Brassinosteroid gene regulatory networks at cellular resolution

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Brassinosteroids (BRs) are plant steroid hormones that regulate diverse processes such as cell division and cell elongation. BRs control thousands of genes through gene regulatory networks (GRNs) that vary in space and time. We used time series single-cell RNA-sequencing to identify BR-responsive gene expression specific to different cell types and developmental stages of the Arabidopsis root, uncovering the elongating cortex as a site where BRs trigger a shift from proliferation to elongation associated with increased expression of cell-wall-related genes. Our analysis revealed HAT7 and GTL1 as BR-responsive transcription factors that regulate cell elongation in the cortex. These results establish the cortex as an important site for BR-mediated gene expression and unveil a BR signaling network regulating the transition from proliferation to elongation, illuminating new aspects of spatiotemporal hormone response.

Introduction

During development, cells pass through different states as they acquire identities and progress towards end-stage differentiation (Pierre-Jerome et al., 2018). Gene regulatory networks (GRNs) control this progression and must be tuned according to developmental stage, cell identity, and environmental conditions (Levine and Davidson, 2005; Moreno-Risueno et al., 2010; Shahan et al., 2021). Signaling molecules such as hormones are central players in coordinating these networks, but it has been challenging to disentangle how cell identities, developmental states, and hormone responses influence one another. Recent technological advances in single-cell RNA-sequencing (scRNA-seq) (Seyffarth et al., 2021; Shahan et al., 2021) and tissue-specific gene manipulations (Decaestecker et al., 2019; Wang et al., 2020) make it possible to address this challenge.

Brassinosteroids (BRs) are a group of plant steroid hormones that affect several aspects of development including cell division, cell elongation and differentiation (Clouse et al., 1996; Li et al., 1996; Szekeres et al., 1996). BRs are sensed at the plasma membrane by BR1 family receptors and BAK1/SERK co-receptors (Caño-Delgado et al., 2004; Kinoshita et al., 2005; Li and Chory, 1997; Li et al., 2002), initiating a series of signal transduction events that activate BES1 and BZR1 family transcription factors in the nucleus (Wang et al., 2002; Yin et al., 2002, 2005). BES1 and BZR1 direct BR-responsive GRNs by interacting with other transcription factors to control thousands of genes (Clark et al., 2021; Nolan et al., 2017, 2020; Sun et al., 2010; Yu et al., 2011). This downstream GRN is typically represented singularly without consideration of cell specificity (Clark et al., 2021; Guo et al., 2013; Seyed Rahmani et al., 2021; Yu et al., 2011), but BRs lead to different responses depending on cell type and developmental stage, suggesting that multiple BR GRN configurations exist (Ackerman-Lavert et al., 2021; Fridman et al., 2014; Vilarrasa-Blasi et al., 2014).

The Arabidopsis root is well suited to investigate spatiotemporal BR responses (Jaillais and Vert, 2016). Cell types are found on the radial axis of the root and each cell file forms a developmental timeline, with stem cells in the meristem at the tip of the root and more mature cells towards the shoot in the elongation and maturation zones (Dolan et al., 1993). Loss-of-function mutants in the BR pathway have short roots due to impaired cell division and reduced cell elongation (González-García et al., 2011; Kang et al., 2017; Vukašinović et al., 2021), suggesting that BRs play crucial roles in both the meristem and elongation zone. The optimal concentration of BRs varies between these developmental zones (Vukašinović et al., 2021). Relatively low levels of BRs in the meristem are required for cell cycle progression and orientation of cell division planes, but balanced signaling is important as excess BRs result in exit from the cell cycle, premature elongation, differentiation, and exhaustion of the meristem (González-García et al., 2011; Graeff et al., 2021; Li et al., 2021). BR biosynthesis increases along the longitudinal axis of the root, peaking in the elongation zone, where BRs promote cell elongation (Chaiwanon and Wang, 2015; Vukašinović et al., 2021).

BR signaling also varies among cell types. The epidermis is a major site for BR-induced gene expression that controls...
Fig. 1. scRNA-seq identifies the elongating cortex as a site of BR-response.

(A,B) Two-dimensional uniform manifold approximation and projection (UMAP) embedding of 21,473 BRZ and 22,275 two-hour BL treated cells across 3 biological replicates of scRNA-seq. Colors indicate (A) cell type or (B) developmental stage annotation.

(C) Spatiotemporal response to 2 hour BL treatment among each combination of cell type and developmental stage of the root. Color on the UMAP projection indicates the number of differentially expressed genes (DEGs).

(D) Volcano plot of DEGs in the elongating cortex. Color indicates the direction of regulation. Known markers of BR response including DWF4, RD26, XTH4 and IAA19 are indicated. C/VIF2 and CSI1 (described in this study) are also indicated.

(E,F) UMAP projection colored by BL upregulated DEGs (E) or BL down-regulated DEGs (F).

(G) Expression of C/VIF2 in BRZ and BL scRNA-seq. The color scale represents log normalized, corrected UMI counts.

(H) pC/VIF2-H2B-Venus reporter grown on 1 µM BRZ for 7 days and transferred to 1 µM BRZ or 100 nM BL for 4 hours. Inset shows C/VIF2 signals in the elongating cortex that increase with BL treatment. Propidium iodide-staining is shown in grey, with the color gradient indicating relative C/VIF2-H2B-Venus levels. Scale bars, 100 µm.

BL, brassinolide; BRZ, brassinazole.

meristem size (Ackerman-Lavert et al., 2021; Belkhadir and Jaillais, 2015; Chaiwanon and Wang, 2015; Fridman et al., 2014, 2021; González-García et al., 2011; Hacham et al., 2011; Jaillais and Vert, 2016; Nolan et al., 2020; Planas-Riverola et al., 2019). On the other hand, BR-regulated genes in the stele tend to be repressed and are implicated in differentiation and radial expansion (Fridman et al., 2021; Kang et al., 2017; Vragović et al., 2015). Additionally, BRs promote divisions of quiescent center cells via interactions of BES1 with BRAVO, a quiescent-center-enriched transcription factor (Betegón-Putze et al., 2021; Lozano-Elena et al., 2018; Vilarrasa-Blasi et al., 2014). This suggests that BR GRNs involve context-specific transcription factors, but BR responses have so far been characterized in only a handful of cell types. Since BR responses vary according to both cell type and developmental stage, increased resolution is needed to understand how spatiotemporal BR GRNs are wired.

scRNA-seq can provide such resolution and is a powerful approach to investigate cell- and developmental-stage-specific responses (Seyfferth et al., 2021; Shahan et al., 2021). Droplet-based scRNA-seq enables profiling thousands of Arabidopsis root cells (Denyer et al., 2019; Jean-Baptiste et al., 2019; Ryu et al., 2019; Shulse et al., 2019; Wendrich et al., 2020; Zhang et al., 2019). Using this technology, we constructed a single-cell reference atlas that captures the major cell types and developmental stages of the
Arabidopsis root (Shahan et al., 2022). This atlas allows the investigation of fine-scale transitions at a resolution beyond the morphologically defined meristem, elongation, and maturation zones and facilitates the analysis of additional single-cell experiments through annotation label transfer (Shahan et al., 2022; Stuart et al., 2019). Although scRNA-seq has been used to profile mutants (Denyer et al., 2019; Ryu et al., 2019; Shahan et al., 2022) and investigate plant responses to the environment (Jean-Baptiste et al., 2019; Shulse et al., 2019), previous studies have profiled a single time point and have not explored the GRNs that mediate stimuli-specific responses.

In this study, we used scRNA-seq to profile BR responses across the majority of cell types and developmental stages of the root. We discovered that BRs strongly affect gene expression in the elongating cortex. A BR scRNA-seq time course and reconstruction of cortex trajectories showed that BRs trigger a shift from proliferation to elongation, which is associated with up-regulation of cell wall-related genes. Accordingly, loss of BR signaling in the cortex had little effect on meristem cell length but reduced cell expansion in the elongation zone. Our time-course data allowed us to infer BR-responsive GRNs across cell types, developmental stages, and time points, which led to the identification of HAT7 and GTL1 as validated regulators of BR response in the elongating cortex. These datasets represent more than 180,000 single-cell transcriptomes, providing a view of BR-mediated GRNs at unprecedented resolution.

Results

Reference-guided scRNA-seq reveals differential brassinosteroid response in the Arabidopsis root

To investigate spatiotemporal BR responses in the root, we used a sensitized system. We first inhibited endogenous BR biosynthesis using Brassinazole (BRZ) (Asami et al., 2000) and then reactivated signaling by treating with Brassinolide (BL), the most active BR (Clark et al., 2021; Grove et al., 1979; Nolan et al., 2020). We treated 7-day-old primary roots for 2 hours with BL or a corresponding mock BRZ control and performed scRNA-seq on protoplasts isolated from 0.5 cm root tips (containing meristem, elongation and early differentiation zones) using the 10X Genomics Chromium system (Methods). Employing Cell preprOcessing Pipeline kaLisTo busTools (COPilot) (Bray et al., 2016; Melsted et al., 2019; Shahan et al., 2022), which streamlines scRNA-seq data processing and quality control, we identified over 43,000 high-quality cells with approximately equal numbers for each treatment (Figures 1A-B and Data S1).

To annotate the cell types and developmental stages, we performed label transfer implemented in Seurat (Stuart et al., 2019) based on our single-cell expression atlas of the Arabidopsis root (Shahan et al., 2022). To follow the developmental progression from the meristem to the elongation zone more closely, we distinguished between two domains of the meristem: the proliferation domain, where cells have a high probability to divide, and the transition domain, where cells divide less frequently but have not yet begun rapid cell expansion (Figures S1A-D and Data S2) (Ivanov and Dubrovsky, 2013; Salvi et al., 2020).

After data integration, the 11 major cell types and eight developmental stages identified were logically arranged in 2D uniform manifold approximation and projection (UMAP) space as we and others have previously described for root datasets (Figures 1A and 1B; (Denyer et al., 2019; Shahan et al., 2022). More specifically, young cells in the proliferation domain group together, indicating their transcriptional similarity. As cells develop, they become transcriptionally distinct and separate in UMAP space, forming four major branches corresponding to the stele, the ground tissue which includes cortex and endodermis, the epidermis and root cap (Dolan et al., 1993; Shahan et al., 2022). Marker genes characteristic of cell types and developmental stages remained enriched as expected, suggesting that although BRs alter the expression of thousands of genes, cell identities can be successfully aligned through integration (Figure S2A and S2B).

scRNA-seq captures spatiotemporal patterns of BR-responsive gene expression

Previous studies have profiled BR-responsive gene expression in bulk tissue or in a handful of cell types, conflating cell type and developmental stage (Chaiwanon and Wang, 2015; Guo et al., 2013; Vragović et al., 2015). To obtain better spatiotemporal resolution of the BR response, we performed differential expression analysis for each combination of cell type and developmental stage using pseudobulk expression profiles (Crowell et al., 2020; see methods). We identified over 8,000 differentially expressed genes (DEGs; Fold-change >1.5, False discovery rate <0.05; Figures 1C-F), which were enriched in BES1 and BZR1 targets and had significant overlap with previously identified BR-regulated genes (Figures S2C, S2D and Data S3).

Strikingly, we found that 37% of DEGs were significantly altered in a single cell type/developmental stage and more than 82% were differentially expressed in 5 or fewer cell type/developmental stage combinations (Figure S2E). This indicates that although BRs broadly influence gene expression, they modulate distinct sets of genes in different spatiotemporal contexts.

Among the tissues with many DEGs was the epidermis, as previously described (Chaiwanon and Wang, 2015; Fridman et al., 2014; González-García et al., 2011; Hacham et al., 2011). Atrichoblasts, or non-hair cells in the epidermis were particularly affected, showing marked changes across both the meristem and elongation zone. Unexpectedly, our data also indicated that BRs strongly influence gene expression in the cortex, especially in the elongation zone (Figures 1C and S2D). The cortex has been linked to plant environmental interactions, including response to water limitation (Longkumer et al., 2021; Verslues and Longkumer, 2022) and hydrotropism (Dietrich et al., 2017; Miao et al., 2021; Takahashi et al., 2002), but how BRs modulate gene expression in the cortex and what processes are affected have not been described. To address these questions, we focused
Fig. 2. Waddington optimal transport traces cortex trajectories along a BR scRNA-seq time series.

(A) UMAP of 52,921 cells across 6 time points of a scRNA-seq BL treatment time course. Mock BRZ control represents time 0. Colors indicate cell type annotation.

(B) UMAP projection colored by developmental stage annotation.

(C) Density plot showing cell wall gene expression score. The shaded region with cell wall expression scores >1 indicates “Cell Wall+” cells.

(D) Bar plot showing the percentage of Cell Wall+ cells in the cortex versus other cell types over the time course. Color indicates developmental stage annotation. Only transition and elongation zones are shown as other zones represent less than 2% of cell wall + cells.

(E) WaddingtonOT (WOT) fate probabilities for all cortex cells along the BL time course. The BL 2 hour time point was used as a reference, therefore all cells have a probability of either 1 or 0 at this time point.

(F) Probabilities for cortex cell wall+ state from WOT trajectories.

(G) Expression trends for select transcription factors differentially expressed along WOT cortex cell wall+ trajectories.

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on BR-mediated gene expression in the elongating cortex.

**Cell wall-related genes are up-regulated by BRs in the elongating cortex**

We found that BR treatment led to approximately 1,000 up-regulated genes and about the same number of down-regulated genes in the elongating cortex (Figures 1D and S2D). Gene ontology (GO) analysis indicated the BR up-regulated genes were strongly enriched for genes related to “cell wall organization or biogenesis”, which is intriguing given the role of BRs in promoting cell elongation (Figure S2F). The cell wall-related DEGs included **CELLULOSE SYNTHASES (CESAs)**, **CELLULOSE SYNTHASE INTERACTIVE1 (CSI1)**, which is required for efficient cellulose synthesis and alignment of cellulose synthase complexes with cortical microtubules (Bringmann et al., 2012; Gu et al., 2010; Li et al., 2012) and cell-wall loosening enzymes such as **EXPANSINS** and **XYLOGLUCAN ENDOTRANSGLUCOSYLASES**. Cell-wall-related genes such as CESAs have been demonstrated to be direct targets of BES1 and BZR1 (Sun et al., 2010; Xie et al., 2011; Yu et al., 2011), but their spatiotemporal regulation, especially in the cortex, has not been reported.

To monitor their responsiveness to BRs, we generated transcriptional reporters for three of the DEGs: **C/VIF2**, **CSI1** and **XTH16**. **CELL WALL / VACUOLAR INHIBITOR OF FRUCTOSIDASE 2 (C/VIF2)** was enriched in the transition domain and elongation zone of the cortex and induced by BL (Figures 1G-H). **CSI1** was broadly induced by BL, especially in the epidermis and cortex (Figures S2G and S2H). **XYLOGLUCAN ENDOTRANSGLUCOSYLASE/HYDROLASE 16 (XTH16)** was enriched in the endodermis and induced by BL (Figures S2I and S2J). These results confirm that our differential expression analysis captures spatiotemporal BR responses and raises the possibility that BR induction of cell-wall-related genes is associated with cortex cell elongation.

**Waddington optimal transport identifies exceptionally responsive cells in a BR scRNA-seq time course**

Analyzing expression trends over a time course can reveal regulators and downstream targets associated with gene expression programs (Schiebinger, 2021; Swift et al., 2021). A recently developed analytical approach for scRNA-seq is Waddington-OT (WOT), which connects snapshots of gene expression along a time course to facilitate trajectory reconstruction (Schiebinger et al., 2019). WOT identifies putative ancestors for a given set of cells at earlier time points and descendants at later time points (Schiebinger et al., 2019; Shahan et al., 2022; Zhang et al., 2021). To better understand how BRs influence cell wall-related gene expression we performed scRNA-seq at six time points beginning with BRZ treatment (time 0) and subsequent BL treatments for 30 minutes, 1 hour, 2 hours, 4 hours or 8 hours (Figures 2A and 2B). These time points capture the rapid root elongation triggered by re-addition of BRs (Chaiwanon and Wang, 2015).

To examine the trajectories leading to activation of cell wall-related genes in the elongating cortex, we applied WOT (Schiebinger et al., 2019) and created a cell wall gene signature using 107 cell wall-related genes that were induced by BL in the elongating cortex (Data S4). We monitored the relative expression of this set of genes, resulting in a “cell wall score” for each cell in the time course (see methods). Cortex cells had a higher cell wall score compared to other cell types, which increased with BL treatment (Figures 2C-D and S3A-B), confirming that the cell wall score represents a BR-responsive module in the cortex. At the 2 hour BL time point, more than 20% of cortex cells had a cell wall score greater than 1, whereas only 5% or fewer cells in other cell types exhibited scores this high (Figure 2C). We therefore designated cells with a cell wall score of at least 1 as “Cell wall+” to indicate their exceptional BR response (Figures 2C-D).

**BR induction of cell wall genes in the cortex is associated with the switch to elongation**

An advantage of WOT analysis is that it does not rely on pre-specified boundaries between developmental zones. We used this property to examine the relationship between developmental stage annotation and cell wall score. Under BRZ treatment, cortex cell wall+ cells were sparse and predominately annotated as transition domain. Upon BL treatment, the annotation of cortex cell wall+ cells shifted to the elongation zone (Figure 2D), suggesting that BR induction of cell wall-related genes is related to the onset of cell elongation.

Using the cells at the 2 hour time point as a reference, we looked at the probability of cells being ancestors or descendants of cortex cell wall+ cells. We also constructed a similar trajectory for the remaining cortex cells, which were designated “cortex cell wall-“. These trajectories illustrated a shift towards the cortex cell wall+ state upon BL treatment that coincided with a change from transition domain to elongation zone annotation (Figures 2D-F). These results provide support for the hypothesis that BRs are involved in initiating elongation of cortex cells via activation of cell wall genes.

**Differential expression along WOT trajectories identifies BR-responsive transcription factors**

To reveal potential regulators of cell wall-related genes in the cortex, we performed probabilistic differential expression analysis along WOT trajectories, contrasting cells assigned to cortex cell wall+ versus cortex cell wall- states at each time point (see methods). Among the DEGs identified were known transcription factors in the BR pathway including **BES1** (Yin et al., 2002), **BIM1** (Yin et al., 2005), and **IBLI** (Zhiponova et al., 2014); Figure 2G and Data S4). We also identified additional transcription factors whose role in the BR pathway has not been examined in detail (Figure 2G). These include **JACKDAW (JKD)**, which is involved in ground tissue specification (Moreno-Risueno et al., 2015), **HAI7/17HB-3**, an uncharacterized class I HD-ZIP TF (Ariel et al., 2007; Henriksson et al., 2005; Mattsson et al., 1992) and **GTL1**, which negatively regulates growth in trichomes.
Fig. 3. Triple receptor mutant bri1-T gene expression changes in cortex and distinct patterns in pGL2-BRI1-GFP/bri1-T.

(A) 7-day old WT, bri1-T and pGL2-BRI1-GFP/bri1-T roots grown under control conditions. Propidium iodide-staining is shown in grey and GFP in green. Scale bars, 100 µm.

(B-C) UMAP projection of scRNA-seq from 14,334 wild-type cells, 12,649 bri1-T cells and 7,878 pGL2-BRI1-GFP/bri1-T cells. Two biological replicates of scRNA-seq were performed for each genotype. Colors indicate cell type annotation (B) or developmental stage (C).

(D) UMAP colored by DEGs for each cell type/developmental stage combination of bri1-T compared to WT.

(E) Volcano plot of DEGs in the elongating cortex from bri1-T compared to WT. Color indicates the direction of regulation.

(F-I) UMAP colored by DEGs for the indicated comparisons.

(J) Multidimensional scaling (MDS) analysis of cortex cells from scRNA-seq. Note that replicates from the same genotype group together, but genotypes are well separated.

(K) Multidimensional scaling (MDS) analysis of atrichoblast cells from scRNA-seq.

(L-M) Gene expression trends for bri1-T vs wild-type DEGs along cortex (L) or atrichoblast (M) trajectories. Scaled expression along cortex pseudotime is plotted for each genotype. Lower bar indicates pseudotime progression calculated by CytoTRACE.
(Breuer et al., 2009, 2012) and root hairs (Shibata et al., 2018, 2021), but has not been linked to BR signaling or the cortex. Expression of HAT7 and GT1I significantly increased in the cortex of our BRZ versus BL pseudobulk differential expression analysis at the 2 hour time point. On the other hand, although JKD was detected in WOT differential expression analysis, it did not pass our criteria for statistical significance in the 2 hour pseudobulk differential expression analysis (Data S3). Using a recombineering line (Moreno-Risueno et al., 2015), we found that JKD was present in the proliferation domain regardless of BRZ or BL treatment status and showed an increase in expression in the transition and elongation zone of the cortex upon BL treatment (Figures S3C and S3D). These results indicate that WOT trajectories can identify BR-responsive transcription factors that may be involved in regulating cell wall-related genes in the cortex.

To generalize our WOT analysis we constructed trajectories for each combination of cell type and developmental stage and performed differential expression analysis between each time point (Data S5). These data represent a valuable resource to generate additional hypotheses regarding spatiotemporal BR responses.

Analysis of the triple receptor mutant bri1-T reveals changes in cortex expression

Since our results indicated that exogenous BRs lead to activation of cell wall-related genes in the elongating cortex, we asked if this is also the case for endogenous BRs. A gradient of BRs is present along the longitudinal axis of the root, with low BR levels in the proliferation domain (Vukašinović et al., 2021). BR biosynthesis increases as cells enter the transition domain and peaks in the elongation zone, shootward of which is a BR signaling maximum (Chaiwanon and Wang, 2015; Vukašinović et al., 2021). Interpretation of this endogenous BR gradient requires receptor BRI1 and its close homologs BRL1 and BRL3 (Caño-Delgado et al., 2004; Friedrichsen et al., 2000; He et al., 2000; Irani et al., 2012; Kinoshita et al., 2005).

To identify differentially expressed genes, we performed two replicates of scRNA-seq on the BR-blind bri1bri1bri3 triple mutant (bri1-T) along with paired wild-type controls (Figures 3A-C). A previous study profiled single cells from bri1-T (Graeff et al., 2021), but these data were from a single replicate and were compared to a wild-type sample from a different study (Wendrich et al., 2020). Pseudobulk differential expression identified the elongating cortex as exhibiting substantial differential gene expression (Figures 3D-G, S4A and Data S3). The genes down-regulated in the elongating cortex of bri1-T were enriched for the GO term “cell wall organization or biogenesis” (Figure S4B). These data indicate that, similar to exogenous application of BRs, endogenous BRs promote the expression of cell wall-related genes in the elongating cortex.

The epidermis is widely described as the major site for BR-promoted gene expression in the root (Chaiwanon and Wang, 2015; Fridman et al., 2014, 2021; Großeholz et al., 2021; Hacham et al., 2011; Nolan et al., 2020; Vragović et al., 2015; Wei and Li, 2016). Previous studies showed that epidermal expression of BRI1 was sufficient to rescue morphological phenotypes including meristem size and root length of loss-of-function BR mutants such as bri1-T (Fridman et al., 2014; Hacham et al., 2011; Kang et al., 2017). To determine the extent to which BR-regulated gene expression is restored, we performed scRNA-seq on pGL2-BRI1-GFP/bri1-T - a line in which BRI1 is expressed in atrichoblast cells of the epidermis of bri1-T (Hacham et al., 2011; Kang et al., 2017; Vragović et al., 2015). We identified over 8,000 DEGs in comparison with wild type (Figures 3H and S4C-D) and in comparison with bri1-T (Figures 3I and S4E-H), indicating that gene expression remains dramatically perturbed and that this is far from a complete rescue of the bri1-T phenotype.

To further examine these gene expression changes, we aggregated cell-level counts for cortex cells and separately for atrichoblast cells of each sample and performed Multidimensional scaling (MDS) analysis. Samples of the same genotype appeared grouped across replicates for both cell types, but the pGL2-BRI1-GFP/bri1-T replicates formed a distinct cluster from either wild-type or bri1-T (Figure 3J).

Together, our scRNA-seq of BR treatment and bri1-T indicate that the cortex represents a site of BR-mediated gene expression. The observation that pGL2-BRI1-GFP/bri1-T has a distinct pattern of gene expression from WT or bri1-T suggests that alternative means of probing tissue-specific BR responses could be informative.

Tissue-specific CRISPR of BRI1 confirms a role for the cortex in BR-mediated cell expansion

To selectively block BR signaling in cell types of interest we performed tissue-specific CRISPR (Decaestecker et al., 2019) of BRI1. We used a bri1 mutant complemented with pBRI1-BRI1-mCitrine (Figure 4A) into which we introduced Cas9 driven by tissue-specific promoters to knock out BRI1 either in the epidermis and lateral root cap (pWER-BRI1-CRISPR) or in the cortex (pCO2-BRI1-CRISPR). mCitrine signals were absent in the expected locations of the tissue-specific CRISPR lines, confirming their efficacy and specificity (Figures 4A-C and S5A-C).

Since our scRNA-seq data indicated that exogenous BRs promote the expression of cell wall-related genes in the elongating cortex, we hypothesized that loss of BR signaling in the cortex would affect final cell size. Indeed, pCO2-BRI1-CRISPR lines displayed significantly shorter mature cortex cells, while meristematic cortex cell length was relatively unaffected (Figures 4C-E).

In contrast, epidermal knockout of BRI1 in pWER-BRI1-CRISPR lines resulted in both reduced meristem cell size and reduced mature cortex cell length (Figures 4C-E), which is consistent with the reported role of epidermal BR signaling (Chaiwanon and Wang, 2015; Fridman et al., 2021; Hacham et al., 2011; Vragović et al., 2015). These results indicate that in addition to the epidermis, BR signaling in the cortex is required to promote cell expansion in the elongation zone.

HAT7 and GT1I are BR responsive regulators along cor-
Fig. 4. Tissue-specific CRISPR of BRI1 confirms role for cortex in BR-mediated cell expansion.

(A) Overview of BRI1 tissue-specific CRISPR approach. A bri1 mutant complemented with pBRI1-BRI1-mCitrine (1) was used as background to introduce tissue-specific Cas9 along with gRNAs targeting BRI1 (2). This allows for visualization of BRI1 knockout in specific cell layers, such as the cortex when pCO2-BRI1-CRISPR is used (3).

(B) Appearance of Cas9-tagRFP in the cortex is associated with loss of BRI1-mCitrine signal, confirming tissue-specific knockout.

(C) Confocal images of BRI1 tissue-specific CRISPR lines. Control indicates a broad expression pattern of BRI1-mCitrine in pBRI1-BRI1-mCitrine/bri1. BRI1-mCitrine signals are shown in green and propidium iodide staining (PI) in magenta (upper panels). White arrows specify tissues with absence of BRI1-mCitrine signal; epidermis for pWER-BRI1-CRISPR and cortex for pCO2-BRI1-CRISPR. Mature root sections illustrating changes in cell size and length (lower panels). Cortex cells are pseudocolored to indicate their position.

(D) Quantification of meristematic cortex cell length, defined as the first 20 cells of individual roots starting from the quiescent center. Control indicates pBRI1-BRI1-mCitrine/bri1 complemented line.

(E) Quantification of mature cortex cell length. For (D) and (E), all individual data points are plotted. Magenta horizontal bars represent the means and error bars represent s.d. Significant differences between each line and wild type were determined by one-way ANOVA and Dunnett’s multiple comparison tests. *** P<0.001, ** P<0.01 and * P<0.05. n.s. not significant. Scale bars, for (B) and (C) = 50 µm. TSKO, tissue-specific knockout.
Fig. 5. HAT7 and GTL1 are BR responsive regulators along cortex trajectories.

(A) Upset plot showing a comparison of genes up-regulated by BL in the cortex, down-regulated in the cortex of bri1-T, and differentially expressed along wild-type cortex trajectories. Red color indicates 163 genes common to all three sets.

(B) Gene expression trends for 163 core BR DEGs along cortex trajectories. Scaled expression along cortex pseudotime is plotted for each time point of the BR time series and for wild type versus bri1-T. Lower bar indicates pseudotime progression calculated by CytoTRACE.

(C-E) Gene expression trends for HAT7, GTL1 or C/VIF2 along the developmental zones of the cortex (y-axis) for each time point of the BR time course (x-axis). Color bar indicates the scaled expression level in the cortex.

(F-G) 7-day old roots expressing pHAT7-HAT7-mCitrine or pGTL1-GTL1-mCitrine reporters under the indicated treatments. Control represents a mock DMSO solvent. For BRZ and BL treatments, plants were grown on 1 µM BRZ for 7 days and transferred to 1 µM BRZ or 100nM BL for 4 hours. Propidium iodide-staining is shown in grey, with the color gradient indicating relative mCitrine levels. Scale bars, 100 µm. BL, brassinolide; BRZ, brassinazole.
tex trajectories
To define a core set of genes associated with BR response along cortex trajectories we first compared genes induced in the cortex by BL treatment with those down-regulated in the cortex of bri1-T. Of the 768 genes in common, we then asked which vary along developmental time in wild-type cortex trajectories (Shahan, Hsu et al 2022). The intersection of these three lists identified a core set of 163 BR responsive DEGs (Figure 5A and Data S3). Consistent with regulation by BRs, 69% of the core DEGs are BES1 and BZR1 direct targets from ChIP experiments (Oh et al., 2014; Sun et al., 2010; Yu et al., 2011). Expression along cortex pseudotime illustrates induction by BL treatment and down-regulation in bri1-T (Figure 5B). Additionally, HAT7 and GTL1 were induced along these trajectories, suggesting a potential role for these transcription factors in controlling BR-regulated gene expression in the cortex (Figure 5C-E and S6A).

To gain insight into their roles, we generated translational reporter lines for HAT7 and GTL1 and monitored their expression. Under control conditions, pHAT7-HAT7-mCitrine lines showed expression in the transition domain and elongation zone of the cortex (Figure 5F), consistent with previous reports (Lee et al., 2006). We also observed HAT7 signals in the epidermis and endodermis, in line with expression patterns in our wild-type scRNA-seq atlas (Shahan et al., 2022). HAT7 expression was decreased when BR biosynthesis was inhibited with BRZ, and restored upon BL treatment (Figures 5G and S6A). pGTL1-GTL1-mCitrine was more broadly expressed, with increasing levels in the cortex and epidermis as cells progress from the transition domain to the elongation zone (Figures 5G and S6A). GTL1-mCitrine expression was reduced by BRZ and increased by BL treatment (Figure 5G). These results confirm that BRs promote the expression of HAT7 and GTL1 coinciding with the onset of cell elongation. Furthermore, HAT7 and GTL1 are direct targets of BES1 and BZR1 (Oh et al., 2014; Sun et al., 2010; Yu et al., 2011), suggesting that they may be part of the BR-directed GRN activated as cells progress from proliferation to elongation.

Previous studies have inferred global (Guo et al., 2013; Seyed Rahmani et al., 2021; Sun et al., 2010; Xie et al., 2019a; Yu et al., 2011) or temporally resolved GRNs (Clark et al., 2021) for BR response, but they lack cell type and developmental-stage specificity. To infer GRN configurations across our BR time series we used CellOracle (see methods; Data S8) focusing on BL DEGs and associated transcription factors.

Analysis of network importance scores such as centrality measures is a powerful approach to prioritize candidate regulators among DEGs (Iacono et al., 2019; Kamimoto et al., 2020). Since the cell wall signature peaked at 2 hours after BL treatment, we prioritized transcription factors with high network centrality scores in the elongating cortex at this time point. HAT7 was the top-ranked transcription factor in terms of degree centrality and three close homologs: HB13, HB20 and HB23 were also among the top 10 transcription factors (Figures 6A-B and Data S8). Together HAT7, HB13, HB20 and HB23 make up the alpha clade HD-ZIP I transcription factors in Arabidopsis (Ariel et al., 2007; Henriksson et al., 2005; Mattsson et al., 1992). Genetic analysis of HAT7 has yet to be carried out, but HB13, HB20 and HB23 have been implicated in root development (Perotti et al., 2019; Silva et al., 2016), shoot development (Ribone et al., 2015), and stress responses (Ebrahimian-Motlagh et al., 2017; Gao et al., 2014; Harris et al., 2011; Perotti et al., 2021).

We used CRISPR to generate hat7 loss-of-function mutants but did not observe strong phenotypes in terms of cortex cell elongation (Figure S6D-F). Since HB13, HB20 and HB23 are induced by BRs and predicted to regulate cell wall-related genes in our GRNs (Figures 6B, S6A-B and Data S10), we next generated hat7 hb13 hb20 hb23 quadruple mutants via multiplex CRISPR (Stuttmann et al., 2021). Mature cortex cell length was reduced by approximately 25% in two independent quadruple mutants (Figures 6C-E and S6E), providing strong evidence that HAT7 and its homologs are required for cell elongation. Despite the decrease in final cell length, the root length of the quadruple mutant was not dramatically reduced (Figure SSD), suggesting that the decrease in cell length is at least partially compensated for by increased cell production.

We next investigated GTL1, which was the 5th highest ranked TF in the BL 2 hour elongating cortex GRN in terms of degree centrality (Figures 6A-B). Given that GTL1 was shown to function redundantly with its close homolog DF1 in terminating root hair growth (Shibata et al., 2018, 2021), we examined gtl1 and df1 single mutants along with gtl1 df1 double mutants. gtl1 and df1 displayed only subtle changes in mature cortex cell length. However, gtl1 df1 double mutant had significantly shorter mature cortex cell lengths and shorter roots (Figures 6C-E and S6D-F). As previously reported (Shibata et al., 2018), DF1 was expressed at lower levels as compared to GTL1, making DF1 challenging to detect in scRNA-seq (Figure S6A-B). Despite this, we observed increasing trends of DF1 expression along WOT trajectories in the BL time course, especially in cortex cell wall+ cells (Figure S6B). We also verified that BL treatment results in an increase in pDF1-DF1-GFP levels as compared to BRZ controls (Figure S6C).

Together, our genetic analysis of HAT7 and GTL1 family transcription factors illustrates the power of GRN-mediated discovery of regulatory factors in spatiotemporal BR response.

**BES1 and GTL1 physically interact and share a common set of target genes**

Since BES1 is known to interface with other transcription factors in controlling BR-regulated gene expression, we compared target genes for BES1 and BZR1 (Oh et al., 2014; Sun et al., 2010; Yu et al., 2011) to ChIP targets of GTL1 and DF1 (Shibata et al., 2018). BES1 and BZR1 share 3,020 common targets with GTL1, significantly more than expected by chance (Figure S7A, P < 0.001, Fisher’s exact test). Similarly, BES1 and BZR1 share 2,490 common targets with DF1 (Figure S7A, P < 0.001, Fisher’s exact test). When compared to BR-regulated genes from scRNA-seq, BES1 and GTL1 targets showed the strongest enrichment in genes up-
Fig. 6. HAT7 and GTL1 are top-ranked regulators in cortex GRNs and affect BR-related phenotypes.

(A) Top 10 transcription factors (TFs) in the CellOracle BL two-hour elongating cortex GRN ranked by out degree. The ranking is indicated by the number inside the circle. Color indicates TF family, with light grey corresponding to any family other than HAT7 or GTL1.

(B) Subnetwork showing cell wall-related genes that are predicted targets of HAT7 and GTL1 in the CellOracle elongating cortex GRN. HB13, HB20 and HB23 are included in the subnetwork since they are connected to HAT7 and cell-wall-related genes.

(C) Propidium iodide-staining of 7-day-old WT, hat7 hb13 hb20 hb23 (line 1-2), and gtl1 df1 roots. Insets show cortex cells entering the elongation zone. Scale bars, 100 μm.

(D) Quantification of cortex cell length along the longitudinal axis of the root. The quiescent center was designated as “0” and each cell number consecutively thereafter. The grey area represents the confidence interval of the smoothed mean estimated with a generalized additive model. Number of roots per genotype: WT=51, gtl1 df1=26, hat7 hb13 hb20 hb23=16.

(E) Quantification of mature cortex cell length. Red horizontal bars represent the means and error bars represent s.d. Significant differences between each line and wild type were determined by one-way ANOVA and Dunnett’s multiple comparison tests. ***P<0.001.
regulated by BRs in the transition domain and elongation zone of the cortex (Figure S7B), with 297 common targets of both BES1 or BZR1 and GTL1 being induced in the elongating cortex by BL treatment.

Given the strong overlap between BES1 and GTL1 targets, we hypothesized that these TFs physically interact to regulate a common set of genes. Co-immunoprecipitation showed that GTL1-FLAG pulled down BES1-GFP (Figure S7C). These results suggest that BRs induce GTL1 and subsequently BES1 and GTL1 interact to control a common set of target genes.

**scRNA-seq reveals cell-type-specific expression underlying gtl1 df1 phenotypes**

Our results indicate that *gtl1 df1* mutants have reduced cortex cell elongation. On the other hand, *gtl1 df1* mutants have longer trichoblasts or root hairs (Shibata et al., 2018). A downstream regulatory network that enables GTL1-mediated growth inhibition has been dissected in trichoblasts (Shibata et al., 2018). To identify the cell-type-specific changes in gene expression underlying *gtl1 df1* cortex phenotypes we performed scRNA-seq on *gtl1* and *df1* single mutants, and on the *gtl1 df1* double mutant. Using pseudobulk differential expression analysis, we detected relatively subtle changes in *gtl1* or *df1* single mutants compared to wild type (Figures 7A-B). In contrast, over 8,000 genes were differentially expressed in *gtl1 df1* double mutants versus wildtype (Figure 7C).

Over 1,000 genes were up-regulated across all developmental stages of the cortex of *gtl1 df1*, and an approximately equal number of genes were down-regulated. The majority of cortex DEGs were affected in the elongation zone (Figures 7D-E, S7D and Data S3). Of the down-regulated genes in the cortex of the double mutant, 226 genes were also up-regulated by BL treatment. Furthermore, 31.3% of the core BR DEGs were down-regulated in the cortex of *gtl1 df1*, whereas only 6.8% were up-regulated (Data S3). The larger proportion of genes up-regulated by BRs but down-regulated in the cortex of *gtl1 df1* suggests that GTL1 and DF1 promote the expression of a subset of BR-induced genes in the cortex.

Plotting *gtl1 df1* DEGs along cortex pseudotime illustrated the down-regulation of several genes involved in cell elongation including CESA5 and AHA2 (Figure 7E). These genes were significantly enriched for the GO term “cell wall organization or biogenesis” (Figure S7E). We next examined *C/VIF2*, because it is induced by BL in the cortex (Figures 1G-H), but its expression decreased in cortex cells of *gtl1 df1* (Figures 7F-G). A p*C/VIF2*-H2B-Venus reporter showed expression of *C/VIF2* in the transition and elongation zone of the wild-type cortex, whereas its expression was reduced in the cortex of *gtl1 df1* mutants (Figure 7H and Video S1).

Taken together, our results establish the elongating cortex as a site of BR-regulated expression associated with cell elongation. Reconstruction of cortex trajectories revealed BR induction of cell wall-related genes that coincide with the switch to elongation and identified HAT7 and GTL1 as BR-responsive regulators of cortex GRNs. The reduced expression of cell wall-related genes in *gtl1 df1* mutants validates our cell-type-specific BR GRNs and identifies a function of GTL1 in promoting cortex cell elongation in response to BRs.

**Discussion**

Understanding how hormone-mediated GRNs are controlled in space and time has the potential to enable the engineering of specific downstream responses to optimize plant growth under a changing environment (Fábregas et al., 2018; Gupta et al., 2020; Nolan et al., 2020). Plant hormones including BRs, auxin, gibberellins, and abscisic acid have been shown to exhibit tissue-specific responses (Ackerman-Lavert and Savaldi-Goldstein, 2020; Bargmann et al., 2013; Geng et al., 2013; Iyer-Pascuzzi et al., 2011; Shani et al., 2013; Ubeda-Tomás et al., 2008, 2009), but how the associated GRNs are modulated in different cell types at particular developmental stages is largely enigmatic. In this study, we profiled BR-responses across cell types, developmental stages and time points of treatment using scRNA-seq, providing a high-resolution map of signaling outputs. We identified the elongating cortex as a spatiotemporal context for BR signaling, where BRs activate cell wall-related genes and promote elongation. We further showed that HAT7 and GTL1 are BR-induced regulators along cortex trajectories that control cell elongation. Our findings reveal spatiotemporal BR responses and the underlying GRNs at unprecedented resolution.

BR signaling is arguably one of the best-characterized signaling pathways in plants (Nolan et al., 2020; Planas-Riverola et al., 2019) and there is considerable evidence for tissue-specific BR responses, especially in roots (Ackerman-Lavert and Savaldi-Goldstein, 2020; Fridman et al., 2021; Graeff et al., 2021; Jaillais and Vert, 2016; Vukašinović et al., 2021). Despite these efforts, characterization of cell-type-specific BR signaling has relied on tissue-specific complementation lines, which have led to conflicting results and have not examined the role of BR signaling in the cortex (Chaiwanon and Wang, 2015; Fridman et al., 2014; Graeff et al., 2020, 2021; Hacham et al., 2011; Kang et al., 2017; Pavelescu et al., 2018; Vragović et al., 2015).

Epidermal expression of the BR receptor BRI1, as well as downstream transcription factor BZR1, have been shown to partially rescue root morphology defects of BR signaling mutants (Chaiwanon and Wang, 2015; Fridman et al., 2014; Graeff et al., 2020, 2021; Hacham et al., 2011; Kang et al., 2017; Pavelescu et al., 2018; Vragović et al., 2015).
Fig. 7. scRNA-seq reveals cell-type-specific expression underlying gtl1 df1 phenotypes

(A-C) UMAP projection of scRNA-seq from 74,810 WT, gtl1, df1, and gtl1 df1 cells. Two biological replicates were profiled for each genotype. Color indicates DEGs for each cell type/developmental stage combination of gtl1 compared to WT (A), df1 compared to WT (B) or gtl1 df1 compared to WT (C).

(D) Volcano plot of DEGs in the elongating cortex from gtl1 df1 compared to WT. Color indicates the direction of regulation.

(E) Gene expression trends along cortex trajectories for DEGs in gtl1 df1 compared to WT. Each row represents the scaled expression of a gene along cortex pseudotime. The lower bar indicates pseudotime progression calculated by CytoTRACE.

(F) Expression of C/VIF2 in wild type and gtl1 df1 scRNA-seq. The color scale represents log normalized, corrected UMI counts.

(G) Gene expression trends plotted along developmental zones of the cortex (y-axis) for WT, gtl1, df1, and gtl1 df1. The color bar indicates the scaled expression level.

(H) 7-day old root images of a pC/VIF2-H2B-Venus reporter in wild type or gtl1 df1 under control conditions. Propidium iodide-staining is shown in grey, with the color gradient indicating relative mCitrine levels. Scale bars, 100 µm.
sect cell-type-specific BR phenotypes.

Using tissue-specific CRISPR we found that the phenotype associated with loss of BRI1 depends on the cell type. Tissue-specific knockout of BRI1 in the cortex reduced final cell size, but did not markedly influence the meristem. This is consistent with the idea that BRs induce the expression of cell wall-related genes in the elongating cortex to promote cell expansion. In contrast, scRNA-seq showed that BRs affect gene expression more broadly across the developmental zones of the epidermis. Accordingly, we observed both meristem cell length and final cell size were reduced when BRI1 was knocked out in the epidermis and LRC. It has been suggested that the cortex could instruct anisotropic growth through its physical connection with the epidermis, but as the outermost tissue, relaxation of the epidermis is required to allow for cell elongation (Baskin and Jensen; Bou Daher et al., 2018). This may explain the apparent widening of cortex cells in pWER-BRI1-CRISPR lines. It is plausible that continued BR signaling in the cortex promotes expansion which is physically limited by the stiff outer epidermal cells that lack BR signaling in pWER-BRI1-CRISPR roots. Future studies could examine how cell elongation is coordinated between these neighboring tissues in the BRI1 tissue-specific CRISPR lines.

Notably, BRI1 driven by its native promoter was still present in the stele of our tissue-specific CRISPR lines when we observed these phenotypic defects, suggesting that, unlike pCVP2-BRI1, native expression of BRI1 in the stele is not sufficient for BR-induced cell elongation and root growth. These results confirm the role of the epidermis in BR-regulated root growth and reveal the function of cortex in BR-mediated cell expansion, demonstrating how scRNA-seq can identify a new spatiotemporal context for hormone signaling.

Our findings also highlight the ability of single-cell genomics to identify context-specific transcription factors, a capability that could be leveraged to precisely engineer plant growth, development, and/or responses to stress. Multiple lines of evidence suggest that HAT7 and GTL1 are BR-responsive regulators along cortex trajectories. HAT7 and GTL1 expression increased in exceptionally responsive cortex cell wall + cells along WOT trajectories. This suggests that HAT7 and GTL1 are associated with high relative expression of BR-induced cell wall-related genes and promote the shift from transition domain to elongation zone. We also found that HAT7 and GTL1 were among a core set of BR-responsive genes along cortex trajectories and are induced by BR treatment but decreased in bri1-1. Examination of HAT7 and GTL1 reporters supported these observations, showing that HAT7 and GTL1 are induced by BRs and increase in expression as cortex cells elongate. HAT7, GTL1, and several close homologs were among the top 10 ranked transcription factors in elongating cortex GRNs. We showed that hat7 hb13 hb20 hb23 quadruple mutants and gtl1 df1 double mutants have reduced mature cortex cell lengths. However, neither of these multiple mutants had a phenotype as severe as complete loss-of-function mutants in the pathway such as bri1-1. Considering that most gene promoters are controlled by multiple transcription factors, it’s likely that additional transcription factors cooperate with GTL1 and/or HAT7 in the regulation of genes involved in cell elongation in the cortex. Advances such as combinatorial CRISPR screens (Gaillochet et al., 2020) pave the way to investigate potential genetic redundancy between HAT7, GTL1, and other regulatory factors identified in our cortex GRNs.

A common theme in BR signaling networks is that BES1 and BZR1 bind to the promoters of downstream transcription factors to control their expression and then subsequently physically interact with these transcription factors to co-regulate target genes (Guo et al., 2013; Li et al., 2009; Nolan et al., 2020; Ye et al., 2017). This type of feed-forward loop could provide a mechanism to amplify the BR signal and/or to direct BES1, a more broadly expressed transcription factor, to drive tissue-specific gene expression by interacting with other more specifically expressed transcription factors. Our findings reinforce this theme and also add spatiotemporal context to such networks. GTL1 is a direct ChIP target of BES1 and BZR1 (Oh et al., 2014; Sun et al., 2010; Yu et al., 2011) and is induced by BRs. BES1 and GTL1 physically interact and share over three thousand common target genes, with the strongest enrichment among genes up-regulated by BRs in the transition and elongation zones of the cortex. Our scRNA-seq analysis showed a corresponding reduction of BR-induced genes in the cortex of gtl1 df1, supporting the hypothesis that GTL1 is required for BR-regulated gene expression and cortex cell elongation.

Although we focused on BR control of cell elongation in the cortex in this study, many additional aspects of spatiotemporal BR responses remain to be explored. Our data represent a rich resource to investigate BR-induced changes in cell identity as has been shown for trichoblasts versus atrioblasts in the epidermis (Cheng et al., 2014; Kuppusamy et al., 2009). Comparison of BR treatment versus mutant datasets could provide insight into the maintenance of procambium identity in the stele, which requires BRI1 in a BR-independent manner (Holzwarth et al., 2018). Our BR GRNs could also be used to dissect the transcriptional responses in other contexts such as the meristem, where a dual function of BRs in promoting auxin levels but inhibiting auxin signaling operates through unknown regulators in the epidermis (Ackerman-Lavert et al., 2021). To facilitate such discoveries, we have made our data publicly available as an interactive browser https://shiny.mdc-berlin.de/ARVEX/ which allows users to query gene expression patterns, DEG tables, WOT results and CellOracle GRNs.

In summary, our scRNA-seq analysis and tissue-specific gene manipulations demonstrated that the cortex is involved in BR-regulation of cell wall-related genes and cell elongation. We further identified HAT7, GTL1 and their homologs as critical regulators that cooperate with BES1 to drive cortex-specific gene expression and the transition to cell elongation in response to BRs. This study provides a rich resource to identify additional regulators that dictate BR actions in the cortex and other tissues in a precise spatiotemporal manner.
Methods

Plant materials and growth conditions
Arabidopsis accession Columbia-0 (Col-0) was used as a wild type. The following lines were previously described: brl1 GABI_134E10 (Jaillais et al., 2011); brl1-116brl1brl3 triple mutant (bri1-T) (Irami et al., 2012); pGL2-BRI1-GFP/bri1-T (Vragović et al., 2015); gtl1-1 (WiscDsLox413-416C9), df1-1 (SALK_106258), and gtl1-1 df1-1 (Shibata et al., 2018); JKD-Ypet recombineering line (Moreno-Risueno et al., 2015). Seeds were sterilized using 50% (v/v) bleach and homogenized to 22°C, 16 hours light/8 hours dark and grown for 7 days unless otherwise indicated. Chemical treatments were conducted by cooling the growth media to approximately 60°C after autoclaving and adding DMSO (a mock solvent), 1 µM Brassinazole (BRZ, SML1406, Sigma) or 100nM Brassinolide (BL, 21594, Cayman Chemical).

bri1-T was maintained as a heterozygote for bri1-116 and homozygous mutants were confirmed as previously described (Kang et al., 2017). Primers listed in Data S11 were used to amplify genomic DNA and the resulting 552bp amplicon was digested with PmeI whereas WT was cut into 314bp and 238bp fragments.

Transgenic reporters
To generate new reporters for BR-responsive genes, we first added the FASTRED seed coat selection cassette (Shimada et al., 2010; Stuttmann et al., 2021) and a MoClo (Engler et al., 2014) Level 1 acceptor site to the binary vector pICH86966 (Addgene plasmid #48075). pHA77-HA77-mCitrine and pGTL1-GTL1-mCitrine were assembled into this FASTRED destination vector using Level 1 BsaI golden gate assembly. To facilitate one-step promoter-reporter construction, we assembled an AarI flanked RFP dropout using the overhangs described in the Mobius (Andreu and Nakayama, 2018) upstream of Venus-H2B followed by the Ubiquitin10 terminator (tUBQ10), a plasma membrane marker (pUBQ10-mScarlet-LT16-Nos), and a constitutive histone maker (pUBQ10-H2B-CFP-t19s). Promoters containing up to ~3kb of sequence upstream of the ATG start codon for the gene of interest were PCR amplified with AarI containing primers and used to replace the AarI-RFP module in golden gate reactions to generate Promoter-Venus-H2B constructs. Our Venus-H2B reporter included a Ubiquitin tag to decrease reporter perdurance. Although the plasma membrane marker and histone marker were included as positive controls in the constructs, they were not further analyzed in this study.

Assemblies were confirmed by restriction digestion and sequencing, transformed into Agrobacterium, and used to transform Arabidopsis via floral dip (Clough and Bent, 1998). FASTRED positive T1 seeds were selected under a fluorescent dissecting scope and only lines with 3:1 segregation of seed coat fluorescence in the T2 generation were used. T2 lines with bright seed fluorescence were typically homozygous in our conditions. Therefore, we used bright T2 seeds or homozygous T3 seeds for experiments. We ensured that reporter signals were consistent across at least three independent transgenic lines.

Generation of mutant lines using multiplex CRISPR
We produced hat7 single mutants and hat7 hb13 hb20 hb23 quadruple mutants using FASTRED multiplex CRISPR constructs containing an intronized version of Cas9 (Grützner et al., 2021; Stuttmann et al., 2021). Two gRNAs were designed per gene using CHOP-CHOP (Labun et al., 2019). gRNA containing oligos were hybridized and cloned into pDGE sgRNA shuttle vectors using BpiI (Data S11). Each of the gRNA containing shuttle vectors were then assembled into pDGE666 (Addgene plasmid # 153231) using BsaI golden gate assembly, sequence verified, and transformed into wild-type Arabidopsis as described above. We selected FASTRED positive T1 seeds and subsequently screened FASTRED negative (putatively Cas9-free) T2 seeds for frameshift mutations using Sanger sequencing coupled with ICE analysis of CRISPR edits (Conant et al., 2022). The edits were similarly confirmed in the T3 generation and at least two homozygous alleles from independent lines were used for experiments.

BRI1 tissue-specific CRISPR
Two gRNA-BRI1 were simultaneously expressed in a tissue-specific manner (Decaestecker et al., 2019). Primers used for cloning of gRNA BRI1-2 (Feng et al., 2013) and gRNA BRI1-3 can be found in Data S11. The entry module pGG-B-AtU6-26-BRI1-2-C and pGG-A-AtU6-26-BRI1-3-B were generated by annealing oligos for each gRNA and ligating into BbsI-digested (New England Biolabs) Golden Gate entry vectors described in (Houbaert et al., 2018). Next, gRNA modules were combined with pGG-C-linker-G plasmid and cloned into pEN-R2-A-G-L3 by restriction-ligation using BsaI enzyme (New England Biolabs) to obtain pEN-R2-gRNA_BRI1-3-gRNA_BRI1-2-L3. This plasmid was combined with pDONR-L1-Cas9p-tagRFP-L2 (Wang et al., 2020), pDONRL4-L1r carrying either WER or CO2 promoters (Marquès-Bueno et al., 2016) and a destination vector pK8m34GW-FAST (Vanholme et al., 2013) in a MultiSite Gateway LR reaction (Thermo Fisher Scientific) to obtain expression clones. Expression clones were introduced into Agrobacterium C58 strain and used to transform pBRI1-BRI1-mCitrine/bri1 plants (Jaillais et al., 2011) by floral dip. T2 generation seeds were selected based on the presence of GFP signal in the seed coat and 7-day-old seedlings were used for phenotypic analysis. For each root used for quantitative analysis, BRI1-mCitrine signal was acquired in order to confirm efficiency of the tissue-specific knockout system. Statistical analyses were conducted in GraphPad Prism v.9 software.

Confocal microscopy
Confocal imaging for the majority of experiments was per-
formed using a Zeiss 880 equipped with a 40X objective. Excitation and detection were set as follows: Venus and mCitrine, excitation at 488 nm and detection at 509-571 nm; GFP, excitation at 488 nm and detection at 493-558 nm; PI staining, excitation at 561 nm and detection at 605-695 nm. Confocal images were processed using the Fiji package of ImageJ (Schindelin et al., 2012). Tile scans were stitched and representative median longitudinal sections for each image are shown. Identical settings were used for images that were directly compared.

For BR11 TSKO confocal, roots were imaged between a block of agar and cover glass in imaging chambers. Image acquisition was performed with a Fluoview1000 inverted confocal microscope (Olympus) equipped with a dry 20X objective (NA 0.75) using 514 nm laser excitation and a spectral detection bandwidth of 500–530 nm for mCitrine and 535 nm laser excitation together with a spectral detection bandwidth of 570–670 nm for PI.

BES1 and GTL1 Co-Immunoprecipitation (Co-IP)

Co-IP experiments were conducted as previously described (Xie et al., 2019b). p35S-FLAG-GTL1 and a p35S-FLAG-GUS negative control were cloned into pGWB412 (Nakagawa et al., 2007) using gateway LR reactions. The following construct combinations were co-transformed into Arabidopsis mesophyll protoplasts: p35S-BES1-GFP + p35S-FLAG-GUS; p35S-FLAG-GTL1 + p35S-FLAG-GUS; p35S-BES1-GFP + p35S-FLAG-GTL1. After overnight incubation, transformed protoplasts were harvested and homogenized in Co-IP buffer (50 mM Tris- HCl, pH 7.5, 150 mM NaCl, 10% (v/v) glycerol, 0.1% (v/v) Nonidet P-40, 1 mM phenylmethylsulfonyl fluoride, 20 mM Mg132, and protease inhibitor cocktail) for 1 h at 4 °C with rotation. 5 µg FLAG M2 antibody (F1804, Sigma) was pre-bound to 40 µL protein G Dynabeads (10003D, Thermo Fisher Scientific) for 30 min in phosphate-buffered saline (PBS) buffer with 0.02% Tween 20 at room temperature. The beads were washed once with the same PBS buffer and resuspended in Co-IP buffer. After protein extraction, 10 µL of anti-FLAG pre-bound Dynabeads was added to each sample for another 1.5 h incubation at 4 °C with rotation. Dynabeads were precipitated using a DynaMagnetic rack (12321D, Thermo Fisher Scientific) and washed twice with Co-IP buffer with Nonidet P-40 and three times with Co-IP buffer without Nonidet P-40. The IP products were eluted in 2XSDS sample buffer and used for immunoblotting with rabbit anti-GFP (A11212, Invitrogen) and rabbit anti-FLAG antibody (F7425, Sigma–Aldrich) at 1:1,000 dilution.

scRNA-seq profiling of Arabidopsis root protoplasts using the 10X Genomics chromium system

scRNA-seq experiments were performed as previously described (Shahan et al., 2022) with minor modifications. Plants were grown for 7 days as described above with the addition of 100 µm nylon mesh (Nitex 03-100/44) on the plates to facilitate root collection. For each sample, ~0.5cm root tips were harvested from 1000-3000 roots and placed into a 35mm petri dish containing a 70 µm cell strainer and 4.5mL enzyme solution (1.5% [w/v] cellulase [ONOZUKA R-10, GoldBio], 0.1% Pectolyase [Sigma P3026], 0.4 M mannitol, 20 mM MES (pH 5.7), 20 mM KCl, 10 mM CaCl2, 0.1% bovine serum albumin, and 0.000194% (v/v) beta-mercaptoethanol). The digestion was incubated on an 85 rpm shaker at 25°C for one hour with additional stirring every 15-20 minutes. The resulting cell solution was filtered twice through 40 µm cell strainers and centrifuged for 5 minutes at 500g in a swinging bucket rotor. The pellet was washed with 2mL washing solution (0.4 M mannitol, 20 mM MES (pH 5.7), 20 mM KCl, 10 mM CaCl2, 0.1% bovine serum albumin, and 0.000194% (v/v) beta-mercaptoethanol), centrifuged again at 500g for 3 minutes, and the pellet resuspended in washing solution at a concentration of ~2000 cells/µL. We loaded 16,000 cells, with the aim to capture 10,000 cells per sample with the 10X Genomics Chromium 3˚ Gene expression v3 or v3.1 kits. Cell barcoding and library construction were performed following the manufacturer’s instructions. cDNA and final library quality were verified using a Bioanalyzer High Sensitivity DNA Chip (Agilent) and sequenced on an Illumina NextSeq 500 or NovaSeq 6000 instrument to produce 100bp paired-end reads.

For BL scRNA-seq, we first grew plants on 1 µM BRZ to deplete endogenous BRs, then transferred plants to either a fresh BRZ plate or 100mM BL. We monitored the efficacy of these treatments using a constitutively expressed 35S-BES1-GFP line. In agreement with previous reports (Gampala et al., 2007; Ryu et al., 2007; Yin et al., 2002), BES1-GFP was predominantly present in the cytoplasm under low BR conditions resulting from BRZ treatment but accumulated in the nucleus following BL treatment (Figure S2A). We performed two separate BL scRNA-seq treatment experiments. The first consisted of a BRZ and 2 hour BL treatment. The second experiment included two additional replicates of BRZ and BL 2 hours along with the other time points in our time course (BL 0.5, 1, 4, and 8 hour treatments). Each of the BL treatments was staggered so that all samples were collected simultaneously. A total of 70,223 cells were recovered from the BL treatment scRNA-seq experiments. Wild type Col-0, bri1-T, and pGL2-BRI1-GFP/bri1-T were similarly profiled in a side-by-side scRNA-seq experiment under control conditions with two replicates per genotype, resulting in 34,861 cells. Lastly, scRNA-seq was performed on Wild type Col-0, gt1 df1, and gt1 df1 in duplicate under control conditions, resulting in 74,810 scRNA-seq
expression profiles.

**scRNA-seq data pre-processing**

Raw sequencing reads were demultiplexed from Illumina BCL files to produce FASTQ files for each sample using CellRanger mkfastq (v3.1.0, 10X Genomics). Reads were then aligned against the Arabidopsis TAIR10 reference genome to generate a gene-by-cell matrix using the scKB script: https://github.com/ohlerlab/scKB, which incorporates kallisto (Bray et al., 2016) and bustools (Melsted et al., 2019, 2021). Quality filtering of cells was performed using the R package COPILOT (Cell prepOcessing Pipeline kaListO busTools) (Shahan et al., 2022). COPILOT uses a non-arbitrary scheme to remove empty droplets and dying or low-quality cells. We used one iteration of COPILOT filtering, which adequately separated high-quality cells from the background in our samples based on an examination of barcode rank plots. To address issues with doublets and outliers, the resulting high-quality cells were further filtered to remove the top 1% of cells in terms of UMI counts, and putative doublets were removed with DoubletFinder using the estimated doublet rate from the 10X Genomics Chromium Single Cell 3’ Reagent Kit user guide.

**Normalization, annotation, and integration of scRNA-seq datasets**

Downstream analysis were carried out using Seurat version 3.1.5. Samples were first individually processed and examined. Data were normalized using SCTransform (Hafemeister and Satija, 2019) and all detected genes were subsequently retained for analysis, except those from mitochondria, chloroplasts or those affected by proteoplasting (absolute log2 fold-change >= 2) (Denyer et al., 2019; Shahan et al., 2022). Principal component analysis (PCA) was performed by calculating 50 principal components using the RunPCA function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function (with approx=FALSE). UMAP non-linear dimensionality reduction was next calculated via the RunUMAP function.

To follow the developmental progression from the meristem to the elongation zone more closely, we updated the root atlas (Shahan et al., 2022) developmental annotation to subdivide the meristem into the proliferation domain and transition domain as previously defined (Ivanov and Dubrovsky, 2013). The previous meristem annotation of the root atlas was based on correlation annotation by comparing each cell from scRNA-seq to bulk data from morphologically defined sections (Brady et al., 2007). On the other hand, HIGH PLOIDY2 was used to mark the meristem in a second bulk expression profile (Li et al., 2016), which corresponds to the proliferation domain defined by Ivanov and Dubrovsky. Therefore, we leveraged correlation-based annotations derived from Li et al., 2016 to re-label the meristem of the atlas. If cells were defined as “meristem” by both Li et al., 2016 and Brady et al., 2007, then they were re-labeled as the proliferation domain. Those that were called elongation in the Li et al., 2016 annotation, but meristem in the Brady et al., 2007 annotation were re-labeled as the transition domain. Finally, cells labeled as elongation in both Brady and Li datasets but annotated as meristem in the root atlas were re-labeled as elongation zone.

Consistent with our annotation, we found that cell cycle-related genes were enriched in the proliferation domain of the atlas (Figure S1B), whereas SMR1 (AT3G10525), a marker of endoreduplication, increased in the transition domain (Bhosale et al., 2018). The developmental annotation of cortex markers CO2 (AT1G62500) and CORTEX (AT1G09750) also matched their expression patterns in the root (Heidstra et al., 2004; Lee et al., 2006).

We used the receiver operating characteristic (ROC) test implemented in Seurat FindMarkers to identify genes enriched in each developmental zone. A largely distinct set of markers was enriched in the transition domain (Figure S1C and Data S2). These included genes involved in vesicle-mediated transport (Figure S1D), in line with the observation that vesicle recycling activity is highest in this region (Baluska et al., 2010).

We transferred the cell type and developmental stage labels from the wild-type atlas (Shahan et al., 2022) to each sample using the Seurat label transfer workflow (Butler et al., 2018; Stuart et al., 2019). To align corresponding cell types and developmental stages, we integrated samples from each experiment using the Seurat reference-based integration pipeline (Butler et al., 2018; Stuart et al., 2019). A sample from the atlas with the highest genes detected (sc_12) was used as a reference (Shahan et al., 2022) and two previously described samples (dc_1 and dc_2) (Denyer et al., 2019) were also included in each integration. PCA and UMAP were subsequently calculated for each integration object using the batch-corrected “integrated” assay as described above. Although sc_12, dc_1 and dc_2 were not used in any subsequent analysis, their inclusion at the integration step helped to generate a comparable UMAP projection among different integration objects that facilitates interpretability.

**Plotting gene expression values on the UMAP projection**

We subsequently examined changes in cell state caused by the BL treatments or in the mutants profiled by plotting the log-normalized, ‘corrected’ counts produced by the SCTransform function (Hafemeister and Satija, 2019) rather than the batch-corrected “integrated” values when visualizing changes in expression.

**Pseudotime estimation and heatmaps of gene expression trends**

Cortex cells were extracted from the integrated Seurat objects (BR time course, bril-T vs wild type and gtl1 df1 vs wild type). Pseudotime was then inferred on the SCT assay of the extracted cortex cells using CytoTRACE v0.1.0 (Gulati et al., 2020). Once the pseudotime was calculated,
the cortex cells were converted into SingleCellExperiment objects (Amezquita et al., 2019) before fitting a NB-GAM model (generalized additive model with a negative binomial distribution) using fitGAM function of tradeSeq R package v1.8.0 (Van den Berge et al., 2020). The model-predicted expression trends were plotted with ComplexHeatmap in R (Gu et al., 2016) (v2.10.0).

**Pseudobulk differential expression analysis**

Recent benchmarks point towards pseudobulk methods, which aggregate cell level counts for subpopulations of interest on a per-sample basis, as top performers for cross-condition comparisons in scRNA-seq (Crowell et al., 2020; Squair et al., 2021). Therefore, we employed a pseudobulk approach implemented in muscat (Multi-sample multi-group scRNA-seq analysis tools) (Crowell et al., 2020) to examine changes in each combination of cell type and developmental stage. Pseudobulk expression profiles were aggregated for each of these subpopulations by summing the raw counts using the aggregateData function. We then performed differential expression testing using the edgeR method (McCarthy et al., 2012) incorporated in the pbDS function. A term for the experimental batch and/or replicate was included in the contrast to adjust for potential batch effects. A gene was considered differentially expressed in a given subpopulation if the false discovery-rate adjusted p-value was <0.05, absolute fold change was >=1.5 and detection frequency was >=10% in one of the conditions. Gene ontology enrichment analysis was conducted on the differentially expressed genes using the R package “gprofiler2” (Kolberg et al., 2020). Comparisons between DEG lists were performed using the GeneOverlap package (version 1.12.0; http://shenlab-sinai.github.io/shenlab-sinai/). p-values for intersections between gene lists were computed using Fisher’s exact test. Visualizations were generated using Seurat (Stuart et al., 2019), ComplexHeatmap (Gu et al., 2016), and ggplot2 (Wickham, 2016).

**WOT differential expression along cortex cell wall + trajectories**

WOT constructs trajectories of cells from a reference time point by minimizing the difference over all genes (Schiebinger et al., 2019). The algorithm requires as input the expression profiles of cells as well as an estimation of their proliferation rate. We estimated proliferation rates using imaging data (Rahni and Birnbaum, 2019), as previously described (Zhang et al., 2021). As only the bottom 0.5 cm of each root was observed at each time, we expect some cells to exit the observed section due to proliferation. We estimated the number of cells that should exit the observed section based on the growth rate with the assumption that the section of root stays in equilibrium and assigned the calculated number of cells with the highest pseudotime a growth rate of zero so that they have no descendents in the observed root section at the next time point. We constructed trajectories using full gene expression profiles and evaluated the quality of the trajectories by checking the proportion of cells whose highest fate probability matched the annotation. We found that for 90% of cells the largest fate assigned by WOT matched the annotation, rising to 97% in the maturation zone where we have the greatest confidence in the annotation.

The cell wall signature was calculated for each cell by taking the sum of Z-scores for each of the 107 BR-induced cell wall-related genes in the signature (GO:0017554), truncated to [-5.5]. We defined the cortex cell wall+ subset as cortex cells with a cell wall score greater or equal to 1. This threshold was chosen as it selected less than 5% of cells from other lineages while still retaining >20% of cortex cells at the 2 hour time point. Any cortex cell that did not belong to the cortex cell wall+ group was labelled as “cortex cell wall-”. We performed differential expression on the cortex cell wall+ and cell wall- subsets at 2 hours, using WOT lineages to also perform differential expression on their putative ancestors and descendants. Statistical significance was evaluated using Welch’s t-Test with adjusted p-values for multiple tests, requiring t_{FDR} < 0.01. Results were ranked by the adjusted expression ratio

$$\frac{p_{\text{max}}}{p_{\text{min}} + \varepsilon}$$

with $\varepsilon = 0.1$, where larger $\varepsilon$ puts more emphasis on genes with non-zero expression in both groups. Using the same process, we also performed differential expression between groups of cells along the same WOT trajectory at adjacent time points.

**Gene regulatory networks**

In order to construct GRNs, we used CellOracle (v0.7.0) for single-cell GRN inference (Kamimoto et al., 2020). In the first stage of the CellOracle pipeline, a base GRN is defined, representing a global set of biologically plausible Transcription factor-Target interactions. We used publicly available scATAC-seq data from Arabidopsis roots (Farmer et al., 2021) GSE155304;GSM4698760; (Farmer et al., 2021) to determine regions of open chromatin. Cell Ranger ATAC (v1.2.0) was used to process raw scATAC seq data to call a peak-by-cell matrix. Cicero (v1.11.1) (Pliner et al., 2018) was implemented to infer a co-accessibility map of chromatin regions. Transcription start sites were then annotated based on the Arabidopsis TAIR10 genome assembly. Finally, peaks with weak co-accessibility scores were filtered following instructions of CellOracle manual (https://morrislab.github.io/CellOracle.documentation/tutorials/base_grn.html). To expand the number of TFs present in the base GRN we also included TF-Target interactions from DNA affinity purification sequencing (DAP-seq) (Bartlett et al., 2017) and a previously constructed integrative gene regulatory network (iGRN) (De Clercq et al., 2021). Our resulting base GRN contained 11.7 million interactions between 1,601 transcription factors and 31,019 target genes.

In the second step of the CellOracle pipeline, a regularized machine learning approach is used to define active edges and their regulatory strength in clusters or subpopulations of scRNA-seq data. In this process, the expression of target genes is predicted based on regulatory transcription factor
levels from the base GRN. Inactive edges with low predictive ability are pruned from the base GRN, revealing context-specific GRN configurations (Kamimoto et al., 2020).

To test CellOracle on Arabidopsis root data, we first inferred GRN configurations for each of the 36 cell type and developmental stage combinations in our WT atlas (Shahan et al., 2022) using the SCT normalized counts. We limited the base GRN to genes dynamically expressed along pseudotime for each cell type plus associated transcription factors (Pruneda-Paz et al., 2014). Each cell type GRN was then constructed with default parameters following the CellOracle manual. To filter network edges with the “filter_links” function, we retained the top 20,000 edges (p-value <0.01) for each subnetwork. This recovered known developmental regulators (Data S6 and Data S7) including MYB36 in the endodermis (Kamiya et al., 2015) and BRN1/BRN2 in the root cap (Bennett et al., 2010), confirming that CellOracle analysis of Arabidopsis root scRNA-seq data can infer GRNs configurations for particular cell identities and states.

We implemented similar procedures to infer context-specific GRN configurations for each cell type, developmental stage and time point of the BR time course samples (sc_43-50). We used transcription factors plus DEGs from BL 2 hour vs. BRZ pseudobulk analysis of each cell type/developmental zone combination. The resulting set of 201 GRN configurations spanned 767,970 edges between 1,164 transcription factors and 7,135 targets (Data S8). Network centrality measures were calculated using the built-in functions of the CellOracle pipeline (Data S9). The data needed to reproduce our results and jupyter notebooks demonstrating the processes are available on ARVE (https://shiny.mdc-berlin.de/ARVEX/).

Data and code availability

Single-cell RNA-seq data have been deposited at GEO: GSE212230. All original code is available at https://github.com/tmnolan/Brassinosteroid-gene-regulatory-networks-at-cellular-resolution.

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COMPETING FINANCIAL INTERESTS

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Fig. S1. Updated developmental annotation distinguishes between the proliferation domain and transition domain of the meristem. Related to Figure 1.

(A) Wild-type atlas of the Arabidopsis root from (Shahan et al., 2022) showing updated developmental stage annotation in which the meristem is divided into the proliferation domain and transition domain.

(B) Expression of markers in wild-type atlas supporting the developmental stage annotation. The color scale on the UMAP projection represents log normalized, corrected UMI counts. In dotplots, the size of the dot represents the percentage of cells in which the gene is expressed.

(C) Comparison of the number of shared markers for each of the developmental zones in the wild-type atlas. Color represents log10 p-values from the indicated overlaps calculated from Fisher’s exact test by GeneOverlap. The number of genes in each intersection is indicated inside each box.

(D) Enriched GO terms for markers of each developmental zone in the wild-type atlas.
Fig. S2. scRNA-seq identifies the elongating cortex as a site of BR-response. Related to Figure 1.

(A) p35S-BES1-GFP was used to monitor the efficacy of the sensitized system for BL scRNA-seq. Plants were grown for 7 days on 1 µM BRZ to deplete endogenous BRs, then transferred plants to either a fresh BRZ plate or 100nM BL. BES1-GFP was predominantly present in the cytoplasm under low BR conditions resulting from BRZ treatment but accumulated in the nucleus following BL treatment.

(B) Dotplots from the WT root atlas, BRZ, and BL 2 hours scRNA-seq showing that cell types and developmental stages are identified through label transfer. One marker gene for each cell type and developmental stage combination is shown. Circle size represents the percentage of cells in which a gene is expressed and color represents the average expression level of each gene. Black boxes denote markers from each cell type. Colors of side annotations indicate cell type and developmental stage.

(C) Comparison of BL 2 hour DEGs from scRNA-seq to BES1 and BZR1 ChIP targets and previous bulk BR RNA-seq datasets. Color represents log10 p-values from the indicated overlaps calculated from Fisher’s exact test by GeneOverlap. The number of genes in each intersection is indicated inside each box.

(E) Number of cell type/developmental stage combinations (subpopulations) in which each BL DEG is differentially expressed. 3,071/8,286 DEGs were significantly altered in only a single sub-population.

(F) Top 10 GO terms among BL up-regulated DEGs from scRNA-seq. Note the strong enrichment for cell wall-related GO terms in BL up-regulated genes in the elongating cortex.

(G) CSI1 expression in BL scRNA-seq data. The color scale on the UMAP projection represents log normalized, corrected UMI counts. In dotplots, the size of the dot represents the percentage of cells in which the gene is expressed.

(H) CSI1-H2B-Venus reporter grown on 1 µM BRZ for 7 days and transferred to 1 µM BRZ or 100nM BL for 4 hours. Inset shows CSI1 signals that are strongest in cortex and epidermis and increase with BL treatment. Propidium iodide-staining is shown in grey, with the color gradient indicating relative CSI1-H2B-Venus levels. Scale bars, 100 µm.

(I) XTH16 expression in BL scRNA-seq data. The color scale on the UMAP projection represents log normalized, corrected UMI counts. In dotplots, the size of the dot represents the percentage of cells in which the gene is expressed.

(J) XTH16-H2B-Venus reporter grown on 1 µM BRZ for 7 days and transferred to 1 µM BRZ or 100nM BL for 4 hours. Inset shows XTH16 signals in the endodermis that increase with BL treatment. Propidium iodide-staining is shown in grey, with the color gradient indicating relative XTH16-H2B-Venus levels. Scale bars, 100 µm.
Fig. S3. Waddington optimal transport identifies JKD as a BR responsive transcription factor along cortex trajectories. Related to Figure 2.

(A) UMAP showing cell type annotation across BL scRNA-seq treatment time course. This panel is repeated from the main text figure as a reference for the panel below.

(B) UMAP projection colored by cell wall signature, calculated as the sum of Z-scores for each of the 107 BR-induced cell wall-related genes in the signature (GO:0071554), truncated to [-5,5].

(C) Expression of JKD in BRZ and BL 2 hour scRNA-seq. The color scale represents log normalized, corrected UMI counts.

(D) pJKD-JKD-YPet grown on 1 µM BRZ for 7 days and transferred to 1 µM BRZ or 100nM BL for 4 hours. Inset shows JKD signals in the elongating cortex that increase with BL treatment. Propidium iodide-staining is shown in grey, with the color gradient indicating relative JKD-Ypet levels. Scale bars, 100 µm.
Fig. S4. Analysis of the triple receptor mutant bri1-T reveals changes in cortex expression and distinct gene expression patterns of pGL2-BRI1-GFP/bri1-T. Related to Figure 3.

(A) Number of DEGs for each cell type/developmental stage combination in bri1-T scRNA-seq compared to WT. Color indicates the number of up-regulated vs down-regulated genes.

(B) GO enrichment of bri1-T vs wild-type DEGs.

(C) Number of DEGs for each cell type/developmental stage combination in pGL2-BRI1-GFP/bri1-T scRNA-seq compared to WT. Color indicates the number of up-regulated vs down-regulated genes.

(D) GO enrichment of pGL2-BRI1-GFP/bri1-T vs wild-type DEGs.

(E) Number of DEGs for each cell type/developmental stage combination in pGL2-BRI1-GFP/bri1-T scRNA-seq compared to bri1-T. Color indicates the number of up-regulated vs down-regulated genes.

(F) GO enrichment of pGL2-BRI1-GFP/bri1-T vs bri1-T DEGs.
Fig. S5. BRI1 CRISPR TSKO. Related to Figure 4.

(A) Two individual transgenic lines for pWER-BRI1-CRISPR and pCO2-BRI1-CRISPR exhibiting similar BRI1-mCitrine expression patterns. Scale bars, 50 µm.

(B) Seven-day-old BRI1-CRISPR transgenic seedlings with roots shorter than those of the wild-type (Col-0) control and complemented pBRI1-BRI1-mCitrine/bri1. Scale bar represents 1 cm.

(C) Quantification of the root length of transgenic lines shown in (B). All individual data points are plotted. Red horizontal bars represent the means and error bars represent s.d. Significant differences between transgenic lines and the WT control were determined by one-way ANOVA and Dunnett’s multiple comparisons tests. ***P<0.001, **P<0.01, *P<0.05. n.s. not significant.
Fig. S6. HAT7 and GTL1 family transcription factors are BR responsive regulators along cortex trajectories. Related to Figure 5 and Figure 6.

(A) UMAP projections showing expression levels of HAT7 and GTL1 family transcription factors over the BR time series scRNA-seq experiment. The color scale represents log normalized, corrected UMI counts.

(B) Expression trends for indicated transcription factors along WOT cortex cell wall + (cortex +) vs cortex cell wall - (cortex -) trajectories.

(C) pDF1-DF1-GFP grown on 1 µM BRZ for 7 days and transferred to 1 µM BRZ or 100nM BL for 4 hours. Inset shows DF1 signals in the elongating epidermis and cortex that increase with BL treatment. Propidium iodide-staining is shown in grey, with the color gradient indicating relative DF1-GFP levels. Scale bars, 100 µm.

(D) Quantification of the root length in the indicated mutants. hat7 quad 1-2 and 9-9 represent two independent CRISPR mutants of hat7 hb13 hb20 hb23. hat7 quad 1-2 is used as a representative allele throughout the manuscript unless otherwise indicated.

(E) Quantification of meristematic cortex cell length, defined as the first 20 cells of individual roots starting from the quiescent center.

(F) Quantification of mature cortex cell length. For D-F, all individual data points are plotted. Red horizontal bars represent the means and error bars represent s.d. Significant differences between mutants and the wild-type control were determined by one-way ANOVA and Dunnett’s multiple comparisons tests. ***P<0.001, **P<0.01, *P<0.05.
Fig. S7. BES1 and GTL1 interact and share a common set of target genes. Related to Figure 7.

(A) Comparison of BES1 or BZR1, GTL1, and DF1 ChIP targets showing an overrepresentation of shared target genes.

(B) Comparison of ChIP targets from (A) with BL 2 hour vs BRZ DEGs. The top 20 cell type/developmental stage combinations that are enriched for BES1 or BZR1 and GTL1 shared targets are shown. For (A) and (B) color represents log10 p-values from the indicated overlaps calculated from Fisher’s exact test by GeneOverlap. The number of genes in each intersection is indicated inside each box.

(C) Co-Immunoprecipitation demonstrating BES1 interaction with GTL1. GTL1-FLAG immunoprecipitated with anti-FLAG beads pulled down BES1-GFP, whereas a GUS-FLAG negative control did not.

(D) Number of DEGs for each cell type/developmental stage combination in gtl1, df1 or gtl1 df1 scRNA-seq compared to WT. Color indicates the number of up-regulated vs down-regulated genes.

(E) GO enrichment of DEGs in gtl1, df1 or gtl1 df1 scRNA-seq compared to WT.
Supplemental Tables

Data S1. Summary of the scRNA-seq samples reported in this study. Related to Figure 1.

Data S2. Marker genes for updated developmental annotation of Shahan et al WT root atlas. Related to Figure 1.

Data S3. DEGs from pseudobulk analysis of scRNA-seq datasets. Related to Figure 1, Figure 3, and Figure 7.

Data S4. DEGs from WOT analysis of cortex cell wall + scRNA-seq datasets. Related to Figure 2.

Data S5. DEGs from WOT analysis of each cell type and developmental stage from BR time series scRNA-seq. Related to Figure 2.

Data S6. CellOracle GRN inferred from wild-type atlas. Related to Figure 6.

Data S7. CellOracle GRN centrality metrics from wild-type atlas. Related to Figure 6.

Data S8. CellOracle GRN inferred from BR time series. Related to Figure 6.

Data S9. CellOracle GRN centrality metrics from BR time series. Related to Figure 6.

Data S10. Predicted targets of HAT7 and GTL1 family TFs from elongating cortex CellOracle GRNs in BR time series. Related to Figure 6.

Data S11. Oligos used in this study.

Video S1. Time-lapse confocal microscopy showing pC/VIF2-H2B-Venus in wild-type and gtl1 df1.

Note that Supplemental data that exceeds the file size limit can be found at: https://shiny.mdc-berlin.de/ARVEX/