Dual Mechanisms of Coniferyl Alcohol in Phenylpropanoid Pathway Regulation

Mengling Guan1,2, Changxuan Li2, Xiaotong Shan1, Fang Chen3,4, Shufang Wang5, Richard A. Dixon3,4,5 and Qiao Zhao1,2*

1 Shenzhen Key Laboratory of Synthetic Genomics, Guangdong Provincial Key Laboratory of Synthetic Genomics, Shenzhen Institute of Synthetic Biology, Shenzhen Institute of Advanced Technology, Chinese Academy of Sciences, Shenzhen, China, 2 Center for Plant Biology, School of Life Sciences, Tsinghua University, Beijing, China, 3 Department of Biological Sciences, University of North Texas, Denton, TX, United States, 4 BioDiscovery Institute, University of North Texas, Denton, TX, United States, 5 Beijing Advanced Innovation Center for Tree Breeding by Molecular Design, Beijing Forestry University, Beijing, China

Lignin is a complex phenolic polymer that imparts cell wall strength, facilitates water transport and functions as a physical barrier to pathogens in all vascular plants. Lignin biosynthesis is a carbon-consuming, non-reversible process, which requires tight regulation. Here, we report that a major monomer unit of the lignin polymer can function as a signal molecule to trigger proteolysis of the enzyme L-phenylalanine ammonia-lyase, the entry point into the lignin biosynthetic pathway, and feedback regulate the expression levels of lignin biosynthetic genes. These findings highlight the highly complex regulation of lignin biosynthesis and shed light on the biological importance of monolignols as signaling molecules.

Keywords: phenylpropanoid pathway, coniferyl alcohol, KFB01/20/39/50 expression, PAL stability, signaling molecule

INTRODUCTION

The phenolic polymer lignin, along with the polysaccharides cellulose and hemicellulose, are the major components of the secondary cell wall in vascular plants. Lignin is essential for providing strength and hydrophobicity to secondary cell walls. It is synthesized from deamination of the amino acid L-phenylalanine (Phe), which also serves as precursor of numerous specialized metabolites (Bonawitz and Chapple, 2010), by L-phenylalanine ammonia-lyase (PAL). A series of aromatic hydroxylations, O-methylations and side-chain reductions then lead to the formation of the three major lignin monomers (monolignols) p-coumaryl alcohol, coniferyl alcohol (CA) and sinapyl alcohol (Bonawitz and Chapple, 2010). After transport across the plasma membrane, most likely by a passive diffusion mechanism (Perkins et al., 2022; Zhuo et al., 2022), monolignols are oxidatively polymerized by laccases and peroxidases in the apoplastic space (Lee et al., 2013; Zhao et al., 2013).

The biosynthesis of lignin consumes approximately 30% of photosynthetically fixed carbon and is not reversible (Boerjan et al., 2003). Therefore, carbon resources that are channeled through Phe toward the biosynthesis of lignin must be tightly controlled to ensure that the Phe pool remains sufficient for protein synthesis. The regulation of lignin biosynthesis occurs at transcriptional, translational and post-translational levels (Taylor-Teeples et al., 2015; Zhao, 2016; Chezem et al., 2017; Geng et al., 2020). It has been well-documented that most lignin biosynthetic
genes contain the so-called AC elements, including AC-I (ACCTACC), AC-II (ACCAACC), and AC-III (ACCTAAC) in the promoters that are recognized by MYB transcription factors (Raes et al., 2003). In Arabidopsis, MYB58 and MYB63 are the major transcription activators essential for lignin biosynthesis (Zhou et al., 2009). Recently, MYB20, MYB42, MYB43, and MYB85 have been identified as novel transcriptional activators that not only directly regulate the expression of the lignin biosynthetic genes but also the genes involved in the production of phenylalanine, which serves as precursor of lignin biosynthesis (Geng et al., 2020). Members of subfamily 4 of the R2R3-MYB family have been reported to function as transcriptional repressors of lignin biosynthesis. AmMYB308 and AmMYB330 in Arabidopsis, EgMYB1 in Eucalyptus gunnii and ZmMYB31 and ZmMYB42 in maize have all been shown to repress lignin biosynthetic genes by directly binding to AC cis-elements (Tamagnone et al., 1998; Jin et al., 2000; Formalé et al., 2010).

Plant natural product biosynthesis is known to feature metabolite-mediated regulation. For instance, most amino acid biosynthetic pathways are subjected to tight feedback regulation (Galili et al., 2016). In the aromatic amino acids (AAA) biosynthesis pathway, the enzyme mediating the entry reaction is regulated by the final products (Maeda and Dudareva, 2012). Recently it has been reported that disruption of DHS1 and DHS3 caused hypersensitivity to tyrosine and tryptophan, respectively, while the dhs2 mutant was resistant to tyrosine-mediated growth inhibition (Yokoyama et al., 2021). This suggests that the AAA biosynthesis pathway is regulated by complex metabolite-mediated mechanisms. However, the underlying molecular nature of the regulation is not clear.

Although it is important for plants to tightly control lignin biosynthesis to avoid overconsumption of Phe, how plants monitor lignin biosynthesis remains elusive. It is intuitive that the regulation of lignin biosynthesis might involve sensing of lignin-specific precursors. It was recently shown that CA biosynthesis can be maintained in the absence of guaiacyl or syringyl lignin biosynthesis (Zhuo et al., 2022), suggesting that CA may serve roles beyond simply being a lignin building block. In this study, we explored the function of CA in the regulation of lignin biosynthesis. We showed that CA treatment rapidly induces the expression of the genes encoding Kelch repeat F-box proteins, which mediate PAL stability. In addition, the expression of the lignin biosynthetic genes was subjected to feedback regulation by CA. Together, these data suggested that the major monolignol CA can regulate lignin biosynthesis in multiple ways.

**MATERIALS AND METHODS**

**Plant Materials and Growth Conditions**

All Arabidopsis (Arabidopsis thaliana) lines used were in the Col-0 ecotype background. The kfb01/20/39/50 quadruple mutant was isolated from the progeny of a cross between the previously described kfb01/20/50 (Zhang et al., 2015) and kfb39. The kfb01/20 and kfb39/50 were isolated from the progeny of a cross between Col-0 and kfb01/20/39/50. All materials used in this study including Col-0 and mutants were grown on half-strength (1/2) Murashige and Skoog (MS) medium containing 1.5% sucrose and 0.35% Phytagel (pH 5.8) at 22°C under long-day conditions with a 16-h light/8-h dark cycle.

To investigate the effects of monolignols and vanillyl alcohol (VA) on A. thaliana, 2-week-old seedlings were treated in 1/2 MS liquid medium containing 0.1 mM monolignol or 0.1 mM VA (with an equal volume of solvent as the control) for different time periods before qRT-PCR analysis and PAL stability assay.

**Plasmid Construction**

The CaMV 35S promoter and the coding sequence of YFP were amplified and cloned into pCAMBIA1300 to generate 35Spro:YFP, and then the full-length coding sequences of KFB01, KFB20, KFB39, and KFB50 without the stop codon were separately amplified and cloned into 35Spro:YFP to obtain 35Spro:KFB01−YFP, 35Spro:KFB20−YFP, 35Spro:KFB39−YFP, and 35Spro:KFB50−YFP constructs. To obtain the truncated KFBs−YFP, the DNA fragments encoding KFB01−35, KFB20−8, KFB39−50, and KFB50−31 were separately amplified and cloned into 35Spro:YFP.

To obtain the UBQ10pro:PAL1−HA, UBQ10pro:PAL2−HA, UBQ10pro:PAL3−HA and UBQ10pro:PAL4−HA, the UBQ10 promoter was cloned into pEarlyGate 301 to generate the UBQ10pro:HA construct, and then the coding sequences of PAL1, PAL2, PAL3, and PAL4 without the stop codon were separately amplified and cloned into UBQ10pro:HA. For Co-IP assay, the KFBs−YFP and PALs−HA constructs were separately transformed into Agrobacterium tumefaciens strain GV3101, which was used to transiently express proteins in Nicotiana benthamiana leaves by the injection method.

**Monolignol Metabolite Quantification**

To determine the monolignol metabolites, 3-week-old Arabidopsis seedlings were separated into leaf and root samples, and inflorescence stems were collected from plants that had just started to flower. Tissues were harvested and ground in liquid nitrogen. The samples (~120 mg) were extracted with 100% methanol (500 mL) in a cold room for 1 h, vortexed briefly and extracted at room temperature for another hour. The liquid phase was separated by centrifugation at 13,000 g for 10 min and passed through a 0.45 mm minifilter before LCMS analysis. For each sample, materials from about 15 plants were pooled. Two biological replicates were processed for each tissue. The strategy for monolignol quantification was as previously described (Cocuron et al., 2019). Specifically, phenolic compounds were detected and quantified using an Agilent 1290 Infinity II liquid chromatography system coupled to a hybrid Triple Quadrupole 6500 + triple quadrupole from AB Sciex. The extracts were kept at 4°C in an auto-sampler. The metabolites were resolved at 30°C using a reverse phase C18 Symmetry column (4.6 × 75 mm; 3.5 μm) associated with a Symmetry C18 pre-column (3.9 × 20 mm; 5 μm) from Waters.
The liquid chromatography gradient was made of 0.1% (v/v) acetic acid in acetonitrile (A) and 0.1% (v/v) acetic acid in water (B). The total LC-MS/MS run was 15 min with a flow rate of 800 µL/min. The following gradient was applied to separate the phenolic compounds: B = 0–1 min 85%, 1–7 min 42%, 7–9 min 20%, 9–9.1 min 15%, 9.1–12 min 15%, 12–12.1 min 85%, 12.1–15 min 85%. The injection needle was rinsed with 50% aqueous methanol. Five µL of extracts or external standard mixture was injected onto the column.

The detection of monolignols was accomplished using an AB Sciex hybrid Triple Quadrupole/Ion trap mass spectrometer QTRAP 6500+. Electrospray ionization with polarity switch was utilized to acquire mass spectra of the phenolic compounds. In this study, negative ionization (−4500 V) was applied to p-coumaraldehyde, p-coumaryl alcohol, sinapyl alcohol, whereas positive ionization (5000 V) was used for coniferaldehyde, CA, and sinapaldehyde. The settling time between each polarity was 15 ms. Metabolites were simultaneously detected as precursor ion/product ion pair using multiple reaction monitoring (MRM): coniferaldehyde (m/z 179 > 147); CA (m/z 163 > 131); p-coumaraldehyde (m/z 147 > 119); p-coumaryl alcohol (m/z 149 > 131); sinapaldehyde (m/z 209 > 177); sinapyl alcohol (m/z 209 > 194). The source parameters such as curtain gas (40 psi), temperature (550°C), nebulizer gas (50 psi), heating gas (60 psi), and collision activated dissociation (Medium) were kept constant during MRM. The dwell time in the mass spectrometer was set to 20 ms. Analyst 1.7 software from AB Sciex was used to record and process the LC-MS/MS data. Monolignols were identified and quantified using a mixture of known external standards run at the same time as the biological extracts.

RNA Analysis
For the RNA-seq analysis, 2-week-old Arabidopsis seedlings of ecotype Col-0 were separately treated with 0.1 mM CA (or EtOH control) for 15, 30 min, 1 and 4 h, and then the seedlings from each treatment were divided into three biological replicates. The total RNA of each sample was extracted using the RNeasy MinElute Cleanup Kit (74204, Qiagen, Germany) before sending to BGI Genomics (The Beijing Genomics Institute) for RNA-seq analysis.

For qRT-PCR analysis, total RNA was isolated using TRIzol™ Reagent (15596018, Invitrogen, United States) and treated with DNase I (AMPD1-1KT, Sigma-Aldrich, United States). Complementary DNA (cDNA) was obtained using SuperScript™ III Reverse Transcriptase (18080093, Invitrogen, United States). The qRT-PCR reaction was set with TB Green™ Premix Ex Taq™ II (RR820L, Takara, China). GAPDH was used as the control gene for normalization. The primers used for qRT-PCR analysis are listed in Supplementary Table 1.

For RT-PCR analysis, quantitative copies of the total RNA extracted from 2-week-old seedlings of Col-0 and kfb01/20/39/50 were reverse-transcribed. The RT-PCR was performed with Taq DNA Polymerase (AP111-02, Transgen, China), and TUA4 was used as the control gene. Primers are listed in Supplementary Table 1.
RESULTS

Exogenous Coniferyl Alcohol Inhibits the Growth of Arabidopsis Seedlings
To determine the endogenous amount of lignin monomers, we quantified the three free forms of monolignols and their corresponding aldehydes in Arabidopsis. Although the aldehydes were barely detectable, p-coumaryl alcohol and CA accumulated to significant levels, approximately 7 nmol/g fresh weight for CA in stem tissue (Table 1). The lower levels of sinapyl alcohol may be due to the formation of other sinapate derivatives such as sinapylmalate, sinapoylgucose, etc. in Arabidopsis (Fraser and Chapple, 2011).

To investigate the possibility of the lignin monomers as signal molecules, we first treated the Arabidopsis seedlings with coniferyl alcohol (CA) to explore whether the exogenous monolignol affected plant growth. Five-day-old Arabidopsis seedlings were transplanted to 1/2 MS solid medium containing different concentrations of CA, and allowed to grow for 10 days before observing the phenotype. Compared to the different concentrations of CA, and allowed to grow for 10 days before observing the phenotype. Compared to the 0.1 mM CA treatment, and differentially expressed genes with FDR < 0.001 and fold-change ≥ 2 or ≤ 0.5 were identified. Compared with the ethanol control, 1124, 1243, 1188, and 1353 genes were up-regulated respectively, while 1139, 1195, 1122, and 3269 genes were down-regulated after 15, 30 min, 1, and 4 h treatment of CA respectively (Supplementary Data Sets 1–4). CA could specifically induce the expression of lignin biosynthesis transcriptional repressors MYB4 and MYB7 after only 15 min treatment, while the transcript levels of the lignin biosynthesis genes PAL1, 4-coumarate:CoA ligase 1 (4CL1), hydroxycinnamoyl transferase (HCT), caffeoyl shikimate esterase (CSE), ferulate 5-hydroxylase (F5H), caffeic acid 3-O-methyltransferase (COMT) and cinnamyl alcohol dehydrogenase (CAD4) were all significantly decreased at 4 h after exposure to CA (Figures 1A,B). The results suggested that CA treatment caused the down-regulation of lignin biosynthesis genes by inducing the expression of MYB4/7.

The Expression of KFB01/20/39/50 Rapidly Responds to Coniferyl Alcohol
Interestingly, genes encoding the Kelch repeat F-box (KFB) proteins KFB01/20/39/50 that regulate the protein stability of PAL (Zhang et al., 2013, 2015) were also rapidly induced by CA treatment (Figures 1C–F). The transcript levels of KFB39 was significantly induced after as soon as 10 min, and induced by 30-fold after 30 min treatment. The expression of other KFB genes was induced to a lesser extent (Figure 1C). To determine the specificity of CA induction of the KFB genes, seedlings were also treated with vanillyl alcohol (VA), which has a similar chemical structure to CA but with a shortened side-chain (Supplementary Figure 2). qRT-PCR analyses showed that the transcripts of KFB genes were induced as early as 10 min by CA but not VA treatment or solvent control (Figures 1C–F). We also compared the ability of other monolignols (p-coumaryl alcohol, caffeyl alcohol, and sinapyl alcohol) to induce KFB39 transcripts. p-Coumaryl alcohol and sinapyl alcohol significantly induced KFB39 transcripts, but caffeal alcohol, the precursor of C-lignin only reported to date in some seed coats (Chen et al., 2012), did not (Supplementary Figure 3). This result suggests that all the classical monolignols found in vegetative tissues are signaling molecules of lignin biosynthesis through induction of the KFB genes.

To test if the induction of the KFB genes by CA treatment is dependent on MYB4 and/or its homologs MYB7 and MYB32, the triple mutant myb4/7/32 was treated with CA or dimethyl sulfoxide (DMSO) as control. KFB gene transcript levels were enhanced in myb4/7/32 by CA treatment (Figures 1G–J), indicating that CA induction of the KFB genes is independent of MYB4 and its homologs.

The Mediator complex subunits MED5a and MED5b have been shown to be involved in the regulation of lignin biosynthesis and also in the expression of the KFB genes (Bonawitz et al., 2014; Kim et al., 2020; Wang et al., 2020). The transcript levels of KFB39 and KFB50 were significantly decreased in the med5 mutant. To determine whether MED5 is involved in monolignol signaling, the med5a/5b and ref4-3 (a semidominant point mutation in MED5a) were treated with CA. KFB transcripts were not induced in med5a/5b and KFB39 transcripts were increased only weakly in ref4-3 (Figures 1G–J). Therefore, we conclude that MED5a

| Metabolite            | Leaf (pmol g⁻¹ fresh weight) | Root (pmol g⁻¹ fresh weight) | Stem (pmol g⁻¹ fresh weight) |
|-----------------------|-----------------------------|-------------------------------|-----------------------------|
| p-Coumaraldehyde      | 0.64 ± 0.06                 | 32.41 ± 0.65                  | 8.81 ± 1.84                 |
| p-Coumaryl alcohol    | 662.29 ± 13.62              | 28460.42 ± 907.87             | 5702.78 ± 166.44            |
| Coniferaldehyde       | 302.34 ± 23.00              | 267.12 ± 3.28                 | 195.596 ± 9.52              |
| Coniferyl alcohol     | 938.89 ± 83.29              | 28773.57 ± 1151.53            | 6974.56 ± 369.49            |
| Sinapaldehyde         | 3.47 ± 1.03                 | 11.11 ± 4.53                  | 12.64 ± 0.96                |
| Sinapyl alcohol       | 45.61 ± 19.50               | 142.25 ± 23.47                | 1087.56 ± 24.83             |

Values shown are mean ± SD from two biological replicates (n = 2).
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FIGURE 1 | Coniferyl alcohol acts as a signal molecule to regulate the phenylpropanoid pathway. **(A)** The expression of MYB4/7 was induced by coniferyl alcohol (CA). The transcript levels of MYB4/7 were assayed in 2-week-old Col-0 seedlings treated with solvent DMSO or 0.1 mM CA for 15 min. **(B)** Relative transcript levels of lignin biosynthetic genes are decreased after CA treatment. The transcript levels of PAL1, 4CL1, HCT, CSE, FSH, COMT, and CAD4 were assayed in 2-week-old Col-0 seedlings treated with solvent DMSO or 0.1 mM CA for 4 h. qRT-PCR assays showing the expression of KFB01 **(C)**, KFB20 **(D)**, KFB39 **(E)**, and KFB50 **(F)** in Col-0 seedlings. **(C–F)** Two-week-old Col-0 seedlings were separately treated with solvent DMSO, 0.1 mM VA (vanillyl alcohol) or 0.1 mM CA for different time intervals. qRT-PCR assays showing the expression of KFB01 **(G)**, KFB20 **(H)**, KFB39 **(I)**, and KFB50 **(J)** in response to CA in Col-0, rel4-3, med5a/5b and myb4/7/32 seedlings. **(G–J)** Two-week-old seedlings were treated with solvent DMSO or 0.1 mM CA for 15 min. **(K)** The induction of KFB39 expression by CA does not rely on proteolysis. The transcript level of KFB39 after 1 h treatment with 0.1 mM CA with or without 60 µM MG132 in 2-week-old Col-0 seedlings. All data represent average transcript level values (±SD) relative to DMSO-treated plants. Asterisks indicates a statistically significant difference between the treated and DMSO control groups (Student’s t-test, *P < 0.05, **P < 0.01, n = 3).
and MED5b are essential for the expression of KFB genes, even though they may not be required for CA sensing.

The Induction of KFB39 Expression by Coniferyl Alcohol Does Not Require Proteolysis

26S Proteasome-mediated proteolysis is a predominant route for protein degradation (Vierstra, 2009). To test whether the induction of KFB39 expression by CA is dependent on proteolysis, we performed qRT-PCR analysis on 2-week-old Col-0 seedlings after 1 h treatment with 0.1 mM CA with or without 60 µM MG132 (26S proteasome inhibitor). The results showed that the expression of KFB39 was significantly induced by CA, while the addition of MG132 had no significant effect on the induction (Figure 1K), indicating that the induction of KFB39 expression by CA does not require proteolysis.

The Induction of KFB39 Expression by Coniferyl Alcohol May Not Rely on Protein Phosphorylation or Dephosphorylation

The above studies showed that the expression of KFB39 can be rapidly induced by CA, the mechanism of which is still unclear. Protein phosphorylation is involved in many signal transduction processes by brassinosteroids, cytokinins, etc. (Hwang et al., 2012; Manghwar et al., 2022). In order to determine whether the induction of KFB39 by CA relied on phosphorylation modification of some transcription factors, we conducted a phosphorylation proteome study on the nuclear-localized proteins from the 2-week-old Col-0 seedlings treated with solvent DMSO or 0.1 mM CA for 1 h. The results showed that the phosphorylation of the threonine residue at position 183 of the transcription factor PLATZ (AT1G32700) disappeared after the seedlings were treated with CA for 1 h, and dephosphorylated PLATZ cannot be enriched by binding to TiO2 beads, while the phosphorylated PLATZ from DMSO treated seedlings can be enriched in this way (Figure 2A). In addition, data on the website ATTED-II v111 show that PLATZ is co-expressed with KFB01, KFB20, KFB39 and KFB50 (Obayashi et al., 2022). In order to investigate whether PLATZ is involved in CA-mediated KFB39 induction we ordered a mutant line for PLATZ (Figure 2B), and examined the expression level of KFB39 in the 2-week-old Col-0 and platz seedlings after 30 min of solvent DMSO or 0.1 mM CA treatment. The results showed that the induction of KFB39 expression by CA was not significantly different between Col-0 and platz (Figure 2C). Thus we conclude that the induction of KFB39 expression by CA does not rely on changes in the protein phosphorylation status of PLATZ. However, this does not rule out the possibility that phosphorylation is involved in the CA signal transduction.

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 FIGURE 3 | Exogenous CA can specifically induce the degradation of PAL proteins in Arabidopsis. (A) Exogenous CA can induce the degradation of PAL proteins in Col-0 but not kfb01/20/39/50. (B) Exogenous VA has no effect on the stability of PAL proteins in Col-0 and kfb01/20/39/50. For (A,B), 2-week-old seedlings were separately treated with solvent DMSO, 0.1 mM CA or 0.1 mM VA in 1/2 MS liquid medium for 1, 3, and 6 h, and the solvent DMSO was used as control. Ponceau staining of RbcL was shown as loading control. (C,D) Quantitative analysis of the band intensity in (A,B). Protein contents were normalized to the corresponding content of RbcL by Ponceau staining. The relative level of PAL (CA/DMSO%) at 0 min was set to 100%. Error bars are means ± SD (Student’s t-test, **P < 0.01, n = 3).

possibility that could be studies by future quantitative proteomic studies of CA-responsive proteins.

L-Phenylalanine Ammonia-Lyase Interact With KFB01/20/39/50
It has been previously reported that KFB01/20/39/50 physically interact with PALs (Zhang et al., 2013, 2015). In order to fully confirm the protein-protein interaction between PAL1/2/3/4 and KFB01/20/39/50, we conducted Co-IP assays using N. benthamiana leaves to transiently express the proteins. The full-length KFBs-YFP and PALs-HA were transiently expressed, and we found that PAL1 and PAL4 can interact with the four KFBs (Supplementary Figures 4A,D). Owing to the weak interaction between PAL2 and PAL3 with the KFBs, we constructed the truncated KFBs (KFBs-ΔF) without the F-box motif in KFBs, and found that PAL2 and PAL3 can interact with the four truncated KFBs (Supplementary Figures 4B,C). Due to the larger than expected molecular weight of KFB50-YFP in the Western blot, mass spectrometry analysis was performed. The results showed that the band was indeed KFB50-YFP, and the covalent modification by the tobacco protein polyubiquitin 11-like (Gene ID: 107766892) caused the larger molecular weight than expected (Supplementary Table 2). It can therefore be concluded that the PAL1/2/3/4 interact with the four KFBs.

Exogenous Coniferyl Alcohol Affects the Stability of L-Phenylalanine Ammonia-Lyase Mainly Through KFB39 and KFB50
It has been reported that KFB01, KFB20, KFB39, and KFB50 redundantly control PAL stability (Zhang et al., 2015). To generate the quadruple mutant, the previously published kfb01/20/50 triple mutant and the T-DNA mutant line for KFB39 were crossed, and RT-PCR analysis confirmed that the expression of all four KFBs was disrupted in kfb01/20/39/50 (Supplementary Figure 5).

Because of the rapid up-regulation of KFBs by exogenous CA, we hypothesized that exogenous CA may trigger PAL degradation. To test this possibility, an anti-PAL antibody was developed against the two peptide sequences that are conserved among the four PAL proteins in Arabidopsis (Supplementary Figure 6A), and the 2-week-old Col-0, pal2/3/4, PAL3-GFP/Col-0 and kfb01/20/39/50 seedlings were used to detect and prove the specificity of the antibody (Supplementary Figure 6B). Then
FIGURE 4 | Exogenous CA induces the degradation of PAL proteins mainly through KFB39 and KFB50. (A) The protein level of PAL in Col-0, kfb01/20/39/50, kfb01/20, and kfb39/50 seedlings after CA treatment. Two-week-old Col-0, kfb01/20/39/50, kfb01/20, and kfb39/50 seedlings were treated with 0.1 mM CA for 3 h and the solvent DMSO was used as control. Ponceau staining of RbcL was shown as loading control. (B) Quantitative analysis of the band intensity in (A). Protein contents were normalized to the corresponding content of RbcL by Ponceau staining. The relative level of PAL (CA/DMSO%) at 0 min was set to 100%. Error bars are means ± SD (Student’s t-test, **P < 0.01, n = 3).

the level of immuno-detectable PAL protein was compared in extracts of 2-week-old Col-0 and kfb01/20/39/50 seedlings after 1, 3, and 6 h of CA, VA or solvent DMSO treatment. The results showed that the protein level of PAL in Col-0 seedlings decreased after 3 and 6 h of CA treatment, but remained stable after VA treatment, while the protein level of PAL in kfb01/20/39/50 seedlings remained stable after CA or VA treatment (Figure 3).

Since the expression of KFB39 was affected by CA induction the most, and KFB01 to a lesser extent, we generated the kfb01/20 and kfb39/50 double mutant based on the homology relationship of the four KFBs (Zhang et al., 2015). Then the protein level of PAL was determined in the 2-week-old Col-0, kfb01/20/39/50, kfb01/20 and kfb39/50 seedlings after 3 h of solvent DMSO or 0.1 mM CA treatment. The results showed that the protein level of PAL in the Col-0 and kfb01/20 seedlings decreased to approximately 40%, but remained unchanged in kfb01/20/39/50 and kfb39/50 seedlings after 3 h of CA treatment (Figure 4). Therefore, CA-induced PAL degradation mainly depends on KFB39 and KFB50.

Coniferyl Alcohol Has No Effect on the Interaction Between Kelch Repeat F-box and L-Phenylalanine Ammonia-Lyase

It has been reported that ligands can promote the interaction between F-box receptors and their substrates, such as the receptor of auxin TIR1 (Dharmasiri et al., 2005; Kepinski and Leyser, 2005). In order to investigate whether the affinity between KFBs and PAL proteins is affected in the presence of the small molecule CA, we carried out in vitro GST pull-down assays. PAL1-His, GST-KFB20 and GST-KFB39 were separately purified from E. coli host, and GST pull-down assays with different concentrations of CA were conducted. The results showed that the affinity between KFB20 and KFB39 with PAL1 remained unchanged under the treatment of different concentrations of CA (Figure 5), indicating that CA has no effect on the interaction between KFBs and PAL proteins.

Mutants With Compromised Coniferyl Alcohol Biosynthesis Have Higher L-Phenylalanine Ammonia-Lyase Accumulation

Based on the above-mentioned experimental results, we speculated that the content of CA in plants has impact on the protein level of endogenous PAL. Cinnamyl alcohol dehydrogenases (CADs) are required for the reduction of aldehydes to monolignols, and the disruption of CAD4/5 results in weakly reduced CA in Arabidopsis (Jourdes et al., 2007; Anderson et al., 2015). UGT72E2 encodes a glycosyltransferase which catalyzes the formation of monolignol glucosides from monolignols, and the CA content in UGT72E2 OE
is slightly lower than that in Col-0 (Lanot et al., 2006; König et al., 2014). C3H encodes p-coumaroyl shikimate 3’ hydroxylase, and ref8 with compromised C3H activity exhibits sharply reduced lignin content compared to Col-0 (Kim et al., 2011; Bonawitz et al., 2014). We examined the relative content of endogenous PAL in 2-week-old Col-0, cad4/5, UGT72E2 OE and ref8-2 seedlings. As shown in Figure 6, the protein level is slightly elevated in UGT72E2 OE, but at similar level in cad4/5. In sharp contrast, PAL protein accumulation is significantly higher in ref8-2. These data suggest that plants with reduced CA biosynthesis have higher PAL protein content.

**DISCUSSION**

In this study, we have shown that CA, the major monolignol, can function as a signal molecule to regulate the phenylpropanoid pathway in multiple ways. CA treatment can rapidly induce the expression of the genes encoding Kelch repeat F-box proteins, which mediate PAL stability. On the other hand, the expression of the lignin biosynthetic genes was feedback-regulated by CA, and this may be caused by CA-induced MYB4/7.

**KFB01/20/39/50 Play Different Roles in Regulating L-Phenylalanine Ammonia-Lyase Protein Stability**

It has been previously reported that KFB01, KFB20, KFB39, and KFB50 all interact with PALs and mediate the protein stability. However, protein sequence alignment suggests that KFB01 and KFB20 are closer than the other two (Zhang et al., 2015). The response of KFB39 expression to CA is much more rapid and striking compared to the other KFBs (Figures 1C–F). As shown in Figure 4, KFB39 and KFB50 are the more important in terms of CA-mediated PAL degradation, whereas CA-induced KFB01/20 contributes little to PAL degradation mediated by CA. This may be due to the fact that the expression levels of these KFBs are quite different after CA treatment, especially KFB39, although it is not clear why the induction of KFB01, KFB20, KFB39 and KFB50 is different.

**Coniferyl Alcohol-Induced MYB4/7 Play Important Roles in Regulating the Phenylpropanoid Pathway**

MYB4 and its homolog MYB7 play important roles in the regulation of the phenylpropanoid pathway in Arabidopsis, and many other plant species (Jin et al., 2000; Shen et al., 2012; Fornalé et al., 2014). Here we show that the expression of MYB4/7 can be rapidly and sustainably induced by exogenously applied CA, leading to inhibition of the expression of the lignin biosynthesis genes PAL1, 4CL1, HCT, CSE, F5H, COMT, and CAD4 after 4 h of CA treatment. The response of the lignin biosynthetic genes to CA treatment is much less rapid compared to that of KFB genes, suggesting that the response of the lignin biosynthetic genes is indirect.

**Sensing Coniferyl Alcohol Could Be a Way to Balance Phenylalanine Deamination With Protein Synthesis**

L-Phenylalanine ammonia-lyase catalyzes the first step in phenylpropanoid metabolism in which phenylalanine undergoes deamination to yield trans-cinnamic acid. Phenylalanine-derived
metabolites can constitute up to 30% carbon fixed by photosynthesis. Even though it is well-documented that lignin biosynthesis is subject to tight regulation, how plants sense monolignol production is not clear. We showed that exogenous CA treatment can enhance the expression of KFB39 by 30-fold in 30 min (Figure 1). This leads to PAL degradation and theoretically shifts phenylalanine distribution to protein synthesis. Interestingly, ref8-2 has much more PAL accumulation compared with wild type controls (Figure 6). We have found that exogenous phenylalanine treatment partially rescued the growth defects of the mutant (data not shown). It is possible that the growth defects of ref8-2 are due to overconsumption of phenylalanine as a consequence of compromised PAL degradation.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author/s.

AUTHOR CONTRIBUTIONS

QZ conceived the study. QZ, MG, FC, and RD designed the experiments. MG, CL, XS, FC, and SW performed the experimental work. QZ, MG, and RD wrote the manuscript. All authors contributed to the article and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpls.2022.896540/full#supplementary-material

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