

# *Arabidopsis* JACKDAW and MAGPIE zinc finger proteins delimit asymmetric cell division and stabilize tissue boundaries by restricting SHORT-ROOT action

David Welch,<sup>1</sup> Hala Hassan,<sup>1,3</sup> Ikram Blilou,<sup>1,3</sup> Richard Immink,<sup>2</sup> Renze Heidstra,<sup>1</sup> and Ben Scheres<sup>1,4</sup>

<sup>1</sup>Molecular Genetics Group, Department of Biology, Utrecht University, 3584 CH Utrecht, The Netherlands;

<sup>2</sup>Plant Research International, 6708 PD Wageningen, The Netherlands

In the *Arabidopsis* root, the SHORT-ROOT transcription factor moves outward to the ground tissue from its site of transcription in the stele and is required for the specification of the endodermis and the stem cell organizing quiescent center cells. In addition, SHORT-ROOT and the downstream transcription factor SCARECROW control an oriented cell division in ground tissue stem cell daughters. Here, we show that the JACKDAW and MAGPIE genes, which encode members of a plant-specific family of zinc finger proteins, act in a SHR-dependent feed-forward loop to regulate the range of action of SHORT-ROOT and SCARECROW. JACKDAW expression is initiated independent of SHORT-ROOT and regulates the SCARECROW expression domain outside the stele, while MAGPIE expression depends on SHORT-ROOT and SCARECROW. We provide evidence that JACKDAW and MAGPIE regulate tissue boundaries and asymmetric cell division and can control SHORT-ROOT and SCARECROW activity in a transcriptional and protein interaction network.

[*Keywords:* Meristem; protein movement; stem cells; transcription factor; nuclear localization]

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The intercellular movement of transcription factors has emerged as a regulatory mechanism to direct higher plant development (Barton 2001; Wu et al. 2002). In the *Arabidopsis* root, radial patterning and growth depend upon the transcription factor SHORT-ROOT (SHR) that can move between cells (Helariutta et al. 2000; Nakajima et al. 2001). The root displays a radial organization in which concentric epidermal and ground tissue layers surround a central stele. In addition, a distal stem cell group that extends the tissues of the root is maintained by an organizing center, the quiescent center (QC). Together, the stem cells and the QC form a stem cell niche (Fig. 1a). SHR moves outward from its site of transcription within the stele, and it is required for the specification of QC and ground tissue identity (Helariutta et al. 2000; Nakajima et al. 2001; Sabatini et al. 2003). In *shr* mutants, the asymmetric division of the cortex/endodermis stem cell daughter (CED) does not

occur, resulting in a single layer of ground tissue with cortex identity (Benfey et al. 1993). SHR promotes the expression of a related transcription factor, SCARECROW (SCR), which also patterns the QC and ground tissue (Di Laurenzio et al. 1996; Sabatini et al. 2003).

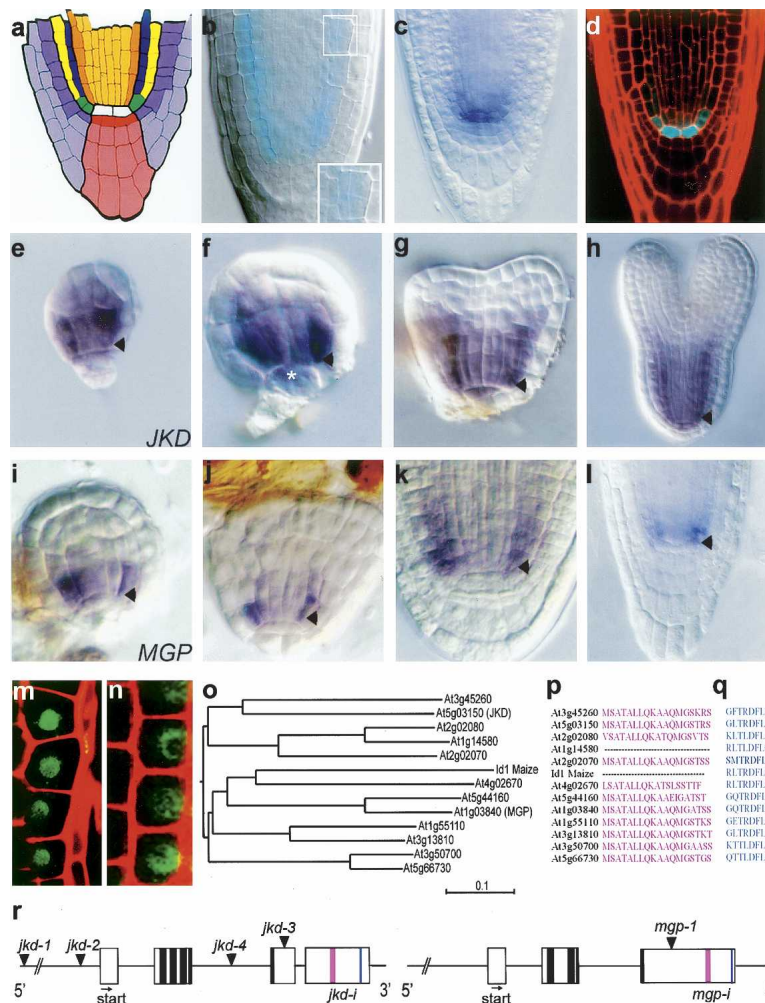
Loss of SCR function produces roots with a single layer of ground tissue with mixed endodermal and cortical identity (Di Laurenzio et al. 1996). Cell-autonomous action of SCR in the QC maintains QC identity and the activity of the surrounding stem cells (Sabatini et al. 2003). Clonal activation and deletion of SCR suggests a second cell-autonomous requirement for SCR in the rotation of the cell division plane and asymmetry of division in the ground tissue and a third role in mature endodermis cells in restricting SHR expression beyond a single ground tissue cell file (Heidstra et al. 2004). Ectopic SHR expression also suggests that in the wild type, SHR movement must be restricted to a single cell layer surrounding the stele (Sena et al. 2004; Nakajima et al. 2001). Recent data reveal that SHR protein interacts with SCR for joint up-regulation of SCR expression, and reveal how SCR can restrict SHR in a feed-forward loop (Levesque et al. 2006; Cui et al. 2007).

<sup>3</sup>These authors contributed equally to this work.

<sup>4</sup>Corresponding author.

E-MAIL [b.scheres@bio.uu.nl](mailto:b.scheres@bio.uu.nl); FAX 31-30-2513655.

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**Figure 1.** *JKD* encodes a C2H2 zinc finger expressed in the QC and ground tissue. (a) Median longitudinal view of the *Arabidopsis* root meristem (RM). (White) QC; (red) columella stem cells; (pink) columella root cap; (light purple) lateral root cap; (purple) epidermis; (green) cortical/endodermal stem cells; (yellow) cortex; (blue) endodermis; (orange) pericycle; (light orange) provascular cells. (b) GUS staining of the DWK15 promoter trap line (*jkd-1*) at 3-dpg; *inset* shows ectopic periclinal cortex division. (c) Whole-mount in situ hybridization with a *JKD* probe in 3-dpg wild-type Col-0 seedling. (d) Longitudinal confocal section of *pJKD::CFP*. (e–h) Whole-mount in situ hybridization with *JKD* in wild-type 32-cell stage (e), globular stage (f), early heart stage (g), and torpedo stage (h) embryos. (i–l) *MGP* expression during embryogenesis (i–k) and in 3-dpg wild-type seedlings (l). Arrowheads mark cortex/endodermis stem cells. Asterix marks the QC cells. (m,n) Longitudinal confocal section of *35S::JKD::GFP* and *35S::MGP::GFP* in *Arabidopsis* root epidermal cells. (o) Phylogenetic tree of *JKD/ID1* protein family. (p,q) Amino acid sequence alignment of conserved C-terminal domains. (r) Schematic representation of *JKD* and *MGP* genes. Dark boxes indicate location of zinc finger domains, pink and blue boxes indicate conserved C-terminal domains. Arrowheads mark the site of T-DNA insertions, arrows mark the translation start.

The question arises as to which factors regulate the three spatially separate roles of SHR and SCR. Here, we describe the *JACKDAW* (*JKD*) and *MAGPIE* (*MGP*) genes, encoding members of a plant-specific family of C2H2 zinc finger proteins, expressed from early embryogenesis onward in a subset of the SHR/SCR expression domain. We show that *JKD* and *MGP* contribute to refining the range of SHR and SCR action. We provide evidence that *JKD* is required for radial patterning and stem cell maintenance and that its activity is counteracted by the SHR target *MGP*, and we demonstrate that *JKD* and *MGP* interact physically with SCR and SHR, indicating how the SHR/SCR feed-forward loop can be differentially regulated in the stem cell domain.

**Results**

*JACKDAW* and *MAGPIE* gene expression in the stem cell region and ground tissue depends on SHR and SCR

The moving transcription factor SHR specifies the QC and the endodermis as a single cell layer and regulates stem cell daughter divisions. To identify new factors in-

involved in this process, we screened ~15,000 T-DNA insertion lines harboring a promoterless β-glucuronidase (GUS) gene adjacent to the right border of the T-DNA (Bechtold et al. 1993) for expression patterns similar to SCR and mutations affecting asymmetric division in the ground tissue. We identified one line, DWK15, with strong GUS staining in the ground tissue and QC (Fig. 1b). DWK15 homozygotes reveal ectopic periclinal divisions in the ground tissue (Fig. 1b and see below). Based on the contrasting phenotype in the ground tissue to *scr* mutants, we designated this mutation as *jackdaw-1* (*jkd-1*). The flanking sequence of the insertion in *jkd-1* identified a T-DNA insertion residing in the promoter region of the *At5g03150* gene, 3.5 kb upstream of the predicted coding region (Fig. 1r).

*JKD* mRNA accumulates in the QC, the ground tissue stem cells, and to a lesser extent in mature cortex and endodermis cells (Fig. 1c). This pattern resembles the GUS staining pattern of the *jkd-1* promoter trap line, and 2.8 kb of the *JKD* promoter fused to CFP essentially recapitulates the expression pattern observed by in situ hybridization (Fig. 1d). *JKD* transcript first accumulates during the 16- to 32-cell stage of embryogenesis, where it is strongest in the lower tier ground tissue or its precur-

cells (Fig. 1e). At the globular stage, *JKD* transcript expands into the lens-shaped upper derivative of the hypophysis (Fig. 1f), and by heart stage (Fig. 1g) it takes on the QC and ground tissue expression found also in later-stage embryos (Fig. 1h). A translational *JKD*:GFP fusion driven by the constitutive CaMV 35S promoter localizes to the nucleus of *Arabidopsis* root cells (Fig. 1m).

The predicted *JKD* protein is a member of a large family of zinc finger proteins (Fig. 1o), defined by the Maize protein INDETERMINATE (ID1), a regulator of flowering time (Colasanti et al. 1998; Kozaki et al. 2004). The protein family is defined by the N-terminal ID domain, a highly conserved amino acid sequence consisting of a putative nuclear localization signal (NLS) followed by two C2H2 and two C2HC zinc finger motifs (Fig. 1r; Kozaki et al. 2004). We additionally identified two C-terminal domains of unknown function conserved in many members of this protein family (Fig. 1p,q). The two middle zinc fingers (Z2 and Z3) of ID1 are required for binding to a specific 11-base-pair (bp) DNA sequence (Kozaki et al. 2004), indicating that one function of ID-like proteins is transcriptional regulation.

We searched the AREX data set (Birnbaum et al. 2003) for ground tissue-expressed *JKD* homologs and identified *At1g03840*, which we designated *MAGPIE* (*MGP*), a gene independently discovered as a *SHR* target (Levesque et al. 2006). Like *JKD*, *MGP*:GFP protein localizes to the nucleus (Fig. 1n). *MGP* transcript accumulates in ground tissue and stele cells of embryos and 2-d post-germination (dpg) wild-type roots but, in contrast to *JKD*, mRNA is absent from the QC (Fig. 1i–l).

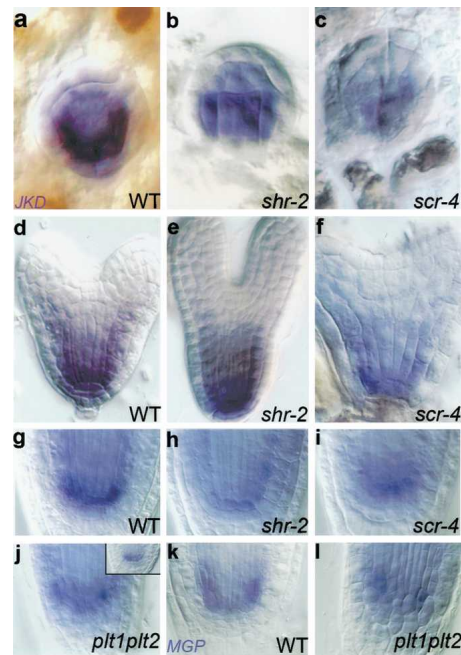
The initiation of *JKD* expression is not affected in *scr* and *shr* mutant embryos (Fig. 2a–f). However, post-embryonic QC and ground tissue stem cell expression of *JKD* is markedly reduced in *shr-1* and *shr-2* roots (Fig. 2h; data not shown) and less reduced in *scr-4* mutants (Fig. 2i; Supplementary Fig. 1g–i).

In contrast to our results with *JKD*, *MGP* transcript in a *shr-2* mutant background is barely detectable by in situ hybridization at any stage (Supplementary Fig. 1b,e), whereas expression is strongly reduced in *scr-4* (Supplementary Fig. 1c,f). RT-PCR analysis confirmed that *SHR* is required for *MGP* expression (Supplementary Fig. 2g,i).

Because both *JKD* and *MGP* transcription are closely associated with the stem cell niche, we assessed whether their expression is controlled by the *PLT1* and *PLT2* genes, which redundantly control QC specification and stem cell activity in a *SHR*- and *SCR*-independent way (Aida et al. 2004). In *plt1,plt2* double mutants, neither *JKD* nor *MGP* transcript accumulation was affected in seedlings (Fig. 2j–l) or embryos (data not shown).

#### *JKD* is required for QC specification and stem cell activity and maintains *SCR* expression

To study the function of the *JKD* gene, we identified three independent *jkd* insertion lines and constructed a line with reduced *JKD* levels by RNA interference (RNAi) referred to as *jkd-i* (see Materials and Methods). Severely reduced levels of *JKD* were detected in *jkd-4*

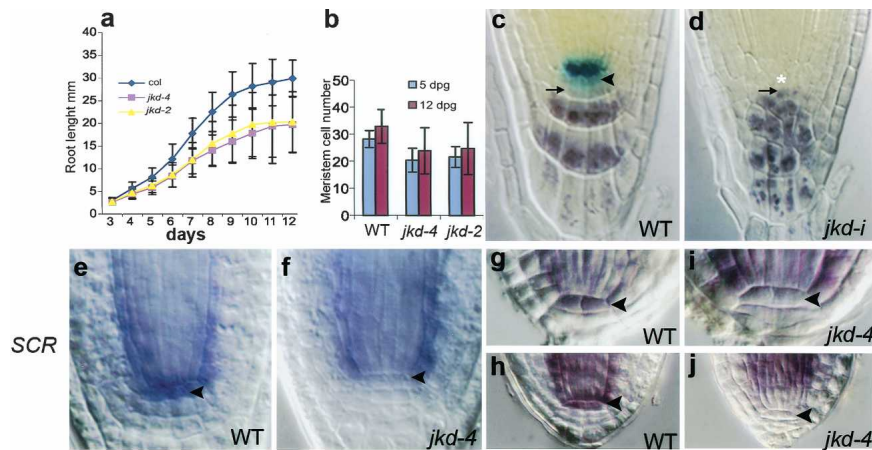


**Figure 2.** Regulation of *JKD* and *MGP* expression in the ground tissue. Whole-mount in situ hybridization of *JKD* (a–f) and *MGP* (g, i) in embryos of wild type (a, d), *shr-2* (b, e), and *scr-4* (c, f), and in 2-dpg seedlings of wild type (g, inset of j, k), *shr-2* (h), *scr-4* (i), and *plt1,plt2* (j, l).

homozygotes, while no signal was detected in *jkd-2* homozygotes (data not shown), indicating that *jkd-2* is a null mutant.

*jkd* mutants display a slight reduction in root length compared with wild-type (Fig. 3a), and ~15% of *jkd-4* and *jkd-2* seedlings show an early initiation of lateral roots (data not shown). To determine if root meristem size is also affected in *jkd* mutants, we measured meristem cell number in wild-type, *jkd-4*, and *jkd-2* seedlings. At both 5 and 12 dpg, the meristem cell number of *jkd-4* and *jkd-2* roots is reduced compared with that of wild type (Fig. 3b).

The reduction in meristem size and root growth suggested a role for *JKD* in QC/stem cell specification and/or maintenance. To investigate *JKD* function in the QC, we examined several independent QC markers in *jkd* mutants. We were unable to detect the QC-specific promoter trap QC46 (Sabatini et al. 1999) in *jkd* mutants (data not shown), while QC25 expression (Sabatini et al. 1999) is normal from germination to 7 dpg but disappears at 8–9 dpg, with a concomitant disorganization of the QC and columella (Fig. 3c,d). Cells at the position of the QC in *jkd* mutants of that age often neighbor columella cells containing starch granules, indicating the absence of columella stem cells (Fig. 3d). Expression of QC25 and QC46 requires cell-autonomous *SCR* activity (Sabatini et al. 2003). We could not detect *SCR* expression in the QC of *jkd-4* mutants (Fig. 3e,f) and detected no QC expression of *pSCR::YFP<sub>ER</sub>* in *jkd* mutant roots (Fig. 4b). To determine at what developmental time point *JKD* is required for QC expression of *SCR*, we analyzed *SCR*



**Figure 3.** *JKD* is required for QC identity and stem cell maintenance. Root length (a) and meristem cell number (b) are reduced in *jkd-4* and *jkd-2* compared with that of wild-type (Col-0). QC25 expression in 10-dpg wild-type (c) and *jkd-i* (d) roots. Lack of lugol staining marks columella stem cells in wild-type roots (arrowhead). Lugol-staining cells (arrow) designate differentiated columella cells, which neighbor defective QC cells (asterisk) in *jkd-i*. *SCR* expression in 2-dpg wild-type (e), and *jkd-4* (f) roots, and in wild-type (g,h) and *jkd-4* (i,j) embryos at heart stage (g,i) and torpedo stage (h,j). *SCR* is absent from *jkd* mutant QC cells (arrowhead) from heart stage onward.

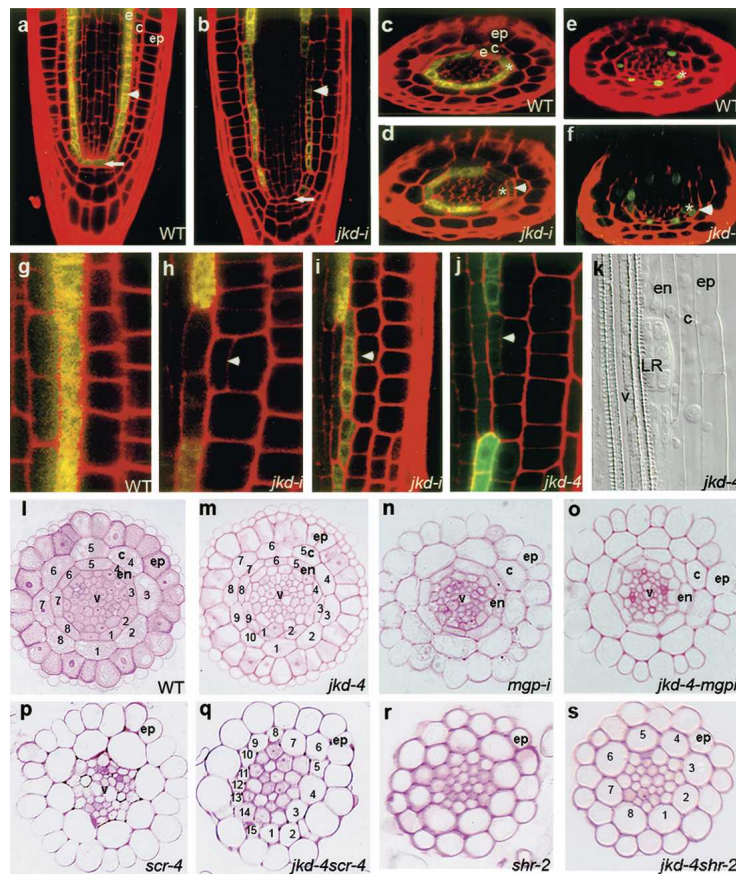
expression in *jkd-4* mutant embryos. From early heart stage onward, QC expression of *SCR* in *jkd-4* mutant embryos is absent (Fig. 3g-i), indicating a requirement for *JKD* in the regulation of *SCR* expression.

In *jkd-4 scr-4* and *jkd-i scr-3* double homozygotes, cells in the root meristem differentiate earlier and root lengths are shorter than in *scr* single mutants (Supplementary Fig. 2a,c), indicating that *JKD* regulates additional factors required for QC and stem cell maintenance. One candidate is the *SHR* protein, which regulates *SCR*-independent factors in the QC (Sabatini et al. 2003). To test whether *JKD* could act independently of *SHR*, we constructed *jkd-4 shr-2* and *jkd-i shr-2* double

mutants. We observed only a slight reduction in root length compared with *shr-2* single mutants (Supplementary Fig. 1b,d; *t*-test,  $P = 0.005$ ), indicating that *JKD* largely but not exclusively acts through *SHR* in the QC.

*JKD and MGP oppositely control SHR-SCR action in the ground tissue*

Wild-type roots contain ground tissue with an inner endodermis and a single outer cortex layer (Figs. 1a, 4a,c). In contrast, we observed ectopic periclinal divisions in the cortex of all *jkd* mutants generating ground tissue with three layers (Fig. 4b,d,f). Close to the root tip,



**Figure 4.** Mutations in *jkd* cause ectopic divisions and misexpression of *SCR* in the ground tissue. Longitudinal (a,b) and radial (c-f) confocal sections of 3-dpg wild-type root carrying *pSCR::YFP* (c) or *pSCR::SCR:GFP* (e) with two layers of ground tissue and *jkd-i* (d,f) roots with ectopic periclinal divisions in cortex cells (arrowhead). Arrow marks QC cells and asterisk mark the endodermal cells. (g-j) Longitudinal confocal sections of *pSCR::FP<sub>ER</sub>* in 3-dpg wild-type (g), *jkd-i* (h,i), and *jkd-4* roots (j). *SCR* is expressed in endodermal cells and is absent from cortex cells in wild type. Ectopic periclinal cortex divisions (arrowhead) in *jkd* mutants coincide with a reduction of *SCR* promoter activity in adjacent endodermis cells and induction of *SCR* promoter activity in the middle ground tissue layer. (k) Lateral root primordium emerging from the pericycle in *jkd-4*. (l-s) Root tissue sections from 10-d-old wild type (l) ([ep] epidermis; [c] cortex; [e] endodermis; [v] vascular bundle); *jkd-4* (m); *mgpi* (n); *jkd-4mgpi* (o) (note that cell number in the ground tissue layers is restored to wild type); *scr-4* (p); *jkd-4scr-4* (q); *shr-2* (r); and *jkd-4shr-2* (s).

patches of ectopic cells revealed that these derive from divisions of cortex cells based on the length of the adjacent cells relative to one another and the position of subsequent divisions (Fig. 4b,h–j). These cell clones join to form a continuous extra layer higher up in the meristem (Supplementary Fig. 2h–k).

To determine whether these ectopic divisions were asymmetric, we analyzed the expression of the *SCR* promoter fused to ER-targeted YFP or GFP, together referred to as *pSCR::FP<sub>ER</sub>*. In wild-type roots, *pSCR::FP<sub>ER</sub>* is found in the QC, ground tissue stem cells, and endodermis (Fig. 4a,c). In both *jdk4* and *jdk-i*, small clones of cortex cells (one to three cells) containing periclinal divisions do not show *pSCR::FP<sub>ER</sub>* expression. However, endodermal cells adjacent to these cells display a reduced level of *pSCR::FP<sub>ER</sub>* expression compared with endodermal cells above and below, and this reduction can be observed prior to the periclinal divisions (Fig. 4d,h–j). The inner cells of larger clones (more than three cells) with ectopic cortical periclinal divisions show expression of *pSCR::FP<sub>ER</sub>* (Fig. 4i,j), demonstrating that the ectopic divisions are asymmetric and that the inner cells take on endodermis characteristics. These new cells express also a SCR protein fusion (Fig. 4f; Supplementary Fig. 2h,i). Interestingly, in many cases, former endodermal cells adjacent to large clones no longer express *pSCR::FP<sub>ER</sub>* (Fig. 4i,j) and eventually acquire pericycle fate based on their capacity to generate lateral root primordia (Fig. 4k). These data, together with the reduction of *pSCR::FP<sub>ER</sub>* in cells adjacent to smaller clones, demonstrate a requirement for *JKD* in the restriction of *SCR* transcription in the ground tissue region.

We did not observe the ectopic periclinal divisions found in cortex cells of *jdk* mutants in *jdk-4 scr-4* and *jdk-4 shr-2* double mutants (Fig. 4p–s), indicating that these divisions require *SHR* and *SCR* and that *JKD* acts in the ground tissue by modifying the activity of *SHR* and *SCR*.

It is of note that the *jdk* mutant showed increased cortex and endodermis cell numbers within layers (Fig. 4l–m; Supplementary Table 1), revealing its requirement for restricting cell number in the ground tissue. This increase in numbers is suppressed in *jdk-4 shr-2* double mutants but not in *jdk-4 scr-4* mutants (Fig. 4p–s), suggesting that *JKD* controls cell numbers in the circumference of the ground tissue through *SHR* but independent of *SCR*.

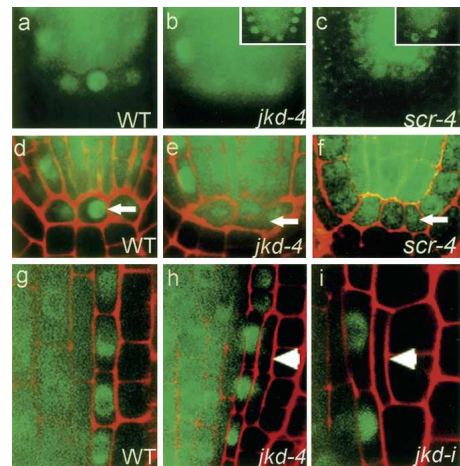
To assess whether MGP acts in a similar process as *JKD*, we analyzed MGP RNAi (*mgp-i*) lines in wild-type and in *jdk-4* background. While *mgp-i* lines reveal no phenotype on their own (Fig. 4n; Supplementary Fig. 2e), combination with *jdk-4* homozygotes largely complements the *jdk-4* ground tissue phenotype and restores *pSCR::SCR::GFP* expression (Fig. 4o; Supplementary Fig. 2g,l,m).

#### *JKD* affects intracellular SHR localization in the QC and SHR expression range in the ground tissue

To determine how *JKD* could affect *SHR* activity in the QC, we analyzed *pSHR::GFP::SHR* expression in the QC

of *jdk-4* and *jdk-i* roots. In contrast to the nuclear-localized GFP in wild-type roots (Fig. 5a,d), we observed cytoplasmic GFP:SHR in the majority of *jdk-4* and *jdk-i* mutant QC cells (Fig. 5b,e; data not shown). Cui et al. (2007) demonstrated a role for *SCR* in sequestering *SHR* within the endodermis. To determine whether loss of *SCR* function in the QC could affect GFP:SHR localization, we examined *pSHR::GFP::SHR* expression in *scr-4* mutants. Despite the low expression of *pSHR::GFP::SHR* in *scr-4* roots, GFP:SHR mostly localizes to both nucleus and cytoplasm in cells at the position of the QC (Fig. 5c,f); however, occasionally we did observe nuclear localization of *SHR* (Fig. 5c, inset), as in *jdk-4* (Fig. 5b, inset). Together, these results suggest that *JKD* controls the nuclear localization of *SHR* in the QC mostly through its effect on the maintenance of *SCR* expression.

Normal levels of *SCR* transcript in the endodermis require functional *SHR* (Helariutta et al. 2000). To determine if the alterations in *SCR* transcription in *jdk* mutants were due to changes in *SHR* expression, we analyzed *pSHR::GFP::SHR* expression in *jdk-4* and *jdk-i* roots. In wild type, GFP:SHR localizes to both cytoplasm and nuclei in stele cells, its site of transcription, and exclusively in the nuclei of endodermis cells (Fig. 5g; Nakajima et al. 2001). In large clones of cortex cells that had divided periclinally, we observed GFP:SHR in the inner cells (Fig. 5h), while smaller clones showed no GFP:SHR (Fig. 5i). All former endodermis cells still express nuclear-localized GFP:SHR (Fig. 5 h,i) at a stage when *SCR* promoter activity is already reduced (cf. Fig. 4h,i). The expansion of *pSHR::GFP::SHR* expression in *jdk* mu-



**Figure 5.** Subcellular localization of *SHR* in the QC and the ground tissue layers of *jdk* mutants. Longitudinal confocal sections of *pSHR::GFP::SHR* in wild type (a,d,g), *jdk-4* (b,e,h), and *scr-4* (c,f). (a–c) Green channel. (d–f) Overlay. Arrows mark GFP:SHR localization in the QC cells. *SHR* protein is nuclear in wild type and cytoplasmic in *jdk-4* and *scr-4*. Insets show nuclear *SHR* localization occasionally observed in both *jdk-4* and *scr-4*. (g) In wild-type roots, GFP:SHR localizes to the cytoplasm and nucleus in the stele and to the nucleus in the endodermis, and is absent from cortex cells. (h,i) Ectopic periclinal cortex divisions (arrowhead) in *jdk* mutants cause *SHR::GFP* expression in two of the three ground tissue layers.

tants is completely suppressed in the *mgpi* background (Supplementary Fig. 2n,o). These data show that the expression of new SCR in *jkd* associates with an expanded SHR domain, but lack of maintenance of SCR expression in the inner ground tissue layer is not caused by down-regulation of SHR.

*JKD and MGP physically interact with SCR and SHR*

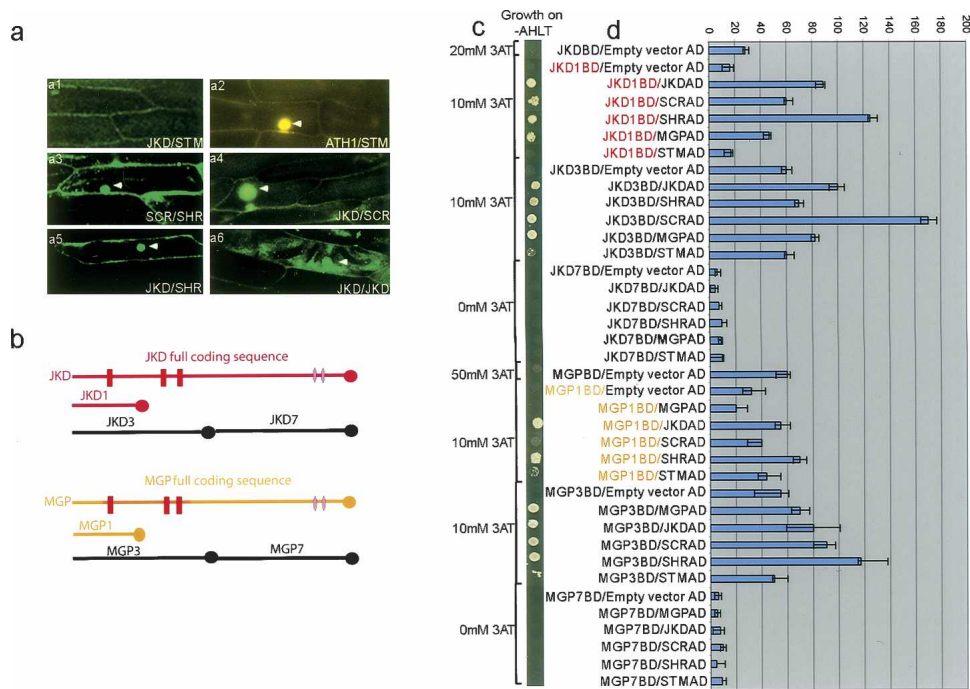
SHR and SCR proteins interact with each other, and a SHR-SCR complex binds promoters of SCR and MGP (Cui et al. 2007). To assess whether JKD and MGP could influence SHR and SCR activity at least in part through protein interaction in the stem cell area where they are jointly expressed, we performed in planta Bimolecular Fluorescence Complementation (BiFC) assays using JKD, MGP, SCR, and SHR cDNAs. We observed no fluorescence following coexpression of free nonfused YFP subfragments and negative control proteins (Supplementary Table 2). Consistent with reported interactions, fluorescence accumulated in nuclei of cells expressing SCR and SHR, and ATH1 and STM (Cole et al. 2006), but notably also in most of the combinations with JKD and MGP, suggesting multiple interactions (Fig. 6; Supplementary Table 2). These interactions were confirmed using the yeast two-hybrid system. We did not observe growth when fusion proteins (BD or AD) were tested with con-

trol empty vectors and when JKD and SCR were tested with a negative control protein. Next, we confirmed that SHR interacted strongly with SCR (Fig. 6c,d; Supplementary Fig. 2; see also Cui et al. 2007). Autoactivation domains in JKD and MGP were identified by comparing full-length cDNAs with fragments and eliminated from the analysis by selecting fragments with low autoactivation potential for subsequent experiments (Supplementary Figs. 2, 4).

Next, we showed that all interactions involving JKD and MGP required their first zinc finger domain (located between 107 and 215 for JKD and 95 and 202 for MGP, designated as JKD1 and MGP1) (Fig. 6b). In pairwise combinations we found that SHR interacted with JKD1 and MGP1 (Fig. 6; Supplementary Fig. 2); JKD1 could interact with itself and with both SCR and MGP1. We detected weak interaction between SCR and MGP1 but no self-interaction for MGP and SCR. Taken together, our data strongly indicate that JKD and MGP can be part of nuclear SHR-SCR complexes.

**Discussion**

Here, we demonstrate that JKD and MGP, two members of a large plant C2H2 zinc finger protein family, regulate the range of SHR action. *JKD* transcription is initiated early in the ground tissue by a SHR- and SCR-indepen-



**Figure 6.** JKD and MGP protein interaction studies. (a) In planta analysis using BiFC. Confocal laser scanning microscopy images of bombarded onion epidermal cells coexpressing JKD with SHOOT MERISTEMLESS (STM) (panel a1), SCR with SHR (panel a3), JKD with SCR (panel a4), JKD with SHR (panel a5), and JKD with itself (panel a6). (Panel a2) A. THALIANA HOMEBOX GENE1 (ATH1) was used as positive control for STM. (b-d) Yeast two-hybrid analysis. (b) Deletions used for mapping JKD and MGP interacting domains. (c) Growth rates on selective medium Ade-His-Leu-Trp-[AHLT] supplemented with different 3-amino triazol (3AT) concentrations; (d) Protein interactions quantified by measuring lacZ activity depicted as units of β-galactosidase. CPRG was used as the substrate. Error bars show standard deviation of the mean of triplicate assays.

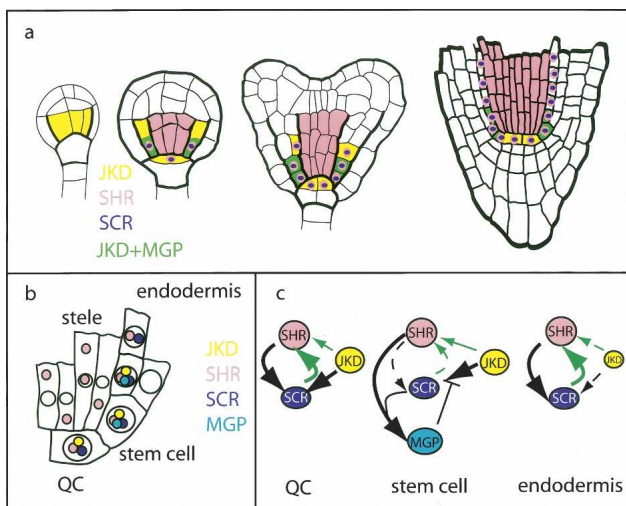
dent mechanism, whereas *MGP* is activated later and depends on SHR and SCR activity. Notably, SHR and SCR can directly bind to the *MGP* promoter, corroborating that *MGP* is a direct target (Levesque et al. 2006; Cui et al. 2007). Later on, maintenance of *JKD* also becomes dependent on SHR and SCR. Together, these transcriptional dependencies create an intriguing pattern of gene expression where the QC and ground tissue layers are marked by *JKD* expression, but *MGP* is transcribed only in those cells that perform asymmetric cell divisions (Fig. 7a). Protein–protein interactions among SHR, SCR, MGP, and JKD indicated by independent methods suggest that the proteins form nuclear complexes in cells where they are coexpressed (Fig. 7b). Our phenotypic analyses show that JKD promotes *SCR* transcription and SHR nuclear localization in the QC, and that it counteracts the occurrence of supernumerary SHR–SCR-mediated asymmetric cell divisions to regulate cell type specification and stable boundary formation. MGP function is harder to assess because single mutants reveal no defects. Even double mutants, where the coregulated and phylogenetically related *NUTCRACKER* gene activity (Levesque et al. 2006) also has been reduced, reveal no phenotype, suggesting extensive redundancy (data not shown). However, *MGP* reduction in the *jkd* background reveals a specific phenotype: In the ground tissue, JKD restricts the range of SHR action through counteracting a MGP-dependent cell division-promoting activity that

is also dependent on SHR and SCR. Given the abundant protein interactions, a plausible scenario is that MGP redundantly facilitates the asymmetric cell division-promoting activity of the SHR–SCR complex while being incorporated in it. To explain the epistasis of *mgp* over *jkd* mutants, JKD would have to inhibit this activity either by competing for binding in the same complex or by interactions within a complex that includes MGP. Future analysis on competitive binding of MGP and JKD on the SHR–SCR complex should distinguish between these possibilities.

In the QC, SCR is required for maintaining its own expression and nuclear localization of SHR (Heidstra et al. 2004; Cui et al. 2007; this study), and in the ground tissue, lower levels of SCR promote extra asymmetric divisions (Cui et al. 2007). Thus, the effects of *jkd* mutants resemble either elimination of SCR (in the QC) or reduction of SCR (in the ground tissue). Therefore, the major function of JKD might be to promote rapid up-regulation of *SCR* by SHR. The delay of this up-regulation by the presence of MGP could specifically poise the stem cell area to perform asymmetric divisions. The roles of JKD in nuclear sequestration of SHR in the QC and in restricting its range of action in the ground tissue might also be explained by restriction of SHR by JKD-regulated SCR, although we cannot exclude that JKD binding to SHR directly affects these processes. It is intriguing that *MGP* itself is a SHR target and that we could not establish a role in *MGP* transcriptional regulation for the second root stem cell regulatory system through the *PLT* genes (Aida et al. 2004). In this context, it is noteworthy that the complex transcriptional regulation and the potential to form different protein complexes indicate different and superimposed dynamic feedbacks that might establish different networks and different activities of SCR in different cells (Fig. 7c). Before all redundant partners in this network have been identified, it seems premature to speculate on the precise dynamics of this system. It is nevertheless gratifying that the presence of different networks in QC, stem cells, and endodermal cells correlates with the different activities in the ground tissue layer and with the different functions of SCR as deduced from clonal analysis.

In conclusion, SHR might be sequestered to the nucleus in different complexes by its downstream target SCR and MGP and by JKD, whose expression is set up independently.

*jkd* and *mgp* mutants have subtle phenotypes, which could argue against a pivotal role for JKD and MGP in control of SHR activity. However, we observed similar genetic interactions between *jkd* and a *mgp* homolog (I. Blilou, H. Hassan, and B. Scheres, unpubl.). Therefore, we suspect extensive redundancy within the JKD clade. Taken together, all available data hint at an elaborate mechanism to restrict the activity of SHR action. Although such a dynamic restriction of the activity range of moving transcription factors seems to be novel, the principle of controlling the movement of instructive factors other than transcription factors is broadly used. In the *Drosophila* wing disc, for example, the Hedgehog



**Figure 7.** JKD and MGP action in QC and ground tissue. (a) Summary of onset of expression of JKD, SHR, SCR, and MGP genes. Only SHR and SCR are drawn in the nucleus for simplicity. (b) Model incorporating the partially cytoplasmic localization of SHR in the stele and the presumed nuclear localization and protein interactions of JKD and MGP. Note the existence of three separate zones in QC and ground tissue with different protein composition. (c) Current view on the interaction network between SHR, SCR, JKD, and MGP revealing differences in QC, stem cells, and mature endodermis. (Black arrows) Transcriptional regulation, (green arrows) regulation of nuclear localization/protein movement. Arrow thickness represents regulation strength.

(Hh) protein, initially expressed in the posterior compartment, moves one cell layer into the anterior compartment where it binds to and up-regulates through a signal transduction cascade the expression of the transmembrane receptor Patched (Ptc). Binding to Ptc serves to limit the movement of Hh beyond this single cell layer (Chen and Struhl 1996; Nybakken and Perrimon 2002). It will be interesting to further investigate whether plants have evolved different restriction mechanisms directly at the level of instructive transcription factors.

## Materials and methods

### Plant material and growth conditions

*Arabidopsis thaliana* ecotypes Wassilewskija (WS) and Columbia (Col) were used. Origins and ecotypes of mutant and transgenic lines are as follows: *jdk-1* (WS), INRA T-DNA lines (Bechtold et al. 1993); *jdk-3*, *jdk-4*, and *jdk-2* (Col), the SALK t-DNA collection (<http://signal.salk.edu>); *scr-3* (Col) and *scr-4* (WS) (Fukaki et al. 1998); *shr-1* (WS) (Benfey et al. 1993); QC25 and QC46 (WS) (Sabatini et al. 2003); *shr-2* and *pSHR::GFP:SHR* (Col) (Nakajima et al. 2001); *pSCR::YFP* (Col) (Aida et al. 2004). Growth conditions of seedlings and embryos are described in Sabatini et al. (1999) and Scheres et al. (1995), respectively.

### Insertion sites of *jdk* mutants

For determination of the insertion site of *jdk-1*, the sequence flanking the left border (LB) of the T-DNA was amplified by Vectorette-PCR (Morrison and Markham 1995). The T-DNA insertion sites in *jdk-3*, *jdk-4*, and *jdk-2* are described in the Salk Institute Web site (<http://signal.salk.edu>) and were confirmed by PCR-based genotyping: *jdk-1*, -3450 bp from the predicted translation start point; *jdk-3*, 1940 bp from the predicted translation start point; *jdk-4*, 1381 bp from the predicted translation start point; and *jdk-2*, -206 bp from the predicted translation start point.

### Transgenic plants

The pGreenII vector set (Hellens et al. 2000; <http://www.pgreen.ac.uk>) was used for plant transformation. For the *35S::JKD:GFP* and *35S::MGP:GFP* translational fusion, xGFP was amplified by PCR from pGIIxGFP using the primers xGFP-L (5'-GCGAGCTCACTAGTAAAGGAGAAGAAGACTTTTCA-3') and xGFP-R (5'-CGGAATTCTCATTGTATAGTTCATCCATGCCATGT-3') and was cloned into a cassette containing the 35S promoter. The coding sequence of *JKD* was amplified by PCR using the primers JKD-HA-0C-L (5'-GCGGATCCATGCAGATGATTCCAGGATATCCAT-3') and JKD-HA-1N-R (5'-GCGAATTCTCAACCCAATGGAGCAAACC-3') from root cDNA and fused in-frame before xGFP. *35S::JKD:GFP* was transferred into pGreen carrying the methotrexate resistance cassette (Heidstra et al. 2004). For *pJKD:CFP*, a 3.5-kb fragment of the *JKD* promoter was PCR-amplified using the primers K15prom-dn-L (5'-GGC GCGCTGTTTCGATATCACATTTTGAC-3') and K15prom-dn-R-mk2 (5'-GCGCGTGCTTGACTCTTTGGTTATGCC-3') and placed before ER-CFP and the NOS terminator and transferred into *pGreenII* carrying the nos-basta resistance cassette. For the *JKD-i* and *MGP-i* constructs, 500-bp fragments of the *JKD* and *MGP* coding sequence were amplified with the primers K15ZFRNAi-L (5'-ATTCTAGACTCGAGCATCATCCCTCCCTGAT-3') and K15ZFRNAi-R (5'-ATATCGATGGTAC

CAACCTTGCGAGTTCTTGAGG-3') for *JKD* and MGPi FW primer (5'-CTCGAGGGATCCGACGCTTTAGCAGAAGAAA CCGC-3') and MGPi REV primer (5'-GGTACCATCGATATTGGTCCGGTAGTAATCGTCTGC-3') for *MGP*, and cloned into the pKANNIBAL vector (Wesley et al. 2001). *JKD-i* and *MGP-i* were transferred into *pGreenII* carrying the norflurazon resistance or methotrexate resistance cassette, respectively (Heidstra et al. 2004).

### Microscopy and mRNA detection

For DIC optics, seedlings and embryos were cleared and mounted according to Willemsen et al. (1998). Starch granules and  $\beta$ -glucuronidase activity were visualized as described (Willemsen et al. 1998). Root length was measured as described (Willemsen et al. 1998). The number of root meristematic cells was obtained by counting cortex cells showing no signs of rapid elongation. For confocal microscopy, roots were mounted in 10  $\mu$ M propidium iodide. In situ hybridization on whole-mount tissues was performed manually as described (Hejatko et al. 2006). Root tissue sections were performed as described (Willemsen et al. 1998).

### Yeast two-hybrid assay

Yeast two-hybrid interactions were studied using the ProQuest Two Hybrid System (Invitrogen Life Technologies). The coding sequences of *JKD*, *MGP*, *SCR*, and *SHR* were amplified and fused to both the pDEST32 BD and pDEST22 AD vectors. Primers used to amplify full-length cDNAs and deletion fragments are described in Supplementary Table 3.

Autoactivation of yeast containing bait was tested using selective medium -His -leu and supplied with 2, 5, 10, 25, 50, 75, and 100 mM 3AT (3-Amino-1,2,4-Triazol, Fluka).

The bait and prey constructs were transformed into the yeast strains Pj694a and Pj694a, respectively. Mating and selection for interactions were performed as described in James et al. (1996).  $\beta$ -Gal activity was measured using CPRG as substrate according to the manufacturer's protocol (Clontech).

### Bimolecular fluorescence complementation assay

For "Split-YFP" analysis (Walter et al. 2004), the coding sequence of Enhanced Yellow Fluorescence Protein (EYFP, Clontech Laboratories) was fragmented into an N-terminal domain of 154 amino acids (YFP/N) and a C-terminal domain of 84 amino acids (YFP/C). Two sets of vectors were generated: one for C-terminal fusions of the two YFP fragments to proteins of interest (pARC235 and pARC236), and one for N-terminal fusions (pARC233 and pARC234). The YFP/N- and YFP/C-encoding fragments were generated by PCR and cloned as XbaI-BamHI fragments into pGD120, which drives expression from the CaMV35S promoter and is suitable for transient assays (Nougalli-Tonaco et al. 2006). An Ochre stop codon was introduced after codon 154 of YFP/N for the pARC233 C-terminal fusion construct. For the pARC236 N-terminal fusion construct, the stop codon of YFP/C was left out. The four pGD120-derived vectors were made Gateway compatible by introducing the RFB Gateway cassette (Invitrogen) into the BamHI-digested and -blunted vector (N-terminal fusion constructs) or XbaI-digested and -blunted vectors (C-terminal fusion constructs). This resulted in pARC233-pARC236. *JKD*, *MGP*, *SCR*, and *SHR* cDNAs were inserted to generate N-terminal YFP fusions (pARC233 and pARC234) and C-terminal fusions (pARC235 and pARC236). YFP fluorescence after bombardment of onion epidermal cells was recorded using a Leica SP2 CLSM.



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## References

- Aida, M., Beis, D., Heidstra, R., Willemsen, V., Blilou, I., Galinha, C., Nusse, L., Noh, Y.-S., Amasino, R., and Scheres, B. 2004. The *PLETHORA* genes mediate patterning of the *Arabidopsis* root stem cell niche. *Cell* **119**: 109–120.
- Barton, M.K. 2001. Giving meaning to movement. *Cell* **107**: 129–132.
- Bechtold, N., Ellis, J., and Pelletier, G. 1993. In planta *Agrobacterium* mediated gene transfer by infiltration of adult *Arabidopsis thaliana* plants. *Compt. Rendus Acad. Sci. III Sci. Vie* **316**: 1194–1199.
- Benfey, P.N., Linstead, P.J., Roberts, K., Schiefelbein, J.W., Hauser, M.T., and Aeschbacher, R.A. 1993. Root development in *Arabidopsis*: Four mutants with dramatically altered root morphogenesis. *Development* **119**: 57–70.
- Birnbaum, K., Shasha, D.E., Wang, J.Y., Jung, J.W., Lambert, G.M., Galbraith, D.W., and Benfey, P.N. 2003. A gene expression map of the *Arabidopsis* root. *Science* **302**: 1956–1960.
- Chen, Y. and Struhl, G.I. 1996. Dual roles for patched in sequestering and transducing Hedgehog. *Cell* **87**: 553–563.
- Colasanti, J., Yuan, Z., and Sundaresan, V. 1998. The indeterminate gene encodes a zinc finger protein and regulates a leaf generated signal required for the transition to flowering in maize. *Cell* **93**: 593–603.
- Cole, M., Nolte, C., and Werr, W. 2006. Nuclear import of the transcription factor SHOOT MERISTEMLESS depends on heterodimerization with BLH proteins expressed in discrete sub-domains of the shoot apical meristem of *Arabidopsis thaliana*. *Nucleic Acids Res.* **34**: 1281–1292.
- Cui, H., Levesque, M.P., Vernoux, T., Jung, J.W., Paquette, A.J., Gallagher, K., Wang, J.Y., Blilou, I., Scheres, B., and Benfey, P.N. 2007. An evolutionarily conserved mechanism delimiting SHR movement defines a single layer of endodermis in plants. *Science* **316**: 421–425.
- Di Laurenzio, L., Wysocka-Diller, J., Malamy, J.E., Pysh, L., Helariutta, Y., Freshour, G., Hahn, M.G., Feldmann, K.A., and Benfey, P.N. 1996. The SCARECROW gene regulates an asymmetric cell division that is essential for generating the radial organization of the *Arabidopsis* root. *Cell* **86**: 423–433.
- Fukaki, H., Wysocka-Diller, J., Kato, T., Fujisawa, H., Benfey, P.N., and Tasaka, M. 1998. Genetic evidence that the endodermis is essential for shoot gravitropism in *Arabidopsis thaliana*. *Plant J.* **14**: 425–430.
- Heidstra, R., Welch, D., and Scheres, B. 2004. Mosaic analyses using marked activation and deletion clones dissect *Arabidopsis* SCARECROW action in asymmetric cell division. *Genes & Dev.* **18**: 1964–1969.
- Helariutta, Y., Fukaki, H., Wysocka-Diller, J., Nakajima, K., Jung, J., Sena, G., Hauser, M.T., and Benfey, P.N. 2000. The SHORT-ROOT gene controls radial patterning of the *Arabidopsis* root through radial signaling. *Cell* **101**: 555–567.
- Hellens, R.P., Edwards, E.A., Leyland, N.R., Bean, S., and Mulineaux, P.M. 2000. pGreen: A versatile and flexible binary Ti vector for *Agrobacterium*-mediated plant transformation. *Plant Mol. Biol.* **42**: 819–832.
- Hejátko, J., Blilou, I., Brewer, P., Friml, J., Scheres, B., and Benková, E. 2006. In situ hybridization technique for mRNA detection in whole mount *Arabidopsis* samples. *Nat. Protoc.* **1**: 1939–1946.
- James, P., Halladay, J., and Craig, E.A. 1996. Genomic libraries and a host strain designed for highly efficient two-hybrid selection in yeast. *Genetics* **144**: 1425–1436.
- Kozaki, A., Hake, S., and Colasanti, J. 2004. The maize ID1 flowering time regulator is a zinc finger protein with novel DNA binding properties. *Nucleic Acids Res.* **32**: 1710–1720.
- Levesque, M., Vernoux, T., Cui, H., Wang, J.Y., Blilou, I., Hassan, H., Nakajima, K., Matsumoto, N., Scheres, B., and Benfey, P. 2006. A whole-genome meta-analysis predicts the SHORT-ROOT developmental subnetwork in *Arabidopsis*. *PLoS Biol.* **4**: e143. doi: 10.1371/journal.pbio.0040143.
- Morrison, J.F. and Markham, A.F. 1995. PCR-based approaches to human genome mapping. In *PCR2—A practical approach* (eds. M.J. McPherson et al.), pp. 165–196. Oxford University Press, Oxford, UK.
- Nakajima, K., Sena, G., Nawy, T., and Benfey, P.N. 2001. Inter-cellular movement of the putative transcription factor SHR in root patterning. *Nature* **413**: 307–311.
- Nougalli Tonaco, I.A., Borst, J.W., de Vries, S.C., Angenent, G.C., and Immink, R.G.H. 2006. In-vivo imaging of MADS box transcription factor interactions. *J. Exp. Bot.* **57**: 33–42.
- Nybakken, K. and Perrimon, N. 2002. Hedgehog signal transduction: Recent findings. *Curr. Opin. Genet. Dev.* **12**: 503–511.
- Sabatini, S., Beis, D., Wolkenfelt, H., Murfett, J., Guilfoyle, T., Malamy, J., Benfey, P., Leyser, O., Bechtold, N., Weisbeek, P., et al. 1999. An auxin-dependent distal organizer of pattern and polarity in the *Arabidopsis* root. *Cell* **99**: 463–472.
- Sabatini, S., Heidstra, R., Wildwater, M., and Scheres, B. 2003. SCARECROW is involved in positioning the stem cell niche in the *Arabidopsis* root meristem. *Genes & Dev.* **17**: 354–358.
- Scheres, B., Di Laurenzio, L., Willemsen, V., Hauser, M.T., Janmaat, K., Weisbeek, P., and Benfey, P.N. 1995. Mutations affecting the radial organization of the *Arabidopsis* root display specific defects throughout the embryonic axis. *Development* **121**: 53–62.
- Sena, G., Jung, J.W., and Benfey, P.N. 2004. A broad competence to respond to SHORT ROOT revealed by tissue-specific ectopic expression. *Development* **131**: 2817–2826.
- Walter, M., Chaban, C., Schütze, K., Batistic, O., Weckermann, K., Näke, C., Blazevic, D., Grefen, C., Schumacher, K., Oecking, C., et al. 2004. Visualization of protein interactions in living plant cells using bimolecular fluorescence complementation. *Plant J.* **40**: 428–438.
- Wesley, S.V., Helliwell, C.A., Smith, N.A., Wang, M.B., Rouse, D.T., Liu, Q., Gooding, P.S., Singh, S.P., Abbott, D., Stoutjesdijk, P.A., et al. 2001. Construct design for efficient, effective and high-throughput gene silencing in plants. *Plant J.* **27**: 581–590.
- Willemsen, V., Wolkenfelt, H., de Vrieze, G., Weisbeek, P., and Scheres, B. 1998. The HOBBIT gene is required for formation of the root meristem in the *Arabidopsis* embryo. *Development* **125**: 521–531.
- Wu, X., Weigel, D., and Wigge, P.A. 2002. Signaling in plants by intercellular RNA and protein movement. *Genes & Dev.* **16**: 151–158.