and sub-cultured every 15 days onto fresh “maintenance medium”.

Protoplasting protocol. Cali were digested for 15 min at 22 °C under hypotonic conditions (2 mM CaCl2, 2 mM MgCl2, 1 mM KCl, 10 mM MES (pH 5.5), 0.2% cellulysin (Calbochem), and 0.2% cellulase RS (Onozuka RS, Yakult Honsha Co.), 0.004% pectolyase Y23 (Kikkoman Corporation), 0.35% bovine serum albumine and mannitol to 600 mM. For enzyme removal, the preparation was washed twice with 2 mM CaCl2, 2 mM MgCl2, 10 mM MES (pH 5.5), and mannitol to 600 mM. For protoplast release, the preparation was incubated with 2 mM CaCl2, 2 mM MgCl2, 10 mM MES (pH 5.5), and mannitol to 280 mM. The suspension was filtered through a 50 µm nylon mesh.

Electrophysiology. Patch-clamp experiments were performed at room temperature with a patch-clamp amplifier (model 200 A, Axon Instruments, Foster City, CA) and a Digidata 1322 A interface (Axon Instruments). Currents were filtered at 5 kHz, digitized at 20 kHz, and analyzed with pCLAMP8.1 and Clampfit 10 software. During patch-clamp recordings, cells were held at a potential of 60 mV and current was measured with a pA scale. The composition of the pipette solution and pressure was applied with a high speed pressure-clamp system (ALA Scientific Instrument, NY), allowing the application of precise and controlled pressure pulses in the pipette 15, 54. For Ca2+ and Ba2+ current recordings, both bath solutions contained: 50 mM CaCl2 or 50 mM BaCl2, respectively, and 3 mM MgCl2. 10 MES-Tris (pH 5.6): while pipettes were filled with: 150 mM CaMES, 2 mM MgCl2, 5 mM EGTA, 4.2 mM CaCl2, and 10 mM Tris-HEPES (pH 7.2), supplemented with 5 mM MgATP. To remove Ca2+, a solution containing: 100 mM TrisCl, 5 mM MgCl2 and 10 mM Tris-MES (pH 5.6) were used. For Gd3+: current recordings, bath solution contained (mM): 50 CaCl2, 5 MgCl2, 10 MES-Tris (pH 5.6) and pipettes were filled with (mM): 150 CaCl2, 2 MgCl2, 5 EGTA, 4.2 CaCl2, and 10 Tris-HEPES (pH 7.2), supplemented with 5 mM MgATP.
inhibitor treatments, 0.25 mM LaCl3 or 0.25 mM GdCl3 were added to the bath solution, osmolality was adjusted with mannitol to 450 mOsmol for the bath solution and to 460 mOsmol for the pipette solution using an osmometer (Type 15, LÖSER Meßtechnik). Gigaohm resistance seals between pipettes (pipette resistance, 0.8–1.5 MΩ) (coated with Sylgard (General Electric) pulled from capillaries (Kimax, KIMAX Glass) and protoplast membranes were obtained with gentle suction leading to the whole-celled configuration, then excised to an outside-out configuration. The current inactivation kinetics were fitted with a mono-

Genomic DNA extraction and genotyping. Plant callus DNA was extracted using a rapid CTAB isolation technique32. dek1-2 and dek1-3 mutants were genotyped using the primers shown in Supplementary Table 1 and Supplementary Fig. 3. The following thermal profile was used: 95 °C for 5 min, 35 cycles of 95 °C for 45 s, 60 °C for 45 s, 72 °C for 1 min, and a final extension step of 5 min at 72 °C.

RNA extraction and quantitative gene expression analysis. Callus material was collected and snap-frozen in liquid nitrogen for gene expression analysis. For each experiment, at least two independent biological replicates were used. Total RNA was extracted using the Spectrum Plant Total RNA Kit (Sigma-Aldrich). Total RNAs were digested with the Turbo DNA-free DNase I (Ambion) according to the manufacturer’s protocol. RNA concentrations were measured with a NanoDrop ND-1000 UV–Vis spectrophotometer (NanoDrop Technologies). One microgram of total RNA was reverse transcribed (RT) using the SuperScript Vilo cDNA Synthesis Kit (Invitrogen) according to the manufacturer’s instructions. PCR reactions were performed in optical 96-well plates in the StepOne Plus Real Time PCR System (Applied Biosystems). Five microliters of a 1:10 dilution of cDNA was amplified in 20 µl of reaction mix. The following thermal profile was used: 95 °C for 10 min, 40 cycles of 95 °C for 15 s, 60 °C for 30 s. Amplicon melting curves, were recorded after cycle 40 by heating from 60 to 95 °C with a ramp speed of 1 °C min⁻¹. Expression levels of each gene, relative to β-Tubulin, were determined using a 2⁻¹⁰⁰ C, where A is the coefficient, τ is the time constant, and C represents the maximum current intensity.

Antibody production. Anti-IaHll rabbit polyclonal antibodies were generated against a peptide from the predicted catalytic domain (RGDKQFTDQEFPPNC) of the Arabidopsis DEK1 protein using the PolyExpress Custom Polyclonal Antibody Production Package Service (GenScript).

Western blot analysis. Callus material for each genotype was snap-frozen in liquid nitrogen, thoroughly ground in pre-cooled porcelain mortars, and proteins were extracted using an extraction buffer containing 50 mM Tris-HCl pH 7.5, 150 mM NaCl, 1% Nonidet P-40, 10% glycerol, 1% deoxycholate, 1 mM EDTA and 1× protease inhibitor cocktail P9599 (Sigma-Aldrich). Samples were incubated on ice for 1 h, centrifuged for 15 min at 14,000 g at 4 °C to remove cell debris, and protein concentrations in supernatants were determined using a Bio-Rad protein assay (Bio-Rad). Equal amounts of proteins were loaded and resolved on 7.5% polyacrylamide/0.1% SDS gels. Proteins were transferred onto nylon membranes using an iBlot II dry blotting system (Invitrogen). Membranes were blocked over night at 4 °C using 1× PBS/0.2% Tween-20 (Sigma-Aldrich) with 3% non-fat milk (Reglaiat), incubated for 2 h at room temperature with rabbit polyclonal anti-IaHll (Genescript) or mouse monoclonal anti-alpha tubulin clone B-5-1-2 (Sigma-Aldrich, catalogue number T5168) primary antibodies at dilutions of 1:1000 and 1:2000, respectively. Secondary horseradish peroxidase-conjugated anti-rabbit and anti-mouse antibodies (Promega, catalogue numbers S3731 and S3721, respectively) were used at a dilution of 1:45,000. Blots were incubated with Super Signal West Femto reagents (Thermo Scientific) according to the manufacturer’s instructions, and then exposed to Super RX film (Fujifilm). Membranes were first incubated with anti-IaHll antibodies, and then re-probed with anti-alpha tubulin antibodies.

Gadolinium sensitivity experiments. Seeds were surface-sterilized with chlorine gas, sown on square plates (in a single row in the upper part of the plates), and stratified for 2 days in the dark at 4 °C. After stratification, seeds were germinated in a growth chamber with 16 h light/8 h dark with plates kept in a vertical position. Plates containing 1/5 strength (0.86 g/l) MS medium (Duchefa) pH 5.7, 1% sucrose and 0.8% phytagel, were supplemented with gadolinium (III) chloride hexahydrate (Sigma-Aldrich). Primary root length was scored after 5 or 6 days of growth with ImageJ software. Different genotypes were sown side by side on the same plate and in multiple combinations to buffer possible positional effects.

Data availability. The authors declare that all data supporting the findings of this study are available within the manuscript and its Supplementary Files or are available from the corresponding authors upon request.
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