Strigolactone elevates ethylene biosynthesis in etiolated Arabidopsis seedlings

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ABSTRACT

The gaseous phytohormone ethylene influences many aspects of plant life, including germination, fruit ripening, senescence, and stress responses. These diverse roles of ethylene occur in part through crosstalk with other phytohormones, which affects ethylene biosynthesis and signaling pathways. We have recently shown that the phytohormones, including gibberellic acid, abscisic acid, auxin, and methyl jasmonate, regulatory the stability of ACC synthases (ACSs), the rate-limiting enzymes in ethylene biosynthesis. Here, we report that treatment of etiolated Arabidopsis seedlings with strigolactone (SL) increases ethylene biosynthesis. SL does not influence ACS stability or ACS gene expression, but it increases the transcript levels of a subset of ACC oxidase (ACO) genes, thereby enhancing ethylene biosynthesis. Taken together with the results of our previous study, these findings demonstrate that most phytohormones differentially regulate ethylene biosynthesis in dark-grown Arabidopsis seedlings by affecting ACS stability and/or the transcript levels of ethylene biosynthesis genes.

The gaseous plant hormone ethylene influences a diverse array of plant growth, development, and biotic or abiotic stress responses through its crosstalk with other phytohormones and signaling pathways. ACC synthases (ACSs) and ACS oxidase (ACO, C-terminal of ACS6). Furthermore, the degradation of ACS7 is hammering in PP2C type-A knockout plants, suggesting the role of PP2Cs in ACS stability regulation. Similarly, a protein phosphatase 2A ROOTS CURL IN 1-N-NAPHTHYLPHTHALAMIC ACID 1 (RCN1), promotes the degradation of a subset of ACS proteins. By contrast to the positive role of phosphorylation on ACS stability, CASEIN KINASE 1.8-mediated phosphorylation of ACS5, ACS9, and ACS6 promotes the degradation of these ACS isoforms. Compared to ACS, less is known about the role of post-translational modification on ACO stability. However, recent studies have shown that the activity of ACO is also regulated by post-translational modification. Sulfhydration is one such regulation that decreases the activity of tomato ACO1 and ACO2 via the sulfhydration of cysteine residues of the ACO isoforms, demonstrating the role of post-translational control on ACO in plants.

In addition to phosphorylation, the degradation of ACS via the ubiquitin-proteasome pathway is another common regulatory mechanism that controls ethylene biosynthesis. ETHYLENE-OVERPRODUCING 1 (ETO1) and its two paralogous ETO1-LIKE 1 (EOL1) and EOL2 specifically target type-2 ACS for degradation via the 26S proteasome pathway. ETO1 contains a BTB/TRP motif and serves as an adaptor protein for the CULLIN3 (CUL3) E3 ligase complex and imparts substrate specificity to CUL3. Type-1 ACS stability is also regulated by the 26S proteasome, but the cognate E3 ligases have not yet been identified. Unlike the type-1 and type-2 ACSs, type-3 ACS does not contain any known regulatory motifs or phosphorylation sites. However, the stability of ACS7 is regulated by a RING-type
E3 ligase XBAT32 and ABI1, a protein phosphatase 2 C, demonstrating that type-3 ACS stability is also under the control of phosphorylation and ubiquitination.16,24

Ethylene biosynthesis appears to be regulated by many external and internal stimuli. However, surprisingly, only a handful of stimuli have been identified and characterized that govern the stability of ACS.1,4 Phytohormone is one of the stimuli that influence the stability of ACS. For example, cytokinins and brassinosteroids (BRs) have long been known as positive regulators of ACS stability.1,4 We have also previously demonstrated a similar regulation of ACS stability by other phytohormones, including gibberellic acid (GA), abscisic acid (ABA), auxin, methyl jasmonate (MJ), and salicylic acid (SA), with distinct effects on different ACS isoforms.1 For instance, SA increases the stability of type-2 ACS, whereas it promotes the degradation of type-1 ACS. MJ and ABA increase type-2 ACS stability, but neither hormone has any effect on the stability of type-1 ACS. Furthermore, none of these phytohormones influences type-3 ACS stability. It is worth noting that the insensitivity of the stability of the type-3 ACS7 isoform to most phytohormones may be related to the nature of the N-terminal-tagged fusion of ACS7 protein used in the study, which blocks the degradation sequence located in the N-terminal domain.25 Further studies on the role of phytohormones on the C-terminal fusion of ACS7 will provide more insights into the phytohormone-mediated regulation of ACS stability.

Strigolactones (SLs), a group of carotenoid-derived phytohormones, were initially characterized as root-derived compounds that stimulate the germination of the seeds of parasitic weeds such as Striga, Orobanche, and Phelipanche species.26–28 SLs also play a pivotal role in plant growth and developmental processes, including the determination of root and shoot architectures, flower development, and leaf senescence.28 Many of these processes are associated with crosstalk between SLs and other phytohormones, including ethylene. For example, leaf senescence is strongly accelerated by the combination of SL with ethylene, not by SL alone, suggesting that SL stimulates leaf senescence through the action of ethylene. Conversely, ethylene treatment enhances SL biosynthesis by increasing expression of the strigolactone biosynthesis genes MORE AXIALLY GROWTH3 (MAX3) and MAX4 during leaf senescence.29 SL and ethylene also regulate root hair elongation. The study showed that ethylene is epistatic to SL and ethylene biosynthesis is required for SL-mediated root hair elongation.30 Roots treated with the ethylene biosynthesis inhibitor AVG do not elongate even in the presence of GR24, a synthetic SL analog, while GR24 alone increases root hair elongation. These results indicate that ethylene plays a positive role in SL biosynthesis. However, a possible role of SL on the biosynthesis of ethylene remains to be established.

We did not examine the role of SL as regulators of the stability of ACS in dark-grown seedlings in our previous study.1 In the present study, we expanded our analysis by examining the effect of SL on the stability of ACS using dark-grown Arabidopsis seedlings expressing Myc-tagged ACS proteins. Similar to several other phytohormones in the previous study, treatment of GR24 increased ethylene biosynthesis in dark-grown wild-type and seedlings expressing Myc-ACS2 (type-1), Myc-ACS5 (type-2), or MycACS7 (type-3). The levels of ethylene in wildtype and Myc-ACS overexpression lines increased in proportion to the increased concentration of GR24 (Figure 1a). We also examined whether the GR24-mediated increase in ethylene biosynthesis is due to an increase in the abundance of ACS proteins by determining the steady-state levels of ACS. An increased concentration of GR24 did not affect the steady-state levels of all three types of ACS proteins (Figure 1b). Furthermore, GR24 did not affect ACS gene expression, regardless of treatment duration (Figure 2). These results indicated that ACS is not the target.

![Figure 1](image_url) GR24 enhances ethylene biosynthesis in dark-grown Arabidopsis seedlings but does not influence ACS stability. (a) Seedlings were grown in capped vials containing growth media with the indicated concentration of GR24 for 3 d and the accumulated ethylene was measured by gas chromatography. Error bars indicate SD; n = 3. (b) Seedlings expressing myc-ACS2, myc-ACS5, or myc-ACS7 were grown on growth media with the indicated concentrations GR24 for 3 d. Total protein extracts from the seedlings were analyzed by immunoblotting using an anti-Myc or an anti-HSC70 antibody as a loading control. The Myc-ACS bands were normalized to the HSC70 control, and these values were then normalized to the “no GR24” control value, which was set to 1. The numbers below the gel panels indicate the relative ratios of the band intensities. * P < .05, Student’s t-test.
of GR24 in the ethylene biosynthesis pathway, although it is possible that GR24 may control the activity of ACS activity via post-translational modification. Furthermore, the involvement of other ACS isoforms that have not been examined in this study may attribute to the GR24-mediated increase in ethylene production. Further study will answer these questions. In contrast to our result, a previous study showed that SL enhances ACS2 gene expression in Arabidopsis roots.\(^\text{30}\) This discrepancy may result from differences in the experimental conditions, including the growth conditions (light-grown vs dark-grown), the concentration of SL, and the treatment duration.

We additionally investigated whether ACS might be a target of SL as increased ACS transcript levels or ACS stability also leads to an increase in ethylene biosynthesis. Unlike its effect on the ACS genes, GR24 increased the transcript levels of a subset of ACS genes (ACS1, ACS2, and ACS4) approximately 2–4 fold within 2 to 4 h after the treatment (Figure 3). By contrast, no changes were observed in the transcript levels of the other ACS genes (ACS3 and ACS5) although a slight decrease in ACS5 transcript was observed at 4 h after the treatment (Figure 3). These results demonstrated that the GR24-mediated increase in ethylene biosynthesis was a result of an increase in ACS gene expression, although the possibility remains for GR24-mediated stabilization of ACS proteins.

The observation of downregulation of ACS transcripts, with no effect on ACS stability, by SL was reminiscent of the effects of MJ and ABA, although these hormones downregulate ACS transcripts.\(^\text{1}\) However, unlike MJ and ABA, which showed different effects on the stability of type-1 and type-2 ACS, SL did not influence the stability of any of the three ACS types in dark-grown seedlings. Taken together, these results demonstrated that...
SL regulates ethylene biosynthesis by enhancing the transcript levels of the ACO genes without influencing ACS stability. Elucidation of the effects of phytohormones on ACS stability in light-grown seedlings or in response to stress conditions will provide more insight into the differential role of most phytohormones on the regulation of ethylene biosynthesis.

Materials and methods

Plant materials and growth conditions

Arabidopsis thaliana Col-0 was used as the wildtype (WT) reference throughout the study. All plants were grown in either long-day or short-day conditions at 22°C ± 2°C or in vitro on Murashige and Skoog (MS) Basal Medium supplemented with 0.8% plant agar (pH 5.7) in a continuous light chamber at 21°C. Transgenic Arabidopsis lines used were previously generated as 35S CaMV constitutive promoter-driven dexamethasone (Dex)-inducible Myc-tagged ACS2, ACS5, or ACS7. Briefly, the coding region of each ACS gene was cloned into the Gateway pENTR vector and transferred to a modified Gateway-compatible version of the binary GVG vector pTA7002 to which a 6-35S cassette was added. Col-0 plants were transformed by the floral dip method and transformants were selected on MS medium containing hygromycin.

Treatment of seedlings with GR24 for protein analysis and RT-PCR

For protein analysis, seedlings expressing Myc-ACS2, Myc-ACS5, or Myc-ACS7 were grown on growth medium containing 20, 40, or 5 nM Dex in the presence or absence of GR24 (PhytoTechnology Laboratories) for 3 d in the dark. For quantitative RT-PCR analysis, 3-day-old dark-grown wild type seedlings were treated with 5 μM GR24 for the indicated times and harvested for analysis.

Protein analysis

Proteins were extracted from 3-day-old etiolated seedlings in 2× SDS buffer (65.8 mM Tris HCl, pH 6.8, 26.3% Glycerol (w/v), 2.1 % SDS, 0.01% Bromophenol blue) and ground with a mini-pestle. Total protein extracts were incubated at 95°C for 3 min and then centrifuged for 3 min at room temperature at 16 000 g in a microcentrifuge. Immunoanalysis was performed using an anti-Myc antibody (Sigma-Aldrich) and HSC70 (Enzo) as a loading control.

RNA extraction and quantitative RT-PCR

Total RNA was prepared using the RNeasy Plant Mini Kit (QIAGEN) and reverse transcribed using SuperScript II reverse transcriptase (Invitrogen) according to the manufacturers’ instructions. Quantitative RT-PCR was performed using PowerUP SYBR Green Master Mix (Applied Biosystems). The primers used are listed in Table S1. Three biological replicates were analyzed with three technical replicates per sample. The relative expression of the candidate genes was normalized to β-tubulin.

Measurements of ethylene production

Ethylene measurements were performed as previously described. Surface-sterilized seeds were germinated in 22 mL gas chromatography vials containing 3 mL of a medium consisting of half-strength MS medium/1% sucrose/0.8% Bacto agar, Dex, and GR24. After 3 d of stratification at 4°C in darkness, the vials containing Arabidopsis seeds were exposed to light for 3 h, capped, incubated in dark for 3 d and the accumulated ethylene was measured by gas chromatography using a Shimadzu GC2010 Plus capillary gas chromatography system with an HS-20 headspace autosampler. Ethylene concentration was calculated in pL•seedling•3 days. All genotypes and treatments were measured from three biological replicates and an average and SD were calculated.

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Author contribution

GMY conceived the experiments; HYL conducted most experiments; GMY and HYL wrote the manuscript.

Data availability

All data discussed in the paper are available in the main text and Supplementary Information.

Disclosure of potential conflicts of interest

No potential conflicts of interest were disclosed.

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Supporting information

Supplemental Table 1. The list of primers used in this study.

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