Data-Driven Modeling of Intracellular Auxin Fluxes Indicates a Dominant Role of the ER in Controlling Nuclear Auxin Uptake

Highlights

- Auxin transport probe NBD-NAA does not enter the nucleus by simple diffusion
- Modulation of auxin transport at the ER affects auxin-mediated responses
- Model of nuclear auxin fluxes interconnects short- and long-term auxin responses
- The ER is the main conduit for nuclear auxin uptake

In Brief

Middleton et al. study how the plant phytohormone auxin enters the nucleus by using quantitative phenotyping in single plant cell systems and bespoke mathematical models that relate controlled perturbations to experimentally measurable responses. Their findings show that auxin predominantly enters the nucleus via the endoplasmic reticulum.
SUMMARY

In plants, the phytohormone auxin acts as a master regulator of developmental processes and environmental responses. The best characterized process in the auxin regulatory network occurs at the subcellular scale, wherein auxin mediates signal transduction into transcriptional programs by triggering the degradation of Aux/IAA transcriptional repressor proteins in the nucleus. However, whether and how auxin movement between the nucleus and the surrounding compartments is regulated remain elusive.

Using a fluorescent auxin analog, we show that its diffusion into the nucleus is restricted. By combining mathematical modeling with time course assays on auxin-mediated nuclear signaling and quantitative phenotyping in single plant cell systems, we show that ER-to-nucleus auxin flux represents a major subcellular pathway to directly control nuclear auxin levels. Our findings propose that the homeostatically regulated auxin pool in the ER and ER-to-nucleus auxin fluxes underpin auxin-mediated downstream responses in plant cells.

INTRODUCTION

The plant hormone auxin, while structurally simple, is known to regulate a multitude of processes in plants—from embryo patterning and development to de novo organ formation and environmental responses (Freire Rios et al., 2014; Wu et al., 2015; Naseem et al., 2015; Taylor-Teeples et al., 2016). Although it is not yet completely understood how auxin can orchestrate such an astonishing range of processes, there has been significant progress in our comprehension of auxin perception and signal transduction.

The best characterized pathway for auxin signaling involves the regulation of genes at the transcriptional level. The current accepted model of auxin-mediated transcriptional responses considers the interplay between auxin and nuclear receptors F-box TIR1/AFBs, transcriptional repressors Aux/IAA, and transcriptional factors ARFs, with 6, 29, and 23 members identified in Arabidopsis thaliana, respectively (Salehin et al., 2015). At elevated nuclear auxin levels, Aux/IAAs and TIR1/AFBs form complex receptor complexes in an auxin-dependent manner. Following complex formation, Aux/IAAs are ubiquitinylated and marked for proteolysis, thus releasing ARFs to drive the transcriptional responses (Dharmasiri et al., 2005; Kepinski and Leyser, 2005). The dynamic range of auxin is tuned by the combinatorial assembly of the various Aux/IAAs with TIR1/AFBs, resulting in a series of co-receptors with different auxin binding affinities (Calderón Villalobos et al., 2012). Upon taking into account the critical role of auxin levels in the assembly of the co-receptor complexes (Tan et al., 2007), key questions arise: How does auxin enter the nucleus? And is this process regulated?
pores without restriction (Wei et al., 2003). However, it remains unclear whether this alone could account for the observed dynamics of auxin-regulated gene expression (Wang and Estelle, 2014) or whether a rate-limited mechanism or mechanisms might be involved.

Novel fluorescent auxin analogs have been developed that mimic native auxin transport and accumulation in vivo without affecting auxin signaling (Hayashi et al., 2014). These 7-nitro-2,1,3-benzoxadiazole (NBD)-conjugated synthetic auxin analogs allowed visualization of subcellular auxin transport and distribution, revealing their preferential accumulation in the endoplasmic reticulum (ER) of tobacco BY-2 and Arabidopsis root cells (Hayashi et al., 2014). Moreover, the ER-residing auxin transport proteins from the PIN and PILS families affect cellular auxin homeostasis and nuclear auxin signaling (Barbez et al., 2012; Dal Bosco et al., 2012; Ding et al., 2012; Mravec et al., 2009). The outer membrane of the nuclear envelope is continuous with the ER, such that the space between the inner and the outer nuclear membranes is directly connected with the lumen of the ER (Meier and Brkljacic, 2010). Altogether, we sought to disentangle the interplay among the ER, the cytosol, and the nucleus in relation to nuclear auxin uptake.

We developed a joint theoretical-experimental approach to unravel this puzzle by resolving two major challenges: the complex organization of the plant due to cellular heterogeneity and the need to quantitatively assess intracellular auxin transport dynamics. Auxin transport between subcellular compartments cannot be measured directly due to the lack of non-invasive methods and tools. Therefore, indirect time-resolved auxin dose responses in single plant cell systems at different scales were combined with modeling approaches. Data-driven mathematical models of intracellular auxin transport were specifically tailored to interpret nuclear auxin uptake. Analysis of the model results indicated a dominant role of ER-to-nucleus auxin fluxes in regulating nuclear auxin levels. Our finding suggests that ER auxin pool and ER-to-nucleus transport rates underpin auxin-mediated signaling responses in plant cells.

RESULTS

Concept of Nuclear Auxin Uptake Exploration

The fundamental idea behind our approach is to infer intracellular auxin fluxes, which cannot be evaluated directly, using bespoke models that relate controlled perturbations to experimentally measurable responses. It was therefore crucial to minimize the complexity of the plant system by using single plant cells (Figure 1A). As controlled perturbations, we modulated intracellular auxin fluxes by overexpressing ER-localized auxin transporters (Figure 1A) and developed assays to measure long-term and short-term responses at different auxin concentrations (Figure 1B). For long-term responses, we monitored expansion of single plant cells, whereas for short-term responses, we used an auxin sensor (Figure 1B). Models describing these responses were independently coupled to a model for the nuclear auxin fluxes (Figure 1C). The coupled models were fitted to the response data to generate estimates for various flux parameters among the nucleus, the ER, and the cytosol. To ensure that our results do not depend on individual parameter estimates (which may be inaccurate), we based our analysis on sets of parameter estimates for which the coupled models behave consistently with the data; these sets define a confidence region (Figure 1D). Histograms of the relative auxin fluxes were generated using estimates from the confidence regions, which were robust to experimental and biological variation (Figure 1D), thus providing a means to elucidate the underlying mechanisms of nuclear auxin transport. We find consistent results from our analysis of the two responses, independently of the timescales involved.

Accumulation of Fluorescent Auxin Analog in the Nucleus Is Not Driven by Diffusion Alone

It is generally considered that a small organic molecule such as auxin should diffuse freely between the cytosol and the nuclear plasm via the nuclear pores. In the context of auxin subcellular distribution, this would mean that the nucleoplasm and the cytosol could be conceptually treated as a single compartment. To test this assumption, we used the NBD-conjugated naphthalene-1-acetic acid (NAA) fluorescent auxin analog, NBD-NAA. NBD-NAA was shown to be active for auxin transport while remaining biologically inactive (Hayashi et al., 2014). The diester form of carboxyfluorescein (CF), 5(6)-carboxyfluorescein diacetate (CFDA), was included in the study as a control diffusion dye. Similar to auxins such as IAA and NAA, CFDA freely passes through the plasma membrane via diffusion, and once in the cell, the dye is hydrolyzed to acetic acid and negatively charged CF, a membrane-impermeable fluorescent probe (Oparka et al., 1994). In contrast to auxins, there are no known CFDA-specific transporter proteins. Both CF and NBD-NAA have carboxylic acid and similar molar masses (376.3 and 408.4 g mol⁻¹, respectively) that are higher than the molar mass of NAA (186.2 g mol⁻¹) yet well below what is considered the size exclusion barrier of nuclear pores (Görlich and Kutay, 1999).

Spatiotemporal changes in probe levels were monitored in living Arabidopsis protoplasts, a versatile analytical system to study cellular and subcellular processes in a uniform single cell population (Yoo et al., 2007). NBD-NAA could be detected in cells within 5 min of incubation (Figures 2A and 2C). The probe largely (maximum value at Pearson’s R value, 0.79) co-localized with the ER-mCherry marker, as reported previously (Hayashi et al., 2014). This, together with a clearly more diffuse pattern, indicated availability of the fluorescent auxin analog in the cytosol as well. Surprisingly, however, the NBD-NAA signal in the nucleus appeared considerably weaker.

To compare changes in fluorescence intensity of NBD-NAA and CF in the nuclear and the extra-nuclear regions (cell excluding nucleus) over time, mCherry-tagged histone H2B (HTB4-mCherry) was transiently expressed as the nucleus marker (Figures 2B and 2C). Spatiotemporal tracking of cells at 5 min intervals showed that CF fluorescence intensity gradually increases in the cells and nuclei over a 30 min period (Figure 2B). The fluorescence intensity of NBD-NAA, however, remained relatively constant in both the nuclei and the whole cells (Figure 2C), suggesting that intracellular auxin levels were rapidly established and auxin-transport machinery constantly maintains a balance between cellular auxin influx and cellular auxin efflux.
To quantitate these ratios, we generated, for each cell, histograms of pixel intensities (distributions) of the whole cell and the nuclear and extra-nuclear regions (Figure S1A). Due to differences in uptake of the fluorescent probes by individual cells, there is significant inter-cellular variability in the median (Figure S1C). In addition, each distribution has a long tail (Figure S1B), so differences are most apparent when one considers the upper and lower quantiles. We therefore used quantile-quantile plots to compare the distribution of the ratios for both CF and NBD-NAA (Figures 2D–2E; Figures S1D and S1E) and normalized these quantiles by the medians to account for the inter-cellular variability. If one assumes that NBD-NAA and CF diffuse freely through the nuclear pores, then the quantiles of fluorescence intensity for the nuclear and extra-nuclear regions should be similar and therefore lie along the diagonal of the quantile-quantile plot.

For NBD-NAA, the 5% quantiles (Figure 2D) fall along the diagonal line, implying that the least bright pixels of nuclear segments are similar in fluorescence intensity to the least bright pixels in the extra-nuclear segments (Figure S1B). However, the 95% quantiles for NBD-NAA (Figure 2D) lie below the line. This means that the brightest pixels in the nuclear segments are less bright than the brightest pixels in the extra-nuclear segments (Figure S1B). The corresponding quantile-quantile plot for CF (Figure S1D) shows that both the 5% and the 95% quantiles lie above the diagonal line, indicating that fluorescence intensities in the nuclear region are stronger than those in the extra-nuclear region. Thus, for both CF and NBD-NAA, the fluorescence intensity distribution in the nuclear region is different from the distribution in the extra-nuclear region.

Quantile-quantile plots were used to analyze time-dependent changes in NBD-NAA by comparing the distribution of NBD-NAA at 5 and 30 min (Figure 2E) for the same compartment. These show that the 5% and 95% quantiles lie along the diagonal line for both the nuclear and the extra-nuclear regions, confirming that NBD-NAA rapidly reaches equilibrium. In contrast, analogous quantile-quantile plots for CF (Figure S1E) show that all quantiles lie above the diagonal line for both regions, reflecting the gradual increase in CF intensity between the two time points.

Our results show that the movement of both CF and NBD-NAA between the nuclear and the extra-nuclear regions (and subsequently their nuclear accumulation over time) is not consistent with the simple conceptual model in which the nucleoplasm...
and the cytosol can be treated as a single compartment, whereby small molecules can rapidly pass through the nuclear pores by passive diffusion. These results therefore indicate that the nuclear uptake of auxin is driven by processes other than diffusion.

Modulation of Auxin Fluxes at the ER Affects Cell Division and Enhances Cell Expansion

We next developed genetic perturbation assays to understand the link between subcellular distribution of physiologically active auxin and downstream signaling responses. ER-localized auxin transporters PIN5 (Mravec et al., 2009) and PIN8 (Dal Bosco et al., 2012; Ding et al., 2012) were selected as potential molecular modulators of auxin fluxes between the cytosol and the ER. The chemically stable synthetic auxin NAA (Dunlap et al., 1986), which is, in contrast to NBD-NAA, a physiologically active compound, was used to study cellular responses to auxin. Due to its hydrophobicity, NAA freely diffuses through the plasma membrane barrier (Delbarre et al., 1996).

We first confirmed co-localization of PIN5 and PIN8 with an ER marker in Arabidopsis and tobacco leaf protoplasts. Previous expression analysis of PIN8 under native regulatory elements (Dal Bosco et al., 2012) showed no detection of this gene in Arabidopsis leaves. Therefore, the Cauliflower mosaic virus (CaMV) 35S promoter was selected to transiently overexpress PIN5-GFP and PIN8-GFP. PIN8-GFP was mainly co-localized with the ER-mCherry marker (Figure S2A). After 48 hr of culture, PIN5-GFP was no longer observed in most cells, while transiently expressed PIN8-GFP or ER-YFP marker could be detected for more than a week (Figures S2B and S2C).

Figure 2. Analysis of NBD-NAA and CF Dynamics in Arabidopsis Protoplast-Derived Cells

(A) Co-localization of NBD-NAA (5 min treatment) and ER-mCherry marker (upper panel). A more diffuse NBD-NAA pattern and probe-specific accumulation sites (lower panel) around the nucleus were surrounded by chloroplasts (Chpl). Scale bars correspond to 10 μm.

(B and C) Time course study reveals distinct dynamics of CF and NBD-NAA in nuclear and extranuclear regions. 3D projections of representative cells incubated in medium supplemented with CFDA (B) or NBD-NAA (C). Image series were acquired at 5 min intervals, and the nucleus was identified using the HTB4-mCherry histone marker.

(D and E) Quantile-quantile plots of fluorescence in the nuclear and extra-nuclear regions using the 5th percentile (triangles) and 95th percentile (crosses) normalized against the median (50th percentile) for NBD-NAA-treated cells (n = 15 per treatment) are as shown (D). Analogous quantile-quantile plots comparing the fluorescence distributions at 5 and 30 min for the nuclear and extra-nuclear regions separately (E).

See also Figure S1.
The expansion and division of single cells are commonly exploited as physiological markers of auxin effects (Campanoni and Nick, 2005). Therefore, these parameters were assessed in immobilized tobacco leaf protoplasts to quantify consequences of perturbed intracellular auxin fluxes at the ER.

To ensure that functional properties of transiently overexpressed PINs in protoplasts were not affected by translational fusion with the fluorescent tags, tagged and non-tagged versions of tested PINs were compared by manual tracking of immobilized transformed cells at 24 hr intervals. In a case of using non-tagged PINs, an additional fluorescence reporter gene encoding the cytosolic mCherry protein was cloned in the expression plasmids. This strategy enabled identification of cells of interest (expressing non-tagged PINs) by detecting the mCherry fluorescence signal (Figure 3B). After 2 days of culture in the presence of 0.5 μM NAA, no statistically significant difference in cell sizes could be found among samples transformed with PIN5, PIN8, or ER-marker control (Figure 3C). In contrast, 3- to 4-day-old cells overexpressing non-tagged PIN8 were significantly larger than the control cells or PIN5-overexpressing cells (Figure 3C). Cells expressing non-tagged or tagged PIN8 (Figure 3B) showed a similar behavior in cultures, thus suggesting that function of PIN8 in protoplasts is not affected upon protein tagging with the yellow fluorescence marker VENUS. Furthermore, cell divisions in PIN8-overexpressing cells were strongly suppressed (Figures 3B and 3D; Figure S2B). This, together with the elevated cell expansion...
behavior, resembles the auxin-starvation phenotype observed in tobacco suspension cells under auxin-deprived conditions (Winicur et al., 1998).

Overall, these results showed that (1) an immobilized protoplast-derived cell model represents a potent experimental platform to quantitatively analyze cell expansion dynamics and (2) perturbation of auxin fluxes at the ER obtained by overexpression of PIN8 (but not of PIN5) affects cell expansion. We therefore chose PIN8 gain-of-function mutant (PINBOX) as a tool for modulating intracellular auxin fluxes in cell expansion studies using immobilized protoplasts.

**Cell Expansion Dynamics Model Indicates a Mechanism Regulating Nuclear Auxin Levels**

Current methods for the immobilization of single plant cells, which cannot grow by adhering to a surface, are not optimized for microscopic multi-time point observations, especially for medium- to large-scale experimental setups. To overcome this, we developed an approach, named protoplast monolayer embedding (PME), for efficient and robust immobilization of protoplast-derived cells directly in multi-well plates (Figure S3A). To quantitatively study cell expansion dynamics from microscopic 4D image sequences, we next developed AutoOvuscule, a computational tool for automated segmentation, tracking, and measurement of cell morphology over time and discriminating living and dead cells (Figure S3B). Immobilized protoplasts can be observed over continuous intervals (days to weeks) under diverse culture conditions (Figures S3C–S3J).

In earlier studies, it was shown that auxin stimulates swelling of freshly prepared protoplasts (Steffens and Lüthen, 2000; Steffens et al., 2001). There, volumetric changes were monitored in cells lacking the cell wall. In this study, we monitored expansion of the cells after recovery of the cell wall, because genetic and pharmacologically induced modulations of the extracellular matrix have demonstrated crosstalk between auxin and cell wall function (Feraru et al., 2011; Steinwand et al., 2014). The cell wall of tobacco protoplasts is recovered within 24 hr, even in auxin-free culture conditions (Figure S4A). Immobilized tobacco leaf protoplasts from wild-type (WT) or PINBOX (Dal Bosco et al., 2012) were used to quantitatively assess cell expansion in response to a range of auxin concentrations (Figures S4B and S4C). Acquired image time series of cultured protoplasts were processed using AutoOvuscule to automatically track individual cells, identify their status (dead or alive), and measure diameters at each time step. Divided and non-divided cells cannot be discriminated using the tool, because daughter cells occupy the same space as a parent cell after the initial cell division (Figure S5A). Therefore, 72 hr was selected as a terminal time point. According to expert-guided image inspection, initial cell divisions could be clearly identified at this time point in WT or PIN8OX PME cultures incubated with 0.5–25 or 5–50 μM NAA, respectively. Cells that had died during the 72 hr observation period were excluded from the analysis. Averaged values of cell diameters were plotted against time for the different experimental conditions (Figure 4A) and visualized as histograms of cell diameters for individual conditions and time points (Figure S5B).

We observed that WT cells cultured in auxin-free conditions initially expanded at a constant rate before decelerating noticeably by 48 hr (Figure 4A). In cells treated with sub- and micromolar concentrations of NAA, the deceleration of expansion occurred even earlier (Figure 4A). In contrast, PINBOX cells treated with concentrations of NAA up to 5 μM expanded at an approximately constant rate, so no significant deceleration could be observed (Figure 4A). Only for cells cultured in the presence of very high concentrations of NAA (10–50 μM) was a deceleration phase observed within 72 hr (Figure 4A). The highest auxin concentration tested (50 μM NAA) was toxic for WT cells, but not PINBOX cells (Figures S4B and S4C). This decreased sensitivity of PINBOX cells to toxic NAA concentration, in concert with auxin-insensitive cell expansion rates at auxin concentrations below 5 μM (Figure 4A; Figures S4B and S4C), consistently indicated that PIN8-mediated changes in the ER auxin pool resulted in significantly decreased responsiveness of cells to exogenous auxin.

We next sought a simple model enabling quantification of the cell expansion dynamics. Because cell diameter measurements at each time point and experimental condition were observed to be normally distributed (Figure S5B), we reasoned that a good description could be obtained by basing the model on the averaged cell expansion data (Figure 4A), rather than developing a more complex model describing the expansion of individual cells.

We assumed that changes in average cell volume are best captured by logistic growth. This model (Figure 4B) contains two parameters: the initial expansion rate (alpha), which defines over what timescale the cell might reach its maximal volume, and the maximal cell volume (V_max). At larger timescales, cell alphas will decrease until steady-state volume V_max is attained (Figure 4C). To fit the model to the data and avoid overfitting, we adopted the simplest assumption, namely, that auxin regulates only one of the two model parameters. Because the relationship between the concentration of exogenous auxin and the value of these parameters was unknown, we fitted each experimental condition separately. We found that auxin regulation of the V_max reproduced the experimental data significantly better (Figure 4A) than regulation of the cell alpha (Figure S5C). Best-fit parameter estimates are illustrated in Table S1.

In both PINBOX and WT cells, V_max decreased with increasing exogenous NAA concentration, whereas V_max was always larger in PIN8OX cells than in WT ones (Figure 4D). WT cells exhibited a strong response of V_max to NAA concentrations below 2.5 μM but a weak response above 2.5 μM. In contrast, the response range of PIN8OX cells was extended. At 10 μM NAA, the V_max of PINBOX cells was comparable to that of untreated WT cells, while at 50 μM NAA, the V_max was comparable to that of WT cells treated with 2.5 μM. This illustrates that modulation of the ER NAA pool results in a substantially altered response of V_max to exogenous NAA.

Our data and the cell expansion model indicate links among exogenous auxin, modulation of the auxin flux at the ER, and cell expansion dynamics. While exogenous auxin showed a pronounced concentration-dependent inhibition of cell expansion, cell expansion per se is a highly complex process in which numerous direct and indirect auxin-triggered responses and auxin-independent events are coordinated over a long period. We therefore further explored how dynamic changes of cytosolic and ER-auxin pools influence nuclear auxin signaling.
Regulation of Auxin-Induced Genes IAA1 and IAA5 Is Affected in PIN8OX Cells

To test how differences in the cell expansion dynamics are correlated with altered nuclear auxin signaling, we analyzed the expression of IAA1 and IAA5, which are among the fastest auxin-induced genes (Abel et al., 1995), in protoplast-derived cells. Cells from Arabidopsis WT or PIN8OX were incubated for 1 hr at various concentrations of NAA. The qRT-PCR analysis
showed that expression levels of both IAA1 and IAA5 in PIN8OX cells were significantly lower than in the WT cells at all concentrations tested (Figures S6A and S6C). This confirmed that PIN8 directly interfered with the auxin cellular responses. Time-resolved analysis of IAA1 and IAA5 at 1 μM NAA showed that prolonged incubations with auxin only partially rescued the expression levels of either gene (Figures S6B and S6D). These results are consistent with reduced expression of auxin responsive genes in PIN8OX previously observed in planta (Dal Bosco et al., 2012). Changes in expression levels of auxin-induced genes in WT and PIN8OX cells indicate that the altered response of Vmax to exogenous NAA in the cell expansion model and the nuclear auxin levels are interconnected.

Mathematical Modeling Predicts the ER as the Main Conduit for Nuclear Auxin Flux

We next sought to understand how alteration of the auxin fluxes at the ER modulated nuclear auxin signaling by developing a mathematical model of nuclear auxin fluxes. Because no methods directly demonstrate transport directionality of intracellular auxin transporters, such as PIN8, we assumed that PIN8 transports auxin from the ER into the cytosol due to the following rationale. Radioactive auxin transport assays performed on whole cells, which were independently performed by different research groups (Ding et al., 2012; Ganguly et al., 2010), consistently showed decreased accumulation of auxin in PIN8OX cells in comparison to the control cells. Therefore, these data exclude accumulation of auxin due to intracellular compartmentalization (which otherwise would lead to the increased radioactivity within the cells) and corroborate the assumption that PIN8 transports auxin from the ER into the cytosol, from which it is exported out of the cells by the efflux machinery.

Our data on subcellular distribution of the fluorescent auxin analog NBD-NAA (Figure 2; Figure S1) suggested that auxin uptake into the nucleus does not occur via simple diffusion. The nuclear auxin flux model (Figure 5A) assumes that auxin is exchanged among the cytosol, the nucleus, and the ER. Thus, there are two fluxes into the nucleus: one via the ER and one via the cytosol. The relative strength of these different fluxes is determined by the model parameters, which are estimated by fitting the model to experimentally derived Vmax values (Figure 4D). To describe the flux of auxin between compartments A and B, we used saturating transport functions (Figure 5A). This approach captures both linear and non-linear transport behavior by adjusting the associated constant KA/B. Additional model parameters (for details, see Supplemental Experimental Cell Reports 22, 3044–3057, March 13, 2018 3051
steady-state and fitted our nuclear auxin flux model to the estimate on a much faster timescale than that of growth (days). We therefore assumed that the nuclear auxin flux model is at steady-state and fitted our nuclear auxin flux model to the estimates of $V_{\text{max}}$ (Figure 4D, solid lines). To understand how differences between WT and PIN8OX $V_{\text{max}}$ curves are reflected in the parameters of the nuclear auxin flux model, we first performed a global parameter scan to estimate the 95% confidence region of the parameter estimates (Table S2). We then calculated the ER-to-nucleus or cytosol-to-nucleus auxin flux ratio (for details, see Supplemental Experimental Procedures) for each parameter set inside the confidence region (Figures 1D and 5C).

In every case, flux from the ER to the nucleus was predicted to be significantly larger than flux of auxin from the cytosol, with a median 10-fold difference between the two (Figure 5C). In other words, the model predicts that of the two fluxes into the nucleus, the ER-to-nucleus one dominates over the cytosol-to-nucleus flux. The corresponding receptor occupancies (Figure 5B, model formulation) were also calculated for each parameter set taken from the confidence region: occupancy in PIN8OX cells is maintained at a level (regardless of the exogenous auxin concentration) that is 2–4 times lower than that in WT (Figure 5D). Similar fold changes in gene expression between WT and PIN8OX cells were determined in our qRT-PCR experiments (Figures S6A–S6D). Hence, our model indicates that the observed differences between PIN8OX and WT cells can be largely attributed to alterations in nuclear auxin transport.

To further explore and to validate our model predictions, we exploited a recently developed quantitative genetically encoded auxin sensor (Wend, et al., 2013). We first generated time-resolved auxin dose-response data of the sensor for WT and PIN8OX cells (Figure 6A). Arabidopsis leaf protoplasts expressing the sensor were incubated in a cell expansion model (Figure 5C), we calculated the ER-to-nucleus flux compared to the cytosol-to-nucleus fluxes. Altogether, the early nuclear auxin response model confirms two key predictions made using the cell expansion dynamics data and the nuclear auxin flux models (Figures 4 and 5), namely, that we next developed an early nuclear auxin response model using the sensor data to ascertain how differences in the sensor degradation kinetics might correspond to altered nuclear auxin levels (Figure 6B). The model takes into account interactions among auxin, its receptors (TIR1/AFBs), and the sensor (for details, see Supplemental Experimental Procedures). As with $V_{\text{max}}$ in the cell expansion model (Figure 4), the relationship between the auxin treatment level and the concentration of nuclear auxin is unknown. Therefore, parameters associated with these interactions, together with the level of nuclear auxin, were estimated by fitting the model to the sensor data (Figure 6B, parameters panel). Sensor responses were measured on timescales at which de novo synthesis of auxin receptors can be assumed to be negligible.

As before, we performed a global parameter scan to approximate the 95% confidence region of the parameter estimates for the early nuclear auxin response model (Table S3). In Figure 6C, we plotted the receptor occupancy for both WT and PIN8OX cells against the concentration of exogenous NAA. In broad agreement with our quantitative analysis of the cell expansion data (Figure 4), receptor occupancy was reduced by 6-fold in PIN8OX cells when compared to WT (Figure 6C).

We next checked predictions of the model for the natural auxin IAA (Table S4). Overall, both the sensor dynamics (Figure S6E) and the resulting receptor occupancy (Figure S6F) were similar to the ones found for NAA (Figures 6A and 6C). We observed that the receptor occupancy measured for IAA saturates at a lower concentration of exogenous IAA than for NAA. This is consistent with the reported lower binding affinity of TIR1 to IAA (Dharmasiri et al., 2005; Kapinus and Leyser, 2005).

Finally, we tested our prediction that auxin enters the nucleus predominantly via the ER. We parameterized the nuclear auxin flux model (Figure 5A; Table S2) by fitting it to the receptor occupancy estimates from the sensor data. The resulting dose-response curves and the corresponding fits for NAA or IAA are shown in Figure 6C and Figure S6F, respectively. Then, as with the cell expansion model (Figure 5C), we calculated the ER-to-nucleus or cytosol-to-nucleus auxin flux ratio for each parameter set inside the 95% confidence region. This revealed strikingly similar histograms for NAA (Figure 6D) and IAA (Figure S6G) flux ratios. According to the median of the flux ratio distribution, the ER-to-nucleus NAA flux is 46-fold stronger than the cytosol-to-nucleus fluxes. This indicates an even larger flux ratio than that predicted by our analysis of the cell expansion dynamics. For IAA, the median flux ratio predicts a 44-fold stronger ER-to-nucleus flux compared to the cytosol-to-nucleus fluxes. Together, the early nuclear auxin response model confirms two key predictions made using the cell expansion dynamics data and the nuclear auxin flux models (Figures 4 and 5), namely, that
the difference in nuclear auxin signaling between PIN8OX and WT cells is established almost immediately after treatment and receptor occupancy in PIN8OX cells is maintained at a low level as exogenous auxin levels are increased. Although the predicted ER-to-nucleus flux rates derived using the sensor are higher than those predicted from the cell expansion dynamics model, both models suggest a dominant role of the ER and the ER-to-nucleus flux in regulating nuclear levels of the phytohormone auxin.

Early Nuclear Auxin Response Model Explains Opposing PIN5 Overexpression Effects

The cell expansion data did not reveal any significant effect of PIN5 overexpression (Figures 3B and 3C), which could be explained by a rapid degradation of PIN5 within 48 hr. However, short-term effects should still be present and therefore detectable with the auxin sensor. PIN5 likely transports auxin from the cytosol to the ER and therefore in the opposite direction of PIN8. This has been demonstrated with radioactive auxin transport assays (Mravec et al., 2009), which show decreased efflux (and hence increased retention) of auxin in PIN5OX cells in comparison to control. A natural question arises: What is the effect of overexpressing PIN5 on nuclear auxin fluxes?

To explore this further, we used the parameter sets sampled from the 95% confidence regions already obtained by fitting the WT and PIN8OX sensor data (Tables S3 and S4) to the early nuclear auxin response model (Figure 6C; Figure S6F) and assumed (consistent with the work of Mravec et al., 2009) that overexpression of PIN5 results in an increase of cytosol-to-ER auxin flux. As a consequence, our model predicts a decrease in cytosol levels of auxin in favor of ER ones. This results in one of two consequences for nuclear auxin uptake. In scenario 1, the increase in ER-to-nucleus flux is greater than the decrease in cytosol-to-nucleus flux, so the net effect is an increase of nuclear auxin levels (and hence increased receptor occupancy). In scenario 2, the opposite occurs, whereby the decrease in cytosol-to-nucleus flux is greater than the increase in ER-to-nucleus flux, resulting in a decrease in nuclear auxin levels (and hence decreased receptor occupancy).
To visualize this, we plotted for each parameter set the predicted gain in ER-to-nucleus flux against the loss in cytosol-to-nucleus flux (Figures 7A and 7B), where blue points correspond to scenario 1 (increase in receptor occupancy) and red points correspond to scenario 2 (decrease in receptor occupancy). Points that lie along the diagonal correspond to the cases in which increase in ER-to-nucleus flux is balanced by decrease in cytosol-to-nucleus flux, leading to no net change in the receptor occupancy and nuclear auxin levels. In WT cells overexpressing PIN5 (WT+PIN5), both scenarios are predicted to be possible, depending on a given parameter set (Figure 7A). In contrast, for PIN8OX cells overexpressing PIN5 (PIN8OX+PIN5) only scenario 1 is predicted to occur, although the magnitude of the increase in nuclear auxin could be rather low and below the level of detection (Figure 7B). Similar predictions were made independent of the concentration or type of auxin used.

We next sought to test these predictions by co-expressing the auxin sensor and PIN5 in WT or PIN8OX cells. For this, we used the early nuclear auxin response model (Figure 6) to obtain fits (within 95% confidence regions) to time-resolved IAA and NAA...
dose-response data (Figures S7A–S7D) to gain information about corresponding receptor occupancies (Figures S7E and S7F). To more clearly visualize the effect of PIN5 overexpression, we plotted (for both NAA and IAA) the difference in receptor occupancies between WT and WT+PIN5 cells (Figure 7C) and the difference in receptor occupancies between PINBOX and PINBOX+PIN5 cells (Figure S7G). Untreated WT+PIN5 cells showed increased receptor occupancy when compared to WT (Figure 7C). This is consistent with the assumption that PIN5 increases cytosol-to-ER flux, leading to an increase in ER-to-nucleus flux and hence an increase in the nuclear steady-state concentration of auxin. However, as the concentration of exogenous auxin is increased, there is clear divergence in the response: overexpression of PIN5 leads to an increase in NAA receptor occupancy, whereas for the IAA receptor, occupancy is reduced when compared to WT (Figure 7C). In contrast, overexpression of PIN5 did not greatly affect receptor occupancy in PIN8OX cells for either IAA or NAA (Figure S7G).

The response of WT+PIN5 cells to NAA (Figure 7C) corresponds to scenario 1 in Figure 7A, whereas the IAA response corresponds to scenario 2. To understand what underpins the difference between these two scenarios in the model, we performed a further inspection of the corresponding parameter sets (for details, see Supplemental Experimental Procedures and Figure S8). This revealed that for parameters corresponding to scenario 2 (which we associate with IAA) the ER-to-nucleus flux rate reaches saturation. As a consequence, overexpression of PIN5 does not lead to a significant increase in the ER-to-nucleus flux, rather to the decrease in cytosol-to-nucleus flux, which results in decreased nuclear auxin levels and hence receptor occupancy. For parameters corresponding to scenario 1 (which we associate with NAA), ER-to-nucleus flux is not saturated, so a decrease in cytosol-to-nucleus flux is compensated by the increase in ER-to-nucleus flux. Thus, the model offers a consistent explanation for opposing effects of PIN5 overexpression in WT cells upon treatment with different auxins (Figure 7C). Overall, our results propose that (1) there are two routes for auxin to enter the nucleus, one via the ER and one via the cytosol; (2) the ER-to-nucleus flux dominates over the cytosol-to-nucleus one; and (3) the ER-to-nucleus flux can be saturated at high concentrations of auxin. In effect, the ER can act as conduit to the nucleus, unless the processes controlling ER-to-nucleus flux become saturated by auxin (for example, by overexpressing PIN5), in which case, it can act as a sink.

**DISCUSSION**

In this work, we focused on unexplored aspects in auxin biology: whether and how auxin transport into the nucleus is regulated. To our knowledge, so far these questions have not arisen, likely due to the general assumption—without experimental evidence—that a small molecule such as auxin can freely diffuse from the cytosol into the nucleoplasm through the nuclear pores.

Taking advantage of auxin transport probes (Hayashi et al., 2014) we revealed unexpected intracellular dynamics of auxin that cannot be explained by simple diffusion alone. This parallels findings on the even smaller signaling molecule Ca^{2+}, which showed distinct Ca^{2+}-mediated responses in the nucleoplasm and the cytosol (Charpentier and Oldroyd, 2013). Furthermore, it has emerged that in addition to size exclusion, the biopolymer matrix within the nuclear pores might function as a selective diffusion barrier for particular molecules by employing filtering mechanisms based on hydrophobic-electrostatic interactions (Lieleg and Ribbeck, 2011). This could affect auxin diffusion through the nuclear pores. Therefore, based on the observed distribution of the NBD-NAA between the nucleus and the extra-nuclear regions, we assumed the existence of an alternative mechanism to regulate nuclear auxin levels.

One of the first attempts to model cellular auxin dynamics in the context of auxin fluxes and ionic transport among apoplast, cytosol, and vacuole was carried out by Steinacher et al. (2012). The proposed framework has led to a number of predictions that remain to be experimentally tested. Here we developed a joint theoretical-experimental approach to circumvent the lack of tools for directly estimating how and what changes in subcellular auxin pools trigger what downstream responses. We exploited the relative simplicity of single plant cell systems and demonstrated that by overexpressing ER-localized auxin efflux facilitators, efficient modulation of the ER auxin pool could be achieved and assessed. Consequences of such modulation were quantitatively monitored using microscopy and synthetic biology tools in time-resolved, dose-response assays. We used mathematical modeling to interconnect auxin-mediated cellular responses, nuclear signaling, and potential strength of nuclear auxin fluxes.

The consistency among the quantitative predictions based on the cell expansion data and the auxin-sensor-derived experimental data strongly suggests that a key step in the regulation of cell expansion is the establishment of nuclear auxin levels inside the cell. Furthermore, the data-driven mathematical models of cell expansion (Figures 4) and of the nuclear auxin fluxes (Figure 5) predicted (1) rapid (on a timescale of minutes) establishment of the difference in nuclear auxin signaling between PINBOX and WT cells after auxin treatment and (2) upon increase of exogenous auxin concentration, receptor occupancy in PINBOX cells remains lower than in WT. These predictions were confirmed using the ratiometric auxin sensor and led to our key model-driven hypothesis summarized in Figure 7D: the ER-to-nucleus flux is a dominant route for nuclear auxin uptake. We then tested our hypothesis by exploring the effect of PIN5 overexpression on WT and PINBOX cells. The model provides a consistent explanation for why PIN5 overexpression results in opposing effects, depending on the type of auxin. In particular, the model indicates that ER-to-nucleus fluxes are mediated by a saturable transport mechanism (Figure S8). This strongly implies that there must exist unidentified molecular players mediating auxin flux between the ER and the nucleus (Figure 7D) that remain to be discovered.

Although ER-residing auxin carriers (Barbez et al., 2012; Dal Bosco et al., 2012; Ding et al., 2012; Mravec et al., 2009) and enzymes involved in auxin deconjugation (Bartel and Fink, 1995; Sanchez Carranza et al., 2016) might modulate auxin levels in the ER, the predicted molecular players should be essential in regulating ER-to-nucleus auxin flux strength. Moreover, the difference in the predicted ER-to-nucleus flux strength derived from the cell expansion and early nuclear auxin response models (10-fold versus 46- to 84-fold, respectively) strongly
corroborates the relevance of other auxin-related processes in the regulation of nuclear auxin content. Upstream auxin regulatory processes, which include auxin synthesis and biochemical modifications, transport in and out of cells and within, collectively forming subcellular auxin pools that define auxin-mediated responses of plant cells during development (Moreno-Risueno et al., 2010) and interaction with the environmental stimuli (Kazan, 2013). In this context, and because auxin is a small and mobile molecule with a strong impact on plant cell responses, particularly auxin-mediated transcriptional processes, the ER can be considered a buffer for auxin between the cytosol and the nucleus, while the predicted ER-to-nucleus auxin flux provides the means for tight regulation of auxin interaction with TIR1/AFB-Aux/IAA machinery and downstream responses. We therefore believe our approach provides a basis for integration of these processes and elaboration of more comprehensive models in the future.

Our results are particularly intriguing in light of growing evidence that in plants, the ER is home to several putative hormone transporters, receptors, and enzymes that regulate homeostasis not only of auxin but possibly of cytokinin (Wulfsbæck, et al., 2011), ethylene (Bisson et al., 2009), jasmonate (Koo, et al., 2014), and other hormones. Therefore, our finding suggests a direct link between ER-based homeostatic mechanisms and nuclear auxin signaling, in which the ER plays the role of the main conduit for nuclear auxin uptake, as well as an important site for hormonal crosstalk.

**EXPERIMENTAL PROCEDURES**

Further details about experimental procedures, mathematical modeling, statistical analysis, and an outline of resources used in this work can be found in Supplemental Experimental Procedures.

**DATA AND SOFTWARE AVAILABILITY**

Code developed as part of this work can be found at https://gitlab.com/wurssb/ER_Controls_Nuclear_Auxin_Uptake. Further information on this code should be directed to A.D.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Supplemental Experimental Procedures, nine figures, and four tables and can be found with this article online at https://doi.org/10.1016/j.celrep.2018.02.074.

**ACKNOWLEDGMENTS**

The authors acknowledge Katja Rapp for technical support, Dr. Xin Yu for providing PIN5 cDNA, and Dr. Claude Becker for HTB4-mCherry construct. We thank Prof. Markus Owen for critically reading the manuscript. We gratefully acknowledge the help of Dr. Thorsten Falk for revising the AutoOvuscule code. This work was supported by Bundesministerium f"ur Bildung und Forschung (BMBF SYSBRA, SYSTEC, Microsystems), the Excellence Initiative Networking Fund of the Helmholtz Association within the Helmholtz Initiative on Synthetic Biology (SO-078) and the Alexander von Humboldt Foundation (1141629).

**AUTHOR CONTRIBUTIONS**

Conceptualization, C.F., A.D., C.D.B., A.M.M., and K.P.; Investigation, A.D., C.D.B., F.R., S.B., A.M.M., and C.F.; Formal Analysis, A.M.M., A.D., C.D.B., and C.F.; Methodology & Software, P.C., R.B., O.R., A.D., C.D.B., S.W., M.D.Z., H.H., and R.U.; Resources, W.W., M.D.Z., H.H., R.U., and K.H.; Writing – Original Draft, A.D., A.M.M., C.F., P.C., and C.D.B.; Writing – Review & Editing, A.D., A.M.M., C.F., C.D.B., K.P., K.H., M.D.Z., R.B., and O.R.; Funding Acquisition, A.D., C.F., O.R., and K.P.

**DECLARATION OF INTERESTS**

The authors declare no competing interests.

Received: September 26, 2016
Revised: December 18, 2017
Accepted: February 19, 2018
Published: March 13, 2018

**REFERENCES**

Abel, S., Nguyen, M.D., and Theologis, A. (1995). The PS-IAA4/5-like family of early auxin-inducible mRNAs in Arabidopsis thaliana. J. Mol. Biol. 251, 533–549.
Barbez, E., Kubes, M., Rolcik, J., Béziat, C., Pencik, A., Wang, B., Rosquete, M.R., Zhu, J., Dobrev, P.I., Lee, Y., et al. (2012). A novel putative auxin carrier family regulates intracellular auxin homeostasis in plants. Nature 485, 119–122.
Bartel, B., and Fink, G.R. (1995). ILR1, an amidohydrolase that releases active indole-3-acetic acid from conjugates. Science 268, 1745–1748.
Bisson, M.M., Bleckmann, A., Allekotte, S., and Groth, G. (2009). EIN2, the central regulator of ethylene signalling, is localized at the ER membrane where it interacts with the ethylene receptor ETR1. Biochem. J. 424, 1–6.
Brunoud, G., Wells, D.M., Oliva, M., Lamie, A., Mirabet, V., Burrow, A.H., Beeckman, T., Kepinski, S., Traas, J., Bennett, M.J., and Vernoux, T. (2012). A novel sensor to map auxin response and distribution at high spatio-temporal resolution. Nature 482, 103–106.
Calderón Villalobos, L.I.A., Lee, S., De Oliveira, C., Ivetc, A., Brandt, W., Armitage, L., Sheard, L.B., Tan, X., Parry, G., Miao, H., et al. (2012). A combinatorial TIR1/AFB-Aux/IAA co-receptor system for differential sensing of auxin. Nat. Chem. Biol. 8, 477–485.
Campanoni, P., and Nick, P. (2005). Auxin-dependent cell division and cell elongation. 1-Naphthaleneacetic acid and 2,4-dichlorophenoxyacetic acid activate different pathways. Plant Physiol. 137, 939–948.
Charpentier, M., and Oldroyd, G.E. (2013). Nuclear calcium signaling in plants. Plant Physiol. 163, 496–503.
Dal Bosco, C., Dovzhenko, A., Liu, X., Woerner, N., Rensch, T., Eismann, M., Eimer, S., Hegemann, J., Papnov, I.A., Ruperti, B., et al. (2012). The endoplasmic reticulum localized PIN8 is a pollen-specific auxin carrier involved in intracellular auxin homeostasis. Plant J. 71, 860–870.
Debarre, A., Muller, P., Imhoff, V., and Guern, J. (1996). Comparison of mechanisms controlling uptake and accumulation of 2,4-dichlorophenoxy acetic acid, naphthalene-1-acetic acid, and indole-3-acetic acid in suspension-cultured tobacco cells. Planta 198, 532–541.
Dharmasiri, N., Dharmasiri, S., and Estelle, M. (2005). The F-box protein TIR1 is an auxin receptor. Nature 435, 441–445.
Ding, Z., Wang, B., Moreno, I., Duplákova, N., Simon, S., Carraro, N., Reem-}

**3056** Cell Reports 22, 3044–3057, March 13, 2018
Dunlap, J.R., Kresovich, S., and McGee, R.E. (1986). The effect of salt concentration on auxin stability in culture media. Plant Physiol. 81, 934–936.

Feraru, E., Feraru, M.I., Kleine-Vehn, J., Martinière, A., Mouille, G., Vanneste, S., Vernhettes, S., Runions, J., and Friml, J. (2011). PIN polarity maintenance by the cell wall in Arabidopsis. Curr. Biol. 21, 338–343.

Freire Rios, A., Yoshida, S., and Weijers, D. (2014). Auxin regulation of embryo development. In Auxin and Its Role in Plant Development, E. Zazimalova, J. Petrasek, and E. Benkova, eds. (Springer-Verlag), pp. 171–189.

Ganguly, A., Lee, S.H., Cho, M., Lee, O.R., Yoo, H., and Cho, H.T. (2010). Differential auxin-transporting activities of PIN-FORMED proteins in Arabidopsis root hair cells. Plant Physiol. 153, 1046–1061.

Görlich, D., and Kutay, U. (1999). Transport between the cell nucleus and the cytoplasm. Annu. Rev. Cell Dev. Biol. 15, 607–660.

Hayashi, K., Nakamura, S., Fukunaga, S., Nishimura, T., Jenness, M.K., Murphy, A.S., Motose, H., Nozaki, H., Furutani, M., and Aoyama, T. (2014). Auxin transport sites are visualized in planta using fluorescent auxin analogs. Proc. Natl. Acad. Sci. USA 111, 11557–11562.

Kazan, K. (2013). Auxin and the integration of environmental signals into plant root development. Ann. Bot. 112, 1655–1665.

Kepinski, S., and Leyser, O. (2005). The Arabidopsis F-box protein TIR1 is an auxin receptor. Nature 435, 446–451.

Koo, A.J., Thireault, C., Zemelis, S., Poudel, A.N., Zhang, T., Kitaoka, N., Brandizzi, F., Matsuura, H., and Howe, G.A. (2014). Endoplasmic reticulum-associated inactivation of the hormone jasmonoyl-L-isoleucine by multiple members of the cytokrome P450 94 family in Arabidopsis. Plant Physiol. 165, 543–551.

Steinacher, A., Leyser, O., and Clayton, R.H. (2012). A computational model of auxin and pH dynamics in a single plant cell. J. Theor. Biol. 296, 84–94.

Steffens, B., and Lüthen, H. (2000). New methods to analyze auxin-induced growth II: The swelling reaction of protoplasts—a model system for the analysis of auxin signal transduction? Plant Growth Regul. 32, 115–122.

Taylor-Teeples, M., Lancot, A., and Nemhauser, J.L. (2016). As above, so below: auxin’s role in lateral organ development. Dev. Biol. 419, 156–164.

Wei, X., Henke, V.G., Strubing, C., Brown, E.B., and Clapham, D.E. (2003). Real-time imaging of nuclear permeation by EGFP in single intact cells. Biophys. J. 84, 1317–1327.

Wend, S., Dal Bosco, C., Kämper, M.M., Ren, F., Palme, K., Weber, W., Dovzhenko, A., and Zurbrüggen, M.D. (2013). A quantitative ratiometric sensor for auxin deprivation induces synchronous Golgi differentiation in suspension-cultured tobacco BY-2 cells. Plant Physiol. 117, 501–513.

Wu, M.F., Yamaguchi, N., Xiao, J., Bargmann, B., Estelle, M., Sang, Y., and Wagner, D. (2015). Auxin-regulated chromatin switch directs acquisition of flower primordium founder fate. eLife 4, e98269.

Winicur, Z.M., Zhang, G.F., and Staehelin, L.A. (1998). Auxin deprivation induces synchronous Golgi differentiation in suspension-cultured tobacco BY-2 cells. Plant Physiol. 117, 501–513.

Yoo, S.D., Cho, Y.H., and Sheen, J. (2007). Alterations in auxin homeostasis suppress defects in cell wall function. PLoS ONE 2, e98193.