ABA represses TOR and root meristem activity through nuclear exit of the SnRK1 kinase

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ABSTRACT

The phytohormone abscisic acid (ABA) promotes plant tolerance to major stresses like drought, partly by modulating plant growth and development. However, the underlying mechanisms are poorly understood. Here, we show that cell proliferation in the Arabidopsis thaliana root meristem is controlled by the interplay between three kinases, SNF1-RELATED KINASE 2 (SnRK2), the main driver of ABA signaling, the SnRK1 energy sensor, and the growth-promoting TARGET OF RAPAMYCIN (TOR) kinase. Under favorable conditions, the SnRK1α1 catalytic subunit is enriched in the nuclei of root meristematic cells and this is accompanied by normal cell proliferation and meristem size. Depletion of SnRK2s in a snrk2.2 snrk2.3 double mutant causes constitutive cytoplasmic localization of SnRK1α1 and a reduction in meristem size, suggesting that, under non-stress conditions, SnRK2s enable growth by retaining SnRK1α1 in the nucleus. In response to elevated ABA levels, SnRK1α1 translocates to the cytoplasm and this is accompanied by inhibition of TOR, decreased cell proliferation and meristem size. Blocking nuclear export with leptomycin B abrogates ABA-driven SnRK1α1 relocalization to the cytoplasm and the inhibition of TOR. Fusion of SnRK1α1 to an SV40 nuclear localization signal leads to defective TOR repression in response to ABA, demonstrating that SnRK1α1 nuclear exit is a premise for this repression. Finally, the SnRK2-dependent changes in SnRK1α1 subcellular localization are specific to the proliferation zone of the meristem, underscoring the relevance of this mechanism for growth regulation.
The phytohormone abscisic acid (ABA) plays major roles in plant stress responses. ABA signals are transduced through a well-established pathway whose main effectors in Arabidopsis are SNF1-RELATED PROTEIN KINASE 2.2 (SnRK2.2), SnRK2.3, and SnRK2.6 (1). ABA promotes plant adaptation partly by modifying developmental programs, having a major impact e.g. on root architecture (2). ABA modulates primary root (PR) and lateral root (LR) growth through interactions with other hormones, ultimately affecting cell division and elongation by poorly understood mechanisms (2). We recently uncovered an intimate connection between ABA and SnRK1 signalling that is crucial for shaping root architecture in a TARGET OF RAPAMYCIN (TOR)-dependent manner (3). TOR is a protein kinase complex that promotes cell proliferation (4). It is highly enriched in the meristems and its inactivation causes reduced root meristem size and defective PR growth (4). SnRK1 is an evolutionarily conserved protein kinase complex that is activated when energy levels decline during stress, conferring protection partly by limiting growth (5). SnRK1 is also activated by ABA, enabling plants to repress growth when water is scarce (3).

Under favorable conditions, the main catalytic subunit SnRK1α1 is sequestered by SnRK2-containing repressor complexes, allowing TOR to be active. In response to ABA, these complexes dissociate, releasing SnRK2 and SnRK1α1, which inhibits TOR and growth (3). Consistent with this model, the snrk2.2 snrk2.3 mutant (snrk2d) shows markedly defective PR growth under favorable conditions due to aberrant repression of TOR activity (3). This defect is fully rescued by the snrk1α1 mutation, demonstrating it is SnRK1α1-dependent (3).

To investigate further how TOR and root growth are controlled by SnRK2 and SnRK1, we examined the root meristems of Col-0 control seedlings, the snrk2d mutant, and its cross with snrk1α1 (snrk2d/1α1) (Fig. 1A). As previously reported (6, 7), ABA treatment reduced the number of root meristematic cells, leading to smaller meristems in Col-0. In contrast, snrk2d showed a reduction in meristem size and cell number already under mock conditions, and this was fully rescued by the snrk1α1 mutation. Consistent with the ABA hyposensitivity of snrk2d and snrk2d/1α1 (3), the addition of ABA did not further decrease meristem size or cell number in these mutants (Fig. 1A). These meristem phenotypes correlate well with the PR length previously observed in these mutants and conditions (3) and suggest that reduced cell proliferation contributes to the reduced PR length of snrk2d in mock, mimicking the situation of Col-0 plants under ABA.
We next hypothesized that regulation of SnRK1 and growth by ABA and SnRK2s could involve changes in SnRK1α1 subcellular localization. Firstly, SnRK1 and SnRK2 are highly enriched in the nuclei of root meristematic cells (3). Secondly, SnRK2-harboring SnRK1 repressor complexes localize to the nucleus in *Nicotiana benthamiana* epidermal cells (3). Thirdly, in planta, the TOR complex subunit RAPTOR1B interacts with SnRK1α1 (3, 8) in the cytosol (9). Fourthly, the subcellular localization of SnRK1α1 is central to its function (10). We therefore monitored SnRK1α1 subcellular localization in mock- or ABA-treated seedlings in two independent lines expressing SnRK1α1-GFP from its own regulatory regions (*SnRK1α1-GFP#1* and *SnRK1α1-GFP#2*). SnRK1α1 was detected in the meristematic, elongating and differentiating zones of the PR both in mock and ABA, displaying a known nuclear and cytoplasmic localization (11) (Fig. 1B). In mock, SnRK1α1 exhibited a ring-shaped pattern characteristic of nuclear proteins that are absent from the nucleolus in meristematic cells (12) (Fig. 1B). In ABA, the nuclear signal appeared to decline specifically in the meristematic zone (Fig. 1B). Electron micrographs confirmed that a ring-shaped nuclear region surrounds a large-sized nucleolus in these cells (Fig. 1C). Closer examination and GFP quantification in the meristematic zone revealed overall similar SnRK1α1-GFP levels in mock- and ABA-treated roots but a reduction in the nucleus to cytoplasm ratio (N/C) from 1.19 (line #1) and 1.2 (line #2) in mock to 0.73 (line #1) and 0.71 (line #2) in ABA (Fig. 1D), suggesting that ABA induces SnRK1α1 relocalization from the nucleus to the cytoplasm. Accordingly, the ABA effect was fully abolished by the nuclear export inhibitor Leptomycin B (LMB; N/C ratio=1.36 and 1.23 in lines #1 and #2; Fig. 1D).

Given the marked reduction in meristem size (Fig. 1A) and PR length (3) of the *snrk2d* mutant, we investigated the role of SnRK2s in SnRK1α1 localization using the cross of *snrk2d* with the *SnRK1α1-GFP#2* line (Fig. 1E). In mock, SnRK1α1-GFP was barely present in the nuclei of *snrk2d* meristematic cells, with most of the GFP signal being cytoplasmatic and yielding a N/C ratio of 0.75 (Fig. 1E), comparable to the ABA-treated *SnRK1α1-GFP#2* control (Fig. 1D). SnRK2.2-GFP suffered from overall protein degradation, but there was no change in its subcellular localization (Fig. 1F). Nuclear exit of SnRK1α1 could be observed in other highly proliferating regions of the root, such as the LR tip (Fig. 1G). However, in differentiated root cells, ABA treatment and SnRK2 depletion did not alter SnRK1α1 subcellular localization (Fig. 1H). Furthermore, the ABA-triggered changes in SnRK1α1 localization remained undetected in nucleus/cytoplasm fractions of whole roots (Fig. 1I). Altogether, these results indicate that the
SnRK2-dependent localization of SnRK1α1 and its ABA-triggered nuclear exit is specific to meristematic cells, consistent with reports that SnRK1α1 subcellular localization is controlled in a cell type-specific manner (13, 14).

Given that SnRK1α1 is required for the inhibitory effect of ABA on TOR and growth (3), we wondered whether SnRK1α1 nuclear exit is a premise for such inhibition. Indeed, pre-treatment with LMB blocked the ABA-triggered repression of TOR activity (Fig. 2A), measured as phosphorylation of ribosomal protein S6 [RPS6S240; (3)]. Moreover, as compared to control plants expressing wild-type SnRK1α1 (control-α1), the repression of TOR by ABA was also defective when SnRK1α1 was fused to an SV40 nuclear localization sequence (NLS) that favors its presence in the nucleus (NLS-α1; Fig. 2B). This demonstrates that SnRK1α1 nuclear exit is necessary for inhibiting TOR in response to ABA. Furthermore, the fact that nuclear export is crucial for repressing TOR (Fig. 2A) but only SnRK1α1 (Fig. 1D), and not SnRK2.2, translocates to the cytoplasm in response to the hormone (Fig. 1F), suggests that the previously reported role of SnRK2s in this process (3, 15) may be indirect via SnRK1α1 regulation.

We conclude that root growth is modulated by ABA through changes in SnRK1α1 subcellular localization, allowing control of TOR activity and cell proliferation in the root meristem in accordance to e.g. water availability (Fig. 2C). When conditions are favorable, SnRK1α1 is sequestered in the nucleus by SnRK2-containing repressor complexes. Dissociation of these complexes in response to ABA releases SnRK1α1, which translocates to the cytoplasm and inhibits TOR activity and growth. This mode of regulation appears to be specific to meristematic cells, where TOR is highly enriched and where active cell proliferation takes place (4). Future work will address whether such specificity stems from the exclusive ability of meristematic cells to form SnRK2-SnRK1α1 complexes, to export SnRK1α1 from the nucleus or both.

REFERENCES


**FIGURE LEGENDS**

**Figure 1.** Impact of ABA and SnRK2s on SnRK1α1 subcellular localization and cell proliferation in the root apical meristem. (A) Meristems of Col-0, snrk2d and snrk2d/1α1 7d-old seedlings with or without ABA treatment (50 µM, 48h). Arrowheads: region for cortical cell number and meristem length quantifications (violin plots). n=23-24; p < 0.05, one-way ANOVA with Tukey’s HSD test. Scale bar: 50 µm. (B) Root apices of 4d-old SnRK1α1-GFP#1 seedlings with or without ABA treatment (50 µM, 3h). Scale bar: 30 µm. Arrowheads: same position between upper and bottom photographs. (C) Electron micrograph of meristematic epidermal cells
with magnification (right panel) showing the cellular ultrastructure. C: cytoplasm; N: nucleus; No: nucleolus. Scale bars: left, 5 µm; right, 1 µm. (D-E) SnRK1α1-GFP subcellular localization in epidermal cells of 4d-old root meristems. Scale bar: 10 µm. Dotted lines: nuclear boundary.

Quantification of SnRK1α1-GFP mean cellular fluorescence and nucleus-to-cytosol (N/C) ratios. n=5-6 (D) or 8 (E); one-way ANOVA with Dunnett’s test. (D) Two independent SnRK1α1-GFP lines (#1 and #2) with or without ABA (50 µM, 3h) or LMB (2.5 µM, 1h prior to ABA addition) treatment. (E) Impact of the presence (control, SnRK1α1-GFP#2) or absence (SnRK1α1-GFP#2; snrk2d) of SnRK2s. (F) Expression and subcellular localization of SnRK2.2-GFP in epidermal cells of 4d-old root meristems treated and quantified as in (D). Scale bar: 10 µm. n=5; two-tailed Student t-test. (G-H) SnRK1α1-GFP subcellular localization in LRs (G) and the PR differentiation zone (H) in seedlings (Line #1 and/or #2) with or without ABA treatment (50 µM, 3h) and in absence (SnRK1α1-GFP#2; snrk2d) of SnRK2s. Scale bar: 30 µm. (G) Nine-day old seedlings. (H) Four-day old seedlings. N/C ratios of differentiated root cells. n=5, two-tailed Student t-test (line #1) or one-way ANOVA (line #2). PI, propidium iodide. ns, non-significant. (I) Representative SnRK1α1 immunoblot from nuclear and cytoplasmic fractions of 14d-old Col-0 whole roots with or without ABA (50 µM ABA, 3h). H3, Histone 3 marker for nuclear fractions.

**Figure 2. Impact of SnRK1α1 subcellular localization on TOR signaling and root growth.** (A-B) Representative immunoblots and quantification of RPS6\(^{5240}\) phosphorylation (phospho-RPS6/total-RPS6) in Col-0 (A) or control-α1 and NLS-α1 (B) seedlings with or without ABA (50 µM, 3h) or LMB (2.5 µM, 1h prior to ABA addition). Images from the same gel were cropped for showing α1 and NLS-α1 contiguously. (A) n=5; (B) n= 3; error bars, SEM; two-tailed Student t-test. (C) Under favorable conditions, SnRK1α1 is sequestered in the nucleus by repressor complexes containing SnRK2 [and a PP2C (3)], enabling TOR activity in the cytoplasm, meristematic cell proliferation and root growth. Dissociation of these complexes in ABA by the hormone-bound PYR/PYL/RCAR receptors releases SnRK1α1 which exits the nucleus and inhibits TOR and growth. TORC1, TOR Complex 1; N, nucleus; C, cytoplasm. Created with BioRender.com.

**Materials and Methods**

Experimental details are provided in SI Appendix.
All study data are included in the paper.

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Author contributions

BBP performed experiments and analyzed data. MC performed the root meristem assays and analyzed the data. TB and FR contributed tools and expertise and supported conceptual work. EBG and BBP conceived the project, designed experiments, analyzed and interpreted data and wrote the manuscript.

Competing interests

The authors declare no competing interests.
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