

Mutation of the Inducible *ARABIDOPSIS THALIANA* CYTOCHROME P450 REDUCTASE2 Alters Lignin Composition and Improves Saccharification^{1[W][OPEN]}

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ARABIDOPSIS THALIANA CYTOCHROME P450 REDUCTASE1 (ATR1) and ATR2 provide electrons from NADPH to a large number of CYTOCHROME P450 (CYP450) enzymes in *Arabidopsis thaliana*. Whereas ATR1 is constitutively expressed, the expression of ATR2 appears to be induced during lignin biosynthesis and upon stresses. Therefore, ATR2 was hypothesized to be preferentially involved in providing electrons to the three CYP450s involved in lignin biosynthesis: CINNAMATE 4-HYDROXYLASE (C4H), *p*-COUMARATE 3-HYDROXYLASE1 (C3H1), and FERULATE 5-HYDROXYLASE1 (F5H1). Here, we show that the *atr2* mutation resulted in a 6% reduction in total lignin amount in the main inflorescence stem and a compositional shift of the remaining lignin to a 10-fold higher fraction of *p*-hydroxyphenyl units at the expense of syringyl units. Phenolic profiling revealed shifts in lignin-related phenolic metabolites, in particular with the substrates of C4H, C3H1 and F5H1 accumulating in *atr2* mutants. Glucosinolate and flavonol glycoside biosynthesis, both of which also rely on CYP450 activities, appeared less affected. The cellulose in the *atr2* inflorescence stems was more susceptible to enzymatic hydrolysis after alkaline pretreatment, making ATR2 a potential target for engineering plant cell walls for biofuel production.

Economic and environmental concerns have stimulated the production and use of renewable materials and fuels. Plant cell wall biomass has the potential to become a renewable feedstock for processing into chemicals and liquid fuels in the biorefinery (Vanholme et al., 2013b). Plant cell walls contain polysaccharides (mainly cellulose and hemicelluloses) that can be

hydrolyzed into fermentable sugars via enzymatic saccharification. However, the presence of lignin in the cell wall severely hampers the saccharification process (Chen and Dixon, 2007; Van Acker et al., 2013). As a consequence, lignin must be removed, or at least partially cleaved, using biomass pretreatments that require chemicals and energy, prior to saccharification. These pretreatments contribute a significant cost to the production of cell wall-based biofuels in the biorefinery. Strategies to alter lignin content or composition are promising for reducing the need for and/or costs of pretreatments (Sticklen, 2006; Chen and Dixon, 2007; Weng and Chapple, 2010; Fu et al., 2011; Jung et al., 2013; Van Acker et al., 2013, 2014).

Lignin is a polymer derived from three differentially ring-methoxylated 4-hydroxycinnamyl alcohols called monolignols. The two major monolignols incorporated into the lignin polymer in angiosperms are coniferyl alcohol and sinapyl alcohol, which give rise to guaiacyl (G) and syringyl (S) units, respectively (Fig. 1). Lignin also incorporates small amounts of *p*-coumaryl alcohol, which gives rise to *p*-hydroxyphenyl (H) units. A number of other hydroxyphenylpropanoids, and other phenolic monomers, can be incorporated into the lignin polymer as well (Baucher et al., 1998; Boerjan et al., 2003; Ralph et al., 2004; Vanholme et al., 2012a). The structural genes involved in lignin biosynthesis have been rather well elucidated, at least in model species

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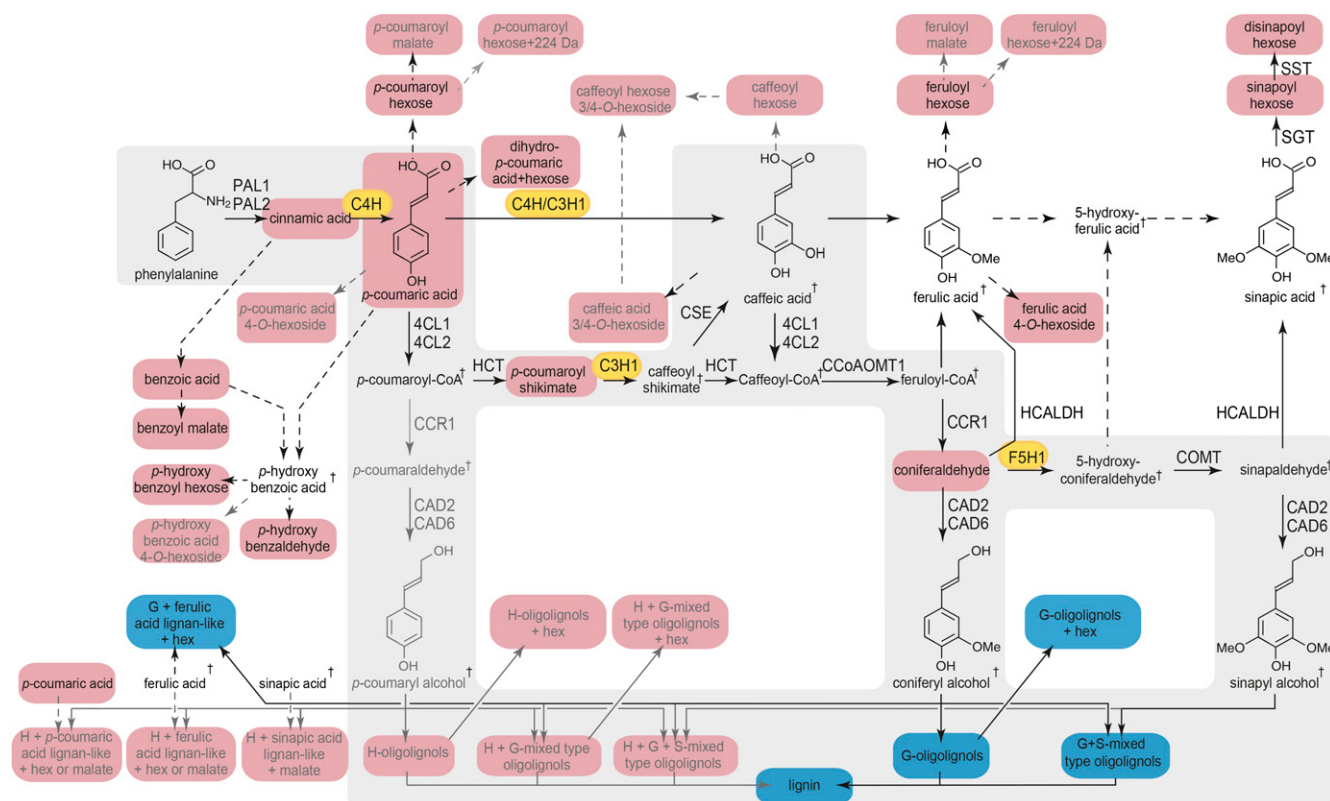


Figure 1. General phenylpropanoid- and monolignol-specific pathways and metabolic map of the changes in phenolic metabolism in *atr2* mutant stems. The main route toward lignin (i.e. the general phenylpropanoid- and monolignol-specific pathway) is marked with a gray background. Metabolites that accumulate in *atr2* mutants are marked in red, and those that decrease in *atr2* mutants are marked in blue. Metabolic routes that are active in stems of wild-type plants are given in black. Metabolic routes of which the metabolites are near or below the detection limit in wild-type plants but detected in *atr2* mutants are given in gray. Metabolites that are below the detection limit in both wild-type and mutant plants are indicated with daggers. The three CYP450s are marked in orange. PAL, PHENYLALANINE AMMONIA LYASE; C4H, CINNAMATE 4-HYDROXYLASE; 4CL, 4-COUMARATE:COENZYME A LIGASE; HCT, *p*-HYDROXYCINNAMOYL-COENZYME A:QUINATE SHIKIMATE *p*-HYDROXYCINNAMOYLTRANSFERASE; C3H, *p*-COUMARATE 3-HYDROXYLASE; CSE, CAFFEYL SHIKIMATE ESTERASE; CCoAOMT, CAFFEYL-COENZYME A *O*-METHYLTRANSFERASE; CCR, CINNAMOYL-COENZYME A REDUCTASE; F5H, FERULATE 5-HYDROXYLASE; COMT, CAFFEIC ACID *O*-METHYLTRANSFERASE; CAD, CINNAMYL ALCOHOL DEHYDROGENASE; HCALDH, HYDROXYCINNAMALDEHYDE DEHYDROGENASE; SGT, SINAPATE 1-GLUCOSYLTRANSFERASE; SST, SINAPYLGLUCOSE:SINAPYLGLUCOSE SINAPYLTRANSFERASE. For each enzymatic step of the general phenylpropanoid- and monolignol-specific pathways, specific gene family members are given that largely contribute to lignification in the inflorescence stem (Sibout et al., 2005; Vanholme et al., 2012b).

such as *Arabidopsis* (*Arabidopsis thaliana*), poplar (*Populus* spp.), and alfalfa (*Medicago sativa*; Bonawitz and Chapple, 2010; Vanholme et al., 2010a), although additional genes are still being discovered (Berthet et al., 2011; Vanholme et al., 2012b, 2013c; Zhao et al., 2013; Petrik et al., 2014).

Starting with Phe, monolignols are synthesized via the general phenylpropanoid- and monolignol-specific pathways (Fig. 1). Up to three hydroxylation steps that are catalyzed by specific CYTOCHROME P450 (CYP450) oxidases occur on the aromatic ring; these are the 4-, 3-, and 5-hydroxylations catalyzed by C4H, C3H1, and F5H1 (Fig. 1; Meyer et al., 1996; Franke et al., 2002; Schillmiller et al., 2009). C4H, C3H1, and F5H1 belong to the class II type CYP450s. Members of this class rely

on CYTOCHROME P450 REDUCTASE (CPR) activity to transfer electrons from NADPH either directly to the CYP450s or via CYTOCHROME B5s (CB5s; Ilan et al., 1981; Hannemann et al., 2007; Jensen and Møller, 2010; Im and Waskell, 2011). Animals and yeast carry only one CPR gene, whereas higher plants often have two or more (Ro et al., 2002), possibly because of the relatively higher number of CYP450 enzymes in plants. Both CYP450s and CPRs are generally accepted to be localized at the membrane of the endoplasmic reticulum (they occur in a 15:1 ratio in rat liver microsomal membranes; Shephard et al., 1983), although a number of CPR-requiring CYP450s have been predicted and/or shown to be localized in the chloroplast (Ro et al., 2002; Bak et al., 2011).

Arabidopsis has 244 CYP450 genes and 28 CYP450 pseudogenes (Bak et al., 2011). In contrast, only three CPR genes have been identified in the Arabidopsis genome. These CPRs are annotated as *ARABIDOPSIS THALIANA* P450 REDUCTASE1 (*ATR1*), *ATR2*, and *ATR3* (Urban et al., 1997; Varadarajan et al., 2010). Of these three genes, *ATR1* and *ATR2* encode genuine CPR proteins that share 64% amino acid identity, according to The Arabidopsis Information Resource genome annotation version 10. *ATR3* has been described as a putative CPR but shares only 25.7% and 24% amino acid identity with *ATR1* and *ATR2*, respectively. Unlike *ATR1* and *ATR2*, recombinant *ATR3* was unable to support human CYP1A2, and there is no experimental evidence to date that it can support CYP450s (Varadarajan et al., 2010). In addition, Arabidopsis has two genes coding for CYTOCHROME B5 REDUCTASE (CBR) proteins. CBR1 is located at the membrane of the endoplasmic reticulum but is mainly involved in electron transport for the biosynthesis of polyunsaturated fatty acids, whereas CBR2 is in the mitochondrial membrane, and there is no evidence yet that it could take part in the endoplasmic reticulum electron transport system (Wayne et al., 2013).

ATR1 and *ATR2* catalyze the same reaction and can reduce C4H in yeast and baculovirus expression systems with the same efficiency (Urban et al., 1997; Mizutani and Ohta, 1998). Despite this observation, there is clear evidence that *ATR1* and *ATR2* have distinct functions in the plant. For example, *ATR1* is constitutively expressed, whereas *ATR2* is induced by wounding and light (Mizutani and Ohta, 1998). In addition, the expression of *ATR2*, but not that of *ATR1*, is correlated with the expression of a number of genes from the phenylpropanoid pathway; microarray analysis of Arabidopsis rosette leaves subjected to cold treatment under high-light conditions displayed a coordinated expression of *ATR2* with phenylpropanoid biosynthetic genes such as *PAL2*, *CCR1*, and *CCoAOMT1* (Soitamo et al., 2008). Also, transcript profiling of Arabidopsis stems during fiber differentiation showed that expression of *ATR2*, in contrast to that of *ATR1*, is coregulated with a set of genes of the general phenylpropanoid- and monolignol-specific pathways, including the three CYP450 genes *C4H*, *C3H1*, and *F5H1* (Ehlting et al., 2005).

Given that the ATR enzymes may need to serve up to 244 different CYP450 enzymes in Arabidopsis, it is surprising that one of them, *ATR2*, is coexpressed with the lignin biosynthetic genes. This coexpression suggests that *ATR2* provides electrons preferentially to the three CYP450s of the phenylpropanoid and monolignol biosynthetic pathways, making *ATR2* a good target for lignin modification. To test this hypothesis, we studied the consequences of the down-regulation of *ATR2* in Arabidopsis on phenolic metabolism, lignin structure, and saccharification potential. As we will show, our data confirm that *ATR2* has a role in the lignin biosynthesis pathway and that *ATR2* mutation reduces the electron donor activity to the three CYP450s involved in lignin biosynthesis, as indicated by the shift in the phenolic

profile and lignin composition. Additionally, the *atr2* mutant biomass has improved saccharification efficiency.

RESULTS

ATR2 Is Coexpressed with Lignin Biosynthetic Genes

We have previously investigated the system-wide responses of lignin biosynthesis perturbation in Arabidopsis (Vanholme et al., 2012b). To this end, a total of 20 mutants, each mutated in a single lignin biosynthetic gene (two mutant alleles for 10 genes), were investigated with transcriptomics and metabolomics (Vanholme et al., 2012b). This enabled the identification of genes that were coexpressed with known lignin biosynthetic genes and, therefore, that are good candidates to be involved in lignin biosynthesis. Two of the candidates, *TRANSALDOLASE2* and *CSE*, were pathway genes subsequently proven via reverse genetics to be involved in lignification (Vanholme et al., 2012b, 2013c). Reevaluation of the transcriptome data showed that both *ATR1* and *ATR2* followed a similar expression profile to those of lignin biosynthetic genes at various stages of wild-type Arabidopsis inflorescence stem development (Fig. 2). Moreover, it is apparent that *ATR2*, but not *ATR1*, is coexpressed with lignin biosynthetic genes in the different lignin biosynthesis mutants (Fig. 2). The potential coexpression of *ATR2* and *ATR1* with lignin biosynthetic genes was further investigated by quantitative reverse transcription (qRT)-PCR in Arabidopsis seedlings treated with isoxaben. This herbicide impairs cellulose biosynthesis, which in turn induces stress-related processes such as ectopic lignification (Caño-Delgado et al., 2003; Duval and Beaudoin, 2009). Wild-type seedlings were grown in liquid cultures for 3 d and subjected to treatment with isoxaben for 6, 24, and 72 h. qRT-PCR analysis showed that the expression levels of *PAL1*, *C4H*, *C3H1*, and *HCT*, which are genes involved in general phenylpropanoid biosynthesis, and *ATR2* were significantly induced after 24 h of isoxaben treatment (Fig. 3). More specifically, a similar increase in expression was observed for *PAL1* and *ATR2* (i.e. the response was already significant after 6 h of treatment). In contrast, *ATR1* expression was not affected by isoxaben treatment (Fig. 3). These data again show that *ATR2* is coexpressed with phenylpropanoid biosynthetic genes.

Growth and Development of *atr2* Mutants

To further investigate the role of *ATR2* during lignin biosynthesis, we isolated two insertion mutants from the SALK collection, SALK_152766 and SALK_026053 (Alonso et al., 2003). The transfer DNA (T-DNA) insertions in the *ATR2* genes were validated by PCR, and the corresponding mutant alleles were designated *atr2-1* and *atr2-2*, respectively (Fig. 4). Allele *atr2-2* contains a T-DNA in the third intron of *ATR2*, whereas *atr2-1* contains a T-DNA in the 12th exon (Fig. 4). Next, *ATR2* expression

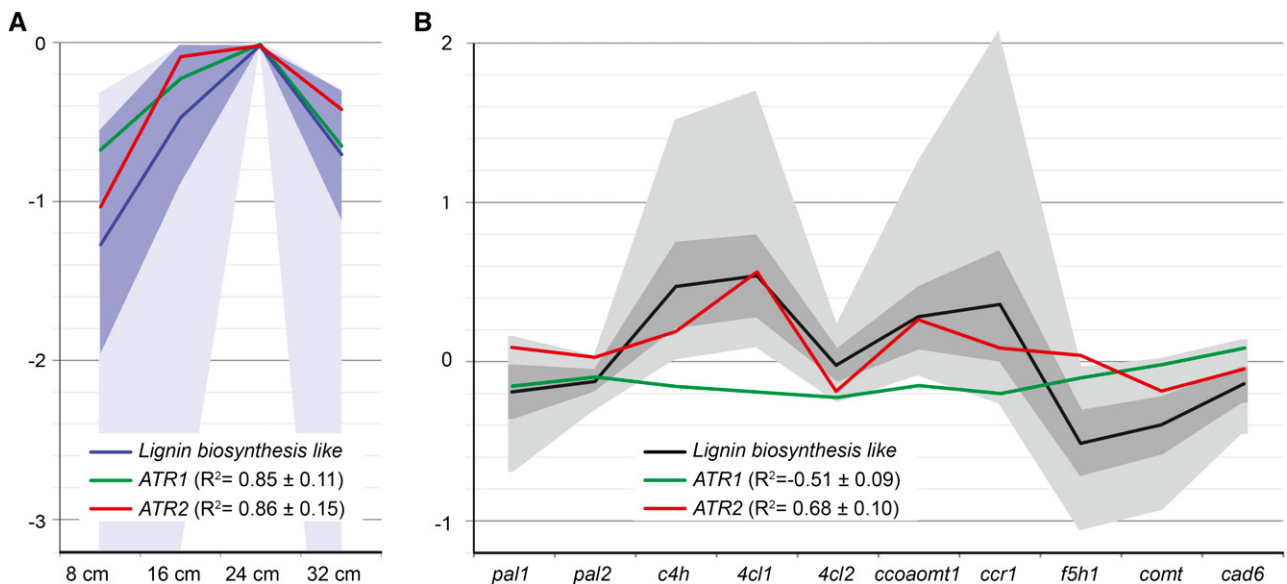


Figure 2. Expression of *ATR2* follows that of lignin biosynthetic genes (evaluation of data from Vanholme et al. [2012b]). A, Both *ATR1* and *ATR2* are coexpressed with lignin biosynthetic genes during inflorescence stem development (at an inflorescence stem height of 8, 16, 24, and 32 cm). The blue line, dark-blue area, and light-blue area indicate average, SD, and spread of the expression, respectively, of the 1,760 probes with a lignin biosynthesis-like expression profile over development (Vanholme et al., 2012b, figure 3, cluster C). The green line indicates expression of *ATR1*, and the red line indicates expression of *ATR2*. R^2 values are average correlation coefficients $\pm$ SD of *ATR1* and *ATR2* with each of the 1,760 probes with a lignin biosynthesis-like expression profile. B, *ATR2*, but not *ATR1*, is coexpressed with genes involved in lignin biosynthesis in lignin mutant backgrounds. The black line, dark gray area, and light gray area indicate average, SD, and spread of the expression, respectively, of the 62 genes with a lignin biosynthesis-like expression profile in the lignin mutants (Vanholme et al., 2012b, figure 9). The green line indicates expression of *ATR1*, and the red line indicates expression of *ATR2*. R^2 values are average correlation coefficients $\pm$ SD of *ATR1* and *ATR2* with each of the 62 genes with a lignin biosynthesis-like expression profile (Vanholme et al. 2012b).

levels in inflorescence stems were determined via qRT-PCR with two sets of primers. Primer set 1 spanned the *atr2-1* T-DNA insertion position, and primer set 2 spanned the *atr2-2* T-DNA insertion position. The *ATR2* transcript levels in *atr2-1* and *atr2-2* mutants were below the detection limit when measured with the respective T-DNA-spanning primers, proving that the native *ATR2* mRNA was absent or close to being absent (Fig. 4). However, residual *ATR2* expression was detected in *atr2-1* mutants using primer set 2 and in *atr2-2* mutants using primer set 1, indicating that residual truncated mRNA fragments of *ATR2* were present in the *atr2* mutants. The expression levels of *ATR1*, the constitutive *CPR*, remained unaltered in both mutant lines, demonstrating that the reduction in *ATR2* transcript levels was not compensated by an increased *ATR1* expression (Fig. 4).

To investigate whether mutation of *ATR2* had any effect on growth, the mutants were grown in soil under short-day conditions for 8 weeks and subsequently transferred to long-day conditions. Short-day conditions lead to the development of large vegetative rosettes, which allow the development of a large inflorescence stem upon a shift to long-day conditions. The final height of the *atr2-1* mutant was not significantly different from that of the wild type, whereas *atr2-2* was 10% smaller at final height (Fig. 5). Analysis of the final

dry weight of the main stem revealed 19% and 17% reductions for *atr2-1* and *atr2-2*, respectively, compared with the wild type (Fig. 5).

In order to examine potential changes in the morphology of the vascular tissues caused by the mutated *ATR2*, transverse sections from the bottom part of stems from fully grown but not yet senesced plants were investigated via microscopy. The morphology of the vasculature as well as the lignin was visualized with Wiesner and Mäule staining and via autofluorescence. Neither

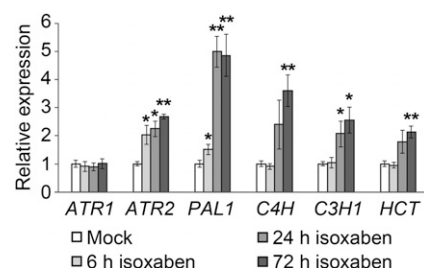


Figure 3. qRT-PCR analysis of wild-type seedlings either untreated or treated with 80 nM isoxaben for 6, 24, and 72 h. Data represent mean values of four biological pools of 20 seedlings each, and error bars represent SE. The relative expression in mock conditions was set to 1 for each gene tested. * $0.05 > P > 0.01$, ** $P < 0.01$.

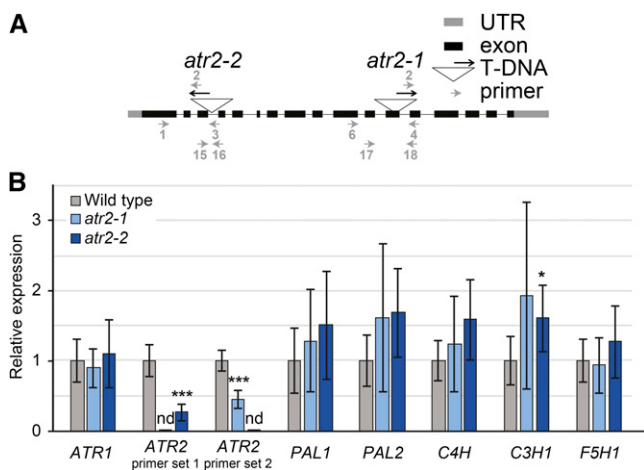


Figure 4. Characterization of *atr2* T-DNA insertion mutants. A, Representation of the *ATR2* genomic structure. Exons are represented as black boxes and introns as gray lines. The T-DNA insertion sites for the two alleles *atr2-1* and *atr2-2* are indicated by triangles. The orientation of each T-DNA is indicated by a black arrow pointing in the direction of the left border. The primers used to confirm the mutants are represented as gray arrows. The numbers of the primers correspond to those given in "Materials and Methods." UTR, Untranslated region. B, Expression analysis of *ATR1*, *ATR2*, *PAL1*, *PAL2*, *C4H*, *C3H1*, and *F5H1* in *atr2-1* and *atr2-2* mutants and the wild type as determined via qRT-PCR. Two primer pairs were used to determine the expression of *ATR2*. Primer sets 1 (i.e. primers 17 and 18) and 2 (i.e. primers 15 and 16) spanned the *atr2-1* and *atr2-2* T-DNA insertion positions, respectively. For each gene tested, the relative expression was normalized to the one of the wild type. nd, Signal below detection limit. * $0.05 > P > 0.01$, *** $P < 0.001$.

collapsed xylem vessels, nor any obvious changes in the intensity of either Mäule or Wiesner staining or autofluorescence, was observed (Fig. 5).

atr2 Mutants Show Altered Levels of Phenylpropanoids, Benzenoids, Lignan-Like Molecules, and Oligolignols

To assess the consequences of the mutation of *ATR2* at the metabolic level, phenolic profiling of methanol-soluble metabolites was performed via ultra-HPLC-mass spectrometry (MS) on inflorescence stems from plants that were around 24 cm tall. This procedure enables the detection of several classes of aromatic compounds, including metabolites of the lignin biosynthetic pathway and derivatives thereof, as well as glucosinolates and flavonols. A total of 1,067 peaks were integrated in the chromatograms of *atr2-1*, *atr2-2*, and wild-type samples (Supplemental Table S1). Principal component analysis showed that *atr2* mutant samples separated from wild-type samples based on a combination of the first and second principal components (Supplemental Fig. S1), indicating metabolic differences between the *atr2* mutants and wild-type plants. Univariate statistical analysis was applied to select peaks with significantly different abundances in the *atr2* mutants as compared with their levels in wild-type

plants. Using stringent filters (i.e. $P < 0.01$, a 2-fold difference in abundance between mutant and the wild type, and a peak intensity of greater than 500), 71 and 73 peaks were found to be significantly lower in abundance in *atr2-1* and *atr2-2*, respectively, of which 61 were in common (Supplemental Fig. S1; Supplemental Table S2); 184 and 229 peaks were found to be significantly higher in abundance in *atr2-1* and *atr2-2*, respectively, of which 177 were in common (Supplemental Fig. S1; Supplemental Table S2). The relatively high fraction of differential peaks that were shared by both mutants indicates that most of the detected differential peaks can be attributed to the defective *ATR2* gene.

Further analysis showed that the 61 peaks that were significantly lower in abundance in both *atr2* mutants could be assigned to 47 compounds (Supplemental Table S3), of which 30 could be structurally characterized based on their mass-to-charge ratio (m/z), retention time, and tandem mass spectrometry (MS/MS) fragmentation spectrum (Table I; Supplemental Fig. S2). Twenty-six of these 30 characterized compounds belonged to the classes of oligolignols, hexosylated oligolignols,

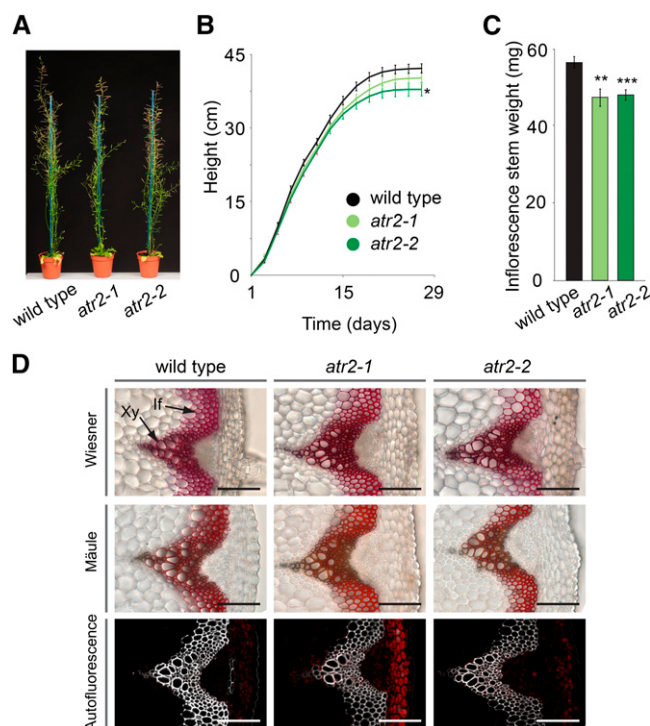


Figure 5. Phenotypic characterization of *atr2* mutants. A, Phenotype of fully grown plants after 8 weeks of short-day growth conditions followed by 5 weeks of long-day growth conditions. B, Growth curves. Height was monitored every 2 d. At the final stage, *atr2-2* was significantly smaller compared with the wild type. C, The main inflorescence stem of *atr2-1* and *atr2-2* showed reduced weight compared with that of the wild type. For B and C, error bars represent SE. * $0.05 > P > 0.01$, ** $0.01 > P > 0.001$, *** $P < 0.001$. D, Transverse stem sections of *atr2* mutants and the wild type. Mäule and Wiesner staining and lignin autofluorescence are shown. If, Interfascicular fibers; Xy, xylem. Bars = 100 μm .

or G(8-O-4/8-5)phenylpropanoic acids/esters. As all of these compounds had at least one G unit, their reduced abundance indicated that the flux toward coniferyl alcohol was reduced in the *atr2* mutants. In addition, the abundances of two aliphatic and two indolic glucosinolates (3-methyl sulfinyl glucosinolate, 2-hydroxy-3-butenylglucosinolate, indol-3-yl methylglucosinolate, and 4-hydroxyindol-3-yl-methylglucosinolate) were lower in both *atr2* mutants. Because several CYP450-dependent reactions are involved in the biosynthesis of these glucosinolates from their precursor amino acids (Supplemental Fig. S3), the reduced abundance in the *atr2* mutants could be a direct result of the lost CYP450 activity. On the other hand, the biosynthesis of all other glucosinolates also relies on two or more CYP450-dependent reactions, but only the four glucosinolates mentioned above were retrieved in the list of compounds with reduced abundance. To investigate if the other glucosinolates could have been missed due to the stringent filters ($P < 0.01$ and 2-fold difference), a targeted search for aliphatic and indolic glucosinolates was performed (Supplemental Fig. S3). However, no general tendency for a reduced abundance of aliphatic or indolic glucosinolates was apparent, indicating that the effect of *atr2* mutation on the CYP450s active in the glucosinolate pathways was moderate or nonexistent or, potentially, only secondary.

Similar to glucosinolate biosynthesis, two CYP450s (C4H and FLAVONOID-3'-HYDROXYLASE) are involved in the biosynthesis of flavonol glycosides (Supplemental Fig. S3), so a reduction in the abundance of flavonol glycosides could be anticipated in *atr2* mutants. However, a targeted search showed that kaempferol glycosides were not differentially abundant between *atr2* and the wild type (Supplemental Fig. S3). There was a tendency for lower quercetin and isorhamnetin glycoside levels (the levels were about 63%–85% in *atr2* mutants as compared with those in the wild type), but these reductions were only significant ($P < 0.05$) in *atr2-1* and not in *atr2-2* (Supplemental Fig. S3).

The 177 peaks that were higher in abundance in both *atr2* mutants as compared with wild-type plants could be assigned to 117 compounds (Supplemental Table S2), of which 63 could be structurally characterized (Table II; Supplemental Fig. S2). Many of these compounds were below the detection limit in wild-type samples and only observed in the *atr2* mutants (Table II; Supplemental Table S1). All identified compounds could be classified into one of the following metabolic classes: phenylpropanoids, benzenoids, oligolignols, hexosylated oligolignols, or H(8-O-4/8-5)phenylpropanoic acids/esters. The majority of the identified phenylpropanoids were esters or hexosides of *p*-coumaric acid, ferulic acid, or caffeic acid. *p*-Coumaric acid and *p*-coumaroyl shikimate, the substrates of the C3H1/C4H complex and C3H1, respectively, were either not or barely detectable in the wild type but accumulated in the *atr2* mutants. Similarly, cinnamic acid and coniferaldehyde, the *in vivo* substrates of C4H and F5H1, respectively, were only detected in *atr2* mutants and not in the wild type. The

benzenoids that accumulated in *atr2* mutants all contained benzoic acid or *p*-hydroxybenzoic acid moieties and were most likely derived via chain shortening of cinnamic and *p*-coumaric acids (Hertweck et al., 2001; Boatright et al., 2004).

All accumulating oligolignols, hexosylated oligolignols, and H(8-O-4/8-5)phenylpropanoic acids/esters contained at least one H unit. This observation indicates a severe increase in the flux toward *p*-coumaryl alcohol (i.e. the precursor of H units) in *atr2* mutants. In some cases, the phenylpropanoic acid/ester moiety in the H(8-O-4/8-5)phenylpropanoic acids/esters class was derived from *p*-coumaric acid. This observation is in line with the increased amount of *p*-coumaric acid-containing phenylpropanoids in *atr2* mutants and indicative of an increased flux toward *p*-coumaric acid or a reduced consumption of this metabolite.

atr2 Mutants Are Affected in Lignin Content and Composition

Fully senesced stems of wild-type plants and the *atr2* mutants were evaluated for lignin content and composition. Soluble compounds were removed from these stems by applying a sequential extraction to produce cell wall residues (CWRs; Jarvis et al., 2000; Van Acker et al., 2013). For both *atr2* mutants and the wild type, the proportion of CWR relative to the dry weight of the stem was similar at approximately 80% (Table III).

The total lignin amount present in these prepared CWRs was estimated by the spectrophotometric acetyl bromide assay. In this method, the cell wall polymers are derivatized by acetyl bromide and dissolved in acetic acid. A reduction in lignin content of 6% was detected in both *atr2-1* and *atr2-2* (Table III).

Senesced stem material of both mutant lines was analyzed further for potential changes in lignin composition. This was first done by thioacidolysis, a procedure that comprises cleavage of the β -O-4-ether bonds of the lignin polymer to release hydroxyphenyl-trithioethyl ether monomers, followed by derivatization for subsequent identification and quantification by gas chromatography coupled to MS. Thioacidolysis showed significant changes in lignin composition in both *atr2* lines, with H units increasing up to 16-fold (Table III). Thioacidolysis-released S units decreased significantly by 35% and 41% in *atr2-1* and *atr2-2* (Table III). Levels of thioacidolysis-released G units were significantly reduced by 13% for the *atr2-2* line compared with the wild type based on CWR (Table III). A similar decrease in thioacidolysis-released G units was also measured for *atr2-1* but was not significant below the 0.05 P value threshold ($P = 0.06$). Consequently, reductions in the S/G ratio by 32% and 25% were observed for *atr2-1* and *atr2-2*, respectively. The degree of lignin condensation inversely correlates with the sum of H, G, and S units released from the lignin by thioacidolysis. This sum was not significantly different from that of the wild type (Table III).

Table 1. Metabolites with a reduced abundance in *atr2* mutants

An overview of the identified compounds of which the abundance was lower in the *atr2-1* and *atr2-2* mutants is given. For each compound, average peak areas of wild-type and *atr2* mutant samples are given. For the structural elucidation of the compounds, see Supplemental Table S2 and Supplemental Figure S2.

No.	Retention Time	m/z	Name	Wild Type		atr2-1		Change in atr2-1:Wild-Type Ratio	atr2-2		Change in atr2-2:Wild-Type Ratio
				Mean	SE	Mean	SE		Mean	SE	
min											
Glucosinolates											
1	1.28	422.025	3-Methylsulfinylpropyl glucosinolate	176,502	7,833	48,551	4,389	0.28	46,856	2,768	0.27
2	3.82	388.077	2-Hydroxy-3-butenylglucosinolate	2,701	293	526	129	0.19	386	51	0.14
3	3.61	895.117	Indol-3-ylmethylglucosinolate (dimer)	10,432	1,004	3,219	456	0.31	2,997	273	0.29
4	2.10	847.149	4-Hydroxyindol-3-ylmethylglucosinolate (dimer) ^a	2,194	129	926	173	0.42	1,064	159	0.49
Oligolignols											
5	8.84	375.145	G(8-O-4)G	274	58	0	0	0.00	0	0	0.00
6	9.14	375.146	G(8-O-4)G	222	57	0	0	0.00	0	0	0.00
7	12.97	339.125	G(8-5)G ^a	2,063	288	33	17	0.02	33	12	0.02
8	15.45	337.109	G(8-5)G ^a	877	79	13	9	0.02	81	27	0.09
9	12.80	369.136	S(8-5)G ^a	208	41	0	0	0.00	0	0	0.00
10	13.12	553.208	G(8-O-4)G(8-5)G	356	71	0	0	0.00	0	0	0.00
11	13.42	553.209	G(8-O-4)G(8-5)G	574	65	0	0	0.00	0	0	0.00
12	14.71	583.218	G(8-O-4)S(8-5)G	7,656	817	78	44	0.01	51	25	0.01
13	15.52	583.219	G(8-O-4)S(8-5)G	2,158	301	0	0	0.00	0	0	0.00
14	16.25	581.203	G(8-O-4)S(8-5)G'	829	100	19	19	0.02	5	5	0.01
15	12.77	555.224	G(8-O-4)G(red8-5)G	456	80	0	0	0.00	5	5	0.01
16	16.66	613.232	G(8-O-4)S(8-8)S	37	15	0	0	0.00	0	0	0.00
17	16.96	583.219	G(8-O-4)S/G(8-8)G/S	1,260	233	0	0	0.00	0	0	0.00
18	18.00	809.306	G(8-O-4)G(8-O-4)S(8-8)S	272	63	0	0	0.00	0	0	0.00
Hexosylated oligolignols											
19	6.06	583.203	G 4-O-hexoside(8-O-4)G ^b	523	80	24	18	0.05	46	30	0.09
20	7.61	613.212	G 4-O-hexoside(8-O-4)S ^b	778	71	46	19	0.06	48	21	0.06
21	8.20	521.203	G(red8-8)G hexoside	412	60	42	17	0.10	53	28	0.13
22	9.04	521.202	G(red8-8)G hexoside	882	54	365	73	0.41	407	77	0.46
23	9.00	565.193	G 4-O-hexoside(8-5)G ^b	3,748	298	576	91	0.15	1,053	136	0.28
24	9.14	567.208	G 4-O-hexoside(red8-5)G ^b	2,975	321	182	81	0.06	286	91	0.10
25	10.68	521.203	G(red8-5)G hexoside	525	65	159	36	0.30	149	23	0.28
G(8-O-4/8-5)phenylpropanoic acid/ester											
26	5.98	551.177	G(8-O-4)feruloyl hexose	2,527	118	744	147	0.29	736	105	0.29
27	11.68	775.246	G(8-O-4)feruloyl hexose + 224 D	7,413	878	2,154	319	0.29	3,243	383	0.44
28	11.89	775.246	G(8-O-4)feruloyl hexose + 224 D	4,438	405	1,318	171	0.30	2,025	235	0.46
29	9.54	533.167	G(8-5)feruloyl hexose	492	39	0	0	0.00	16	10	0.03
30	10.08	533.166	G(8-5)feruloyl hexose	562	44	229	24	0.41	116	20	0.21

^aCompounds detected as fragments of the parent ion. ^bCompounds detected as formic acid adducts.

Changes in both lignin aromatic and side chain regions were further revealed by applying two-dimensional ¹H-¹³C correlation heteronuclear single quantum coherence (HSQC) NMR on whole-plant cell wall samples of the wild type and the *atr2* mutants prepared from senesced main stems (Fig. 6; Supplemental Fig. S4). In contrast to thioacidolysis, which determines the composition of units linked only by ether bonds, NMR analysis of the aromatic region allows the investigation of all monomers irrespective of their mode of linkage. In addition, interunit linkage types can be determined by use of the aliphatic region of the spectrum. Compared with the wild type, the fraction of H units in the lignin of *atr2*

mutants was increased about 10-fold (Fig. 6). The relative fraction of G units was not significantly different, but the relative fraction of S units was reduced in *atr2* mutants by almost 60% compared with the wild type. Accordingly, the S/G ratio decreased significantly from 0.31 in the wild type to 0.13 and 0.11 in the *atr2-1* and *atr2-2* mutants (Fig. 6). The HSQC spectrum fully resolved correlations for the β-aryl ether (β-O-4), phenylcoumaran (β-5), and resinol (β-β) units. The β-syringyl ether linkages, (β-O-4)S, were reduced, reflecting the lower S unit content, whereas those linkages in which the aryl moiety is an H or G unit, (β-O-4)H/G, were seemingly not affected (Fig. 6).

Table II. Metabolites with an increased abundance in *atr2* mutants

An overview of the identified compounds of which the abundance was higher in the *atr2-1* and *atr2-2* mutants is given. For each compound, average peak areas of wild-type and *atr2* mutant samples are given. Infinity (inf) is given when a compound is below detection limit in the wild type. For the structural elucidation of the compounds, see Supplemental Table S2 and Supplemental Figure S2.

No.	Retention Time	<i>m/z</i>	Name	Wild Type		<i>atr2-1</i>		Change in <i>atr2-1</i> :Wild- Type Ratio	<i>atr2-2</i>		Change in <i>atr2-2</i> :Wild Type Ratio
				Mean	SE	Mean	SE		Mean	SE	
<i>min</i>											
Phenylpropanoids											
31	13.73	147.045	Cinnamic acid	0	0	300	99	inf	235	41	inf
32	6.75	163.040	<i>p</i> -Coumaric acid	2	1	1,147	136	593	2,241	438	1,159
33	4.47	325.093	<i>p</i> -Coumaroyl hexose	330	21	10,773	1,695	33	22,378	4,991	68
34	5.05	325.093	<i>p</i> -Coumaroyl hexose	22	12	2,534	408	114	4,305	1,051	194
35	11.66	549.162	<i>p</i> -Coumaroyl hexose + 224 D	0	0	674	134	inf	1,279	245	inf
36	13.36	549.161	<i>p</i> -Coumaroyl hexose + 224 D	0	0	728	196	inf	1,441	331	inf
37	7.80	279.052	<i>p</i> -Coumaroyl malate	0	0	390	73	inf	358	70	inf
38	8.26	279.052	<i>p</i> -Coumaroyl malate	0	0	314	93	inf	260	80	inf
39	7.89	319.083	<i>p</i> -Coumaroyl shikimate	0	0	4,164	347	inf	4,808	540	inf
40	9.49	319.084	<i>p</i> -Coumaroyl shikimate	0	0	198	53	inf	254	44	inf
41	3.00	325.094	<i>p</i> -Coumaric acid 4- <i>O</i> -hexoside	0	0	791	77	inf	1,458	132	inf
42	3.71	327.110	Dihydro- <i>p</i> -coumaric acid + hexose	0	0	564	121	inf	1,173	185	inf
43	3.90	327.108	Dihydro- <i>p</i> -coumaric acid + hexose	5,099	215	12,814	973	3	16,108	572	3
44	3.31	341.089	Caffeoyl hexose	0	0	667	109	inf	1,075	187	inf
45	2.85	341.089	Caffeic acid 3/4- <i>O</i> -hexoside	66	26	2,391	255	36	3,528	264	54
46	3.97	341.089	Caffeic acid 3/4- <i>O</i> -hexoside	266	23	2,085	176	8	2,710	149	10
47	5.44	341.088	Caffeic acid 3/4- <i>O</i> -hexoside	0	0	637	102	inf	712	111	inf
48	2.97	503.141	Caffeoyl hexose 3/4- <i>O</i> -hexoside	0	0	1,053	130	inf	1,683	247	inf
49	5.28	355.104	Feruloyl hexose	377	25	2,362	261	6	3,852	597	10
50	5.80	355.105	Feruloyl hexose	0	0	187	43	inf	265	65	inf
51	12.31	217.051	Feruloyl hexose + 224 D ^a	7	5	203	43	29	397	67	58
52	13.55	355.104	Feruloyl hexose + 224 D ^a	0	0	119	41	inf	277	69	inf
53	8.77	309.062	Feruloyl malate	0	0	1,273	239	inf	577	116	inf
54	9.13	309.062	Feruloyl malate	0	0	1,013	161	inf	810	101	inf
55	3.69	355.105	Ferulic acid 4- <i>O</i> -hexoside	20	10	398	86	20	519	43	26
56	5.48	385.114	Sinapoyl hexose	8,150	713	23,447	2,222	3	27,655	2,809	3
57	6.06	205.051	Sinapoyl hexose ^a	23	10	185	17	8	275	43	12
58	13.22	591.171	Disinapoyl Glc	666	113	4,152	843	6	4,499	765	7
59	10.21	177.056	Coniferaldehyde	0	0	181	45	inf	392	35	inf
Benzenoids											
60	9.16	237.041	Benzoyl malate	500	38	1,581	181	3	2,250	179	5
61	5.44	121.030	<i>p</i> -Hydroxybenzaldehyde	81	13	374	39	5	694	64	9
62	2.64	299.078	<i>p</i> -Hydroxybenzoyl hexose	0	0	320	100	inf	632	199	inf
63	1.85	299.078	<i>p</i> -Hydroxybenzoic acid 4- <i>O</i> -hexoside	1,738	101	5,093	398	3	6,300	292	4
Oligolignols											
64	12.32	309.114	H(8-5)C ^a	0	0	346	75	inf	439	52	inf
65	13.16	279.104	H(8-5)H ^a	0	0	108	47	inf	170	42	inf
66	13.09	523.197	H(8-O-4)G(8-5)G	0	0	518	107	5,318	735	54	7,540
67	14.95	521.182	H(8-O-4)G(8-5)G	0	0	301	95	inf	674	56	inf
68	15.12	521.182	H(8-O-4)G(8-5)G	0	0	708	131	inf	1,386	117	inf
69	15.94	523.197	H(8-O-4)G(8-5)G	0	0	1,223	261	inf	1,913	98	inf
70	16.76	523.198	H(8-O-4)G(8-8)G	0	0	497	143	inf	870	77	inf
71	13.85	493.187	H(8-O-4)H/G(8-5)G/H	0	0	227	69	inf	478	41	inf
72	14.41	553.208	H(8-O-4)S(8-5)G	16	12	2,985	425	190	3,597	117	229
73	15.16	553.208	H(8-O-4)S(8-5)G	10	7	2,341	335	225	2,759	74	265
74	16.46	551.192	H(8-O-4)S(8-5)G'	0	0	2,457	464	inf	4,413	369	inf
75	17.18	551.192	H(8-O-4)S(8-5)G'	0	0	948	225	inf	1,726	161	inf
Hexosylated oligolignols											
76	6.00	621.218	H(8-8)H dihexoside	119	22	11,070	1,461	93	17,861	2,405	150
77	9.00	459.166	H(8-8)H <i>O</i> -4-hexoside	0	0	1,952	261	inf	3,808	406	inf
78	9.60	489.177	H(8-8)G hexoside	0	0	187	50	inf	220	41	inf

(Table continues on following page)

(Table continues on following page.)

Table II. (Continued from previous page.)

No.	Retention Time	m/z	Name	Wild Type		<i>atr2-1</i>		Change in <i>atr2-1</i> :Wild- Type Ratio	<i>atr2-2</i>		Change in <i>atr2-2</i> :Wild Type Ratio
				Mean	SE	Mean	SE		Mean	SE	
79	6.59	651.230	H(8-8)G dihexoside	0	0	569	90	inf	775	111	inf
80	9.16	505.171	H O-4-hexoside(8-5)H ^b	0	0	4,748	538	2,149	8,575	950	222,225
81	7.50	621.219	H 8-O-hexoside (8-5)H 8-O-hexoside	0	0	211	92	inf	591	194	inf
82	8.53	535.181	H/G(8-5)G/H 4-O-hexoside ^b	0	0	1,903	262	inf	3,375	319	inf
83	10.88	461.180	H(red8-5)H hexoside	0	0	129	44	inf	443	65	inf
H(8-O-4/8-5)phenylpropanoic acids/esters											
84	11.20	715.225	H(8-O-4) <i>p</i> -coumaroyl hexose + 224 D	0	0	575	175	inf	1,357	207	inf
85	11.59	715.228	H(8-O-4) <i>p</i> -coumaroyl hexose + 224 D	0	0	187	65	2,149	543	106	6,255
86	10.80	473.145	H 4/9-O hexoside(8-5) <i>p</i> -coumaric acid	0	0	172	44	inf	703	187	inf
87	13.62	427.103	H(8-5) <i>p</i> -coumaroyl malate	0	0	462	159	inf	786	112	inf
88	13.91	427.104	H(8-5) <i>p</i> -coumaroyl malate	0	0	546	197	inf	831	131	inf
89	6.07	521.166	H(8-O-4)ferulic acid hexoside	0	0	228	46	inf	829	195	inf
90	11.52	745.235	H(8-O-4)feruloyl hexose + 224 D	4	4	1,569	397	445	3,297	511	936
91	9.11	475.125	H(8-O-4)feruloyl malate	9	5	3,252	631	348	4,165	474	445
92	9.95	475.126	H(8-O-4)feruloyl malate	3	2	932	277	364	1,107	168	433
93	13.16	457.114	H(8-5)feruloyl malate	0	0	904	184	inf	1,186	160	inf
94	10.84	389.127	H(8-O-4)sinapoyl malate ^a	0	0	292	102	inf	455	67	inf

^aCompounds detected as fragments of the parent ion. ^bCompounds detected as formic acid adducts.

Transcriptional Feedback on *CYP450s* Involved in Lignin Biosynthesis

To investigate the possible feedback caused by *ATR2* deficiency (and the concomitant shifts in phenolic metabolism) on the transcript levels of the lignin-associated *CYP450s*, the expression of *C4H*, *C3H1*, and *F5H1* in inflorescence stems was investigated via qRT-PCR (Fig. 4B). In addition, expression levels of *PAL1* and *PAL2* were used as controls, as these are not *CYP450*-encoding genes but also involved in lignin biosynthesis. No significant differences were observed in the levels of the measured transcripts between the *atr2-1* mutant and the wild type. In the *atr2-2* mutant, *C3H1* expression was significantly increased as compared with the wild type ($P = 0.049$), and the expression of *PAL2* and *C4H* was on the border of being increased significantly, with P values of 0.059 and 0.064, respectively.

atr2 Mutant Biomass Is Easier to Saccharify

Lignin content and composition generally have an impact on cell wall digestibility. Because *atr2* mutants have reduced lignin levels and shifts in lignin composition compared with the wild type, we investigated the saccharification of senesced *atr2* mutant stems. First, the cellulose content of the prepared CWRs was determined by the Updegraff method. In this method, the crystalline cellulose polymer that remains after acetic and nitric acid treatment is broken down to Glc by sulfuric acid; the amount of released Glc is determined spectrophotometrically after derivatization with anthrone. No significant changes in crystalline cellulose levels were detected in the *atr2* mutant lines when compared with the wild type (Table III). Saccharification measures the amount of Glc released from the enzymatic breakdown of cellulose present in the plant cell

Table III. Cell wall characterization of *atr2*

The CWR is the material remaining after sequential extraction of dry senesced stem biomass with hydrophobic and hydrophilic solvents. Lignin content was measured with the acetyl bromide assay. Cellulose content was measured with the Updegraff method. Monomer composition was determined by thioacidolysis. P values are from Student's t test: $*0.05 > P > 0.01$, $**0.01 > P > 0.001$, and $***P < 0.001$; $n = 8$. No diff, No difference.

Parameter	Wild Type		<i>atr2-1</i>		P	Change in <i>atr2-1</i> versus the Wild Type	<i>atr2-2</i>		P	Change in <i>atr2-2</i> versus the Wild Type
	Mean	SE	Mean	SE			Mean	SE		
CWR (%)	77.8	0.6	76.2	0.6	0.08	% No diff	77.0	1.1	0.3	% No diff
Lignin per CWR (%)	20.8	0.4	19.6	0.2	0.03*	−6	19.6	0.2	0.03*	−6
Cellulose per CWR (%)	30.9	1.2	31.2	0.4	0.84	No diff	32.5	1.7	0.5	No diff
H units per CWR ($\mu\text{mol mg}^{-1}$)	0.38	0.03	5.9	0.32	4.6E-07***	+1,454	6.1	0.25	5.0E-08***	+1,516
G units per CWR ($\mu\text{mol mg}^{-1}$)	48.4	1.28	41.9	2.97	0.06	No diff	42.3	1.51	7.8E-03**	−13
S units per CWR ($\mu\text{mol mg}^{-1}$)	23.0	1.23	14.8	1.01	1.5E-04***	−35	13.6	0.50	5.9E-06***	−41
S/G ratio	0.47	0.02	0.32	0.01	2.7E-04***	−32	0.36	0.01	8.7E-05***	−25
(H+G+S) per Lignin ($\mu\text{mol mg}^{-1}$)	344.7	10.9	319.4	20.8	0.3	No diff	317.0	9.5	0.08	No diff

wall. The applied procedure included either no pretreatment or an alkaline (6.25 mM NaOH) pretreatment to loosen the cell wall components, thereby facilitating access of the enzymes to the cell wall polysaccharides. The released Glc was measured after 3, 7, 24, and 48 h. Without pretreatment, a significantly reduced Glc release was observed after 3 h of saccharification for both *atr2* mutants and after 7 h for *atr2-1* only. However, after 48 h of saccharification, *atr2-2* released more Glc than the wild type. The cellulose-to-Glc conversion improved strikingly after alkaline pretreatment; after 24- and 48-h incubation times, 26% and 27% improvements were reached in *atr2-1* and *atr2-2*, respectively, as compared with the wild type (Fig. 7).

DISCUSSION

ATR2 Is Involved in Lignification

ATR2-deficient plants have 6% less lignin with a different composition compared with wild-type plants (Table III; Fig. 6). The composition of lignin has traditionally been an important clue in determining at which position in the pathway the gene in question is acting. A relative increase in H units has been reported in plants in which the flux from *p*-coumaroyl-CoA to caffeoyl-CoA and caffeic acid was disturbed (i.e. in plants with reduced HCT, C3H, and/or CSE activity; Abdulrazzak et al., 2006; Ralph et al., 2006, 2012; Coleman et al., 2008; Vanholme et al., 2013c; Bonawitz et al., 2014). The 10- to 15-fold increase in H units in *atr2* mutants (Table III; Fig. 6), therefore, is likely a direct result of reduced C3H1 activity because of the compromised ATR electron transfer. In addition, the lignin in *atr2* mutants has a reduced S/G ratio (Fig. 6). Commonly, plants with a lower flux toward coniferaldehyde (e.g. *c4h*, *4cl1*, *c3h1*, and *coaomt1*) have lignin with higher S/G ratios, because the remaining flux through the pathway is preferentially used by the rate-limiting enzyme F5H1 and not by cinnamyl alcohol dehydrogenase (Meyer et al., 1996; Van Acker et al., 2013; Bonawitz et al., 2014; Wang et al., 2014). Thus, the reduced S/G ratio in *atr2* mutants strongly suggests that the F5H1 activity is also negatively affected by reduced ATR2 activity (Fig. 6).

Mutation of ATR2 Causes Widespread Changes in Aromatic Metabolism

Phenolic profiling of the *atr2* mutants revealed large differences in the metabolite pool compared with the wild type. Previously, it has been shown that lignin mutants display changes in their phenolic metabolome and that they commonly accumulate the substrate of the deficient enzyme or derivatives thereof (Vanholme et al., 2012b, 2013c). In particular, *c4h* Arabidopsis mutants accumulate cinnamic acid (Schilmiller et al., 2009), *c3h1* Arabidopsis mutants and C3H down-regulated poplar accumulate *p*-coumaroyl shikimate

(Coleman et al., 2008; Bonawitz et al., 2014), and mutants in *F5H1* accumulate coniferaldehyde (Vanholme et al., 2012b). In the *atr2* mutants, cinnamic acid, *p*-coumaroyl shikimate, and coniferaldehyde were all significantly higher in abundance compared with the wild type (Fig. 1; Table II). The accumulation of these compounds can be linked directly to the role of ATR2 in serving electrons to C4H, C3H1, and F5H1.

The accumulating phenylpropanoids and benzenoids in *atr2* mutants were derivatives of *p*-coumaric, caffeic, ferulic, and sinapic acids (Table II). Most of them were detected in their hexosylated or malated form. Hexosylation occurred as 9-*O*-esters, as 3/4-*O*-hexosides, or both. Hexosylation and malation of accumulating phenylpropanoids have been suggested as either a strategy for the plant to detoxify potentially harmful metabolites or a passive process wherein accumulating phenylpropanoids serve as surrogates for the wild-type substrates (Vanholme et al., 2010b). Accumulating metabolites have been commonly detected when individual steps of the lignin biosynthetic pathway were perturbed. It has been shown that *c4h* mutants accumulate metabolites derived from cinnamic acid and ferulic acid (Vanholme et al., 2012b), *f5h1* mutants accumulate metabolites derived from ferulic acid (Vanholme et al., 2012b), and plants with strongly reduced C3H1 activity have increased levels of *p*-coumaric acid-derived metabolites (Bonawitz et al., 2014). The accumulation of *p*-coumarates, caffeates, and ferulates and their benzenoid derivatives in *atr2* mutants are a logical consequence of the reduced activity of C4H, C3H1, and F5H1.

Sinapate esters such as sinapoyl malate and sinapoyl hexose are known to protect wild-type Arabidopsis against harmful UV-B radiation and, therefore, are mainly localized in the epidermis (Chapple et al., 1992; Landry et al., 1995). Paradoxically, sinapate esters accumulated in the *atr2* mutants (Table II), whereas *f5h1*, *c3h1*, and *c4h* mutants have been reported to have fewer sinapate esters (Franke et al., 2002; Vanholme et al., 2012b). This observation might logically be explained by the fact that ATR2 electron donation is important for lignifying cells, but epidermis cells that accumulate sinapate esters do not lignify. Therefore, it is possible that the mutation in ATR2 has no negative effect on the F5H1 activity in the epidermis; electrons might be supplied to C4H, C3H1, and F5H1 by the constitutive ATR1 in this cell type for the biosynthesis of sinapate esters.

In addition to phenylpropanoid and lignin biosynthesis, several other pathways rely on CYP450 enzymes. For instance, *cyp79f1* mutants and Arabidopsis plants down-regulated for *CYP79F1* have reduced levels of short-chain aliphatic glucosinolates (Hansen et al., 2001; Chen et al., 2003), *cyp79f2* mutants were affected in the biosynthesis of long-chain aliphatic glucosinolates (Chen et al., 2003), and *cyp83a1* mutants have reduced levels of both short- and long-chain aliphatic glucosinolates (Hemm et al., 2003). The *cyp79b1*, *cyp79b2*, *cyp83b1*, and *cyp81f2* mutants and the *cyp79b1 cyp79b2* double mutant

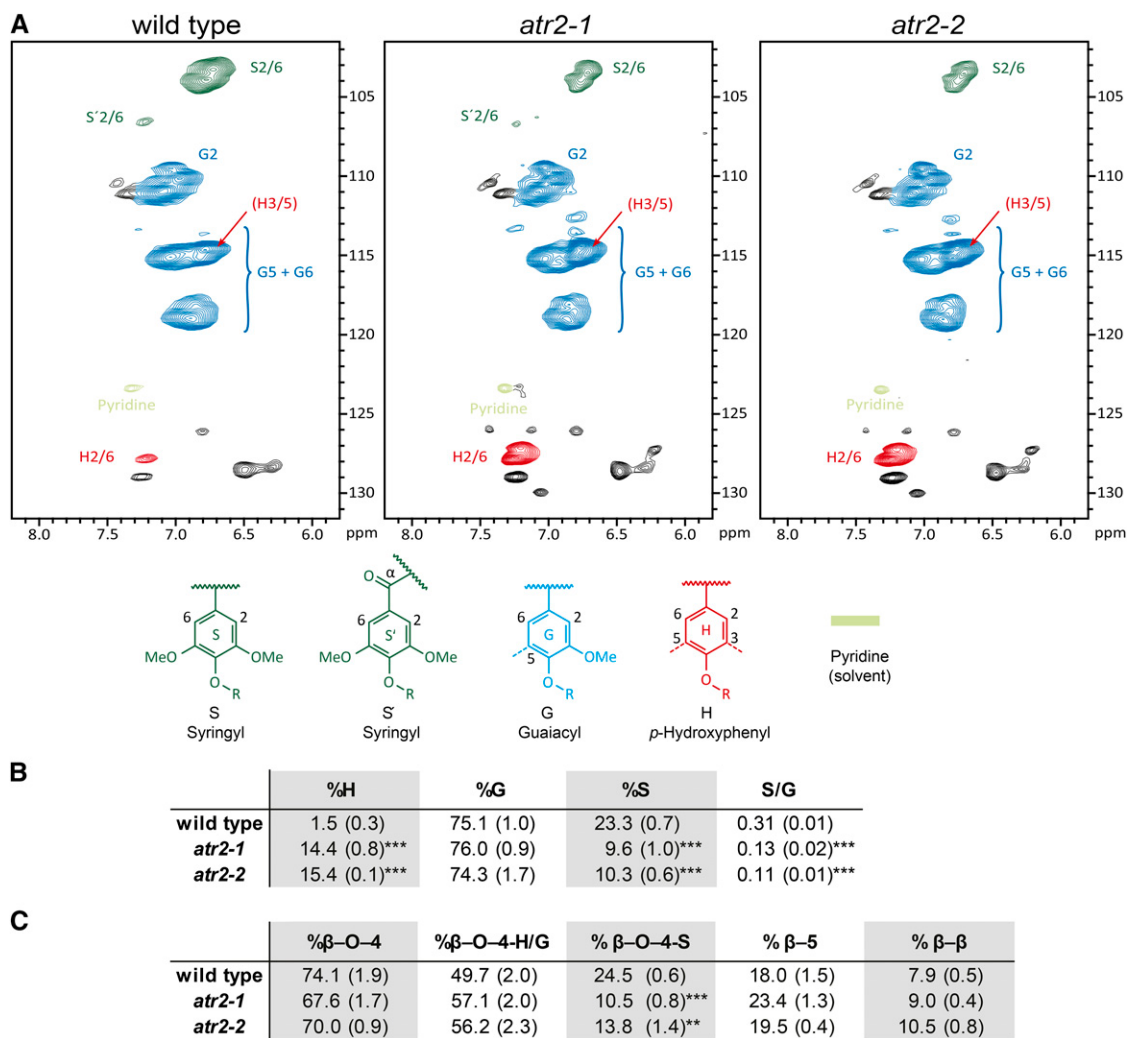


Figure 6. NMR lignin characterization. A, Two-dimensional HSQC NMR spectra of the lignin aromatic region of senesced stem material from the wild type, *atr2-1*, and *atr2-2*. The volume integrals of the S2/6, G2, and H2/6 contour peaks, when scaled (by 2 for G), represents the quantity of each monomer. B, Overview of the relative amounts of the different monoglignol-derived units and the S/G ratio as determined from the HSQC aromatic region. C, Overview of the relative amount (as determined by uncorrected volume integrals) of interunit linkage types (β -O-4, β -5, and β - β) as determined from the HSQC side chain region (for the spectra of the side chain regions, see Supplemental Fig. S4). ** $0.01 > P > 0.001$, *** $P < 0.001$; $n = 3$.

have reduced levels of indolic glucosinolates (Zhao et al., 2002; Naur et al., 2003; Bednarek et al., 2009). In addition, *c4h* and *f3'h* mutants have less flavonol glycosides (Schoenbohm et al., 2000; Vanholme et al., 2012b). On the other hand, *c3h1* mutants accumulate flavonoid glycosides, but this has been attributed to a stress-induced synthesis and not metabolic overflow (Abdulrazzak et al., 2006; Bonawitz et al., 2014). However, a targeted phenolic profiling revealed that a mutation of *ATR2* had only moderate effects on the abundance of glucosinolates and flavonol glycosides (Supplemental Fig. S3). Again, this indicates that *ATR2* has a preferential role in lignin biosynthesis in *Arabidopsis*. However, this does not rule out the possibility that *ATR2* activity is needed for glucosinolate and flavonol synthesis in other cell types or during stress.

The differences in lignin composition and phenolic metabolite levels between the *atr2* mutants and wild-type plants can logically be explained by lower C4H, C3H1, and F5H1 activities in the *atr2* mutants. The ability of *ATR2* to donate electrons to C4H was already proven in vitro in yeast and baculovirus expression systems (Urban et al., 1997; Mizutani and Ohta, 1998). Here, we provided in planta evidence for this interaction. To exclude that these reduced activities were due to a lower expression of the corresponding genes, we determined the expression levels of *C4H*, *C3H1*, and *F5H1* in the inflorescence stem of *atr2* mutants. The expression levels of these genes were not reduced in the *atr2* mutants as compared with the wild type. In contrast, the expression of *C3H1* was significantly higher in *atr2-2* and the expression of *C4H* was on the border of being

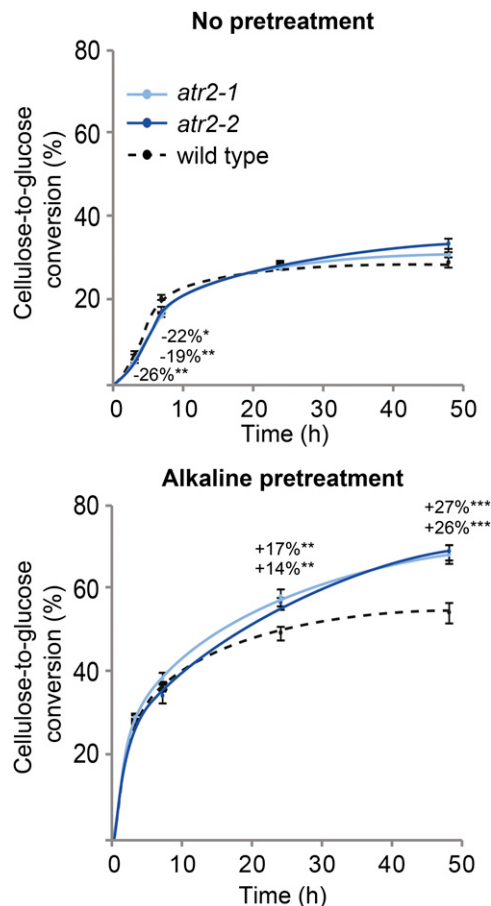


Figure 7. Saccharification data of the *atr2* mutants. Samples were saccharified with either no pretreatment or an alkaline pretreatment (6.25 mM NaOH). The cellulose-to-Glc conversion was measured at 3, 7, 24, and 48 h ($n = 8$). Error bars represent SE. * $0.05 > P > 0.01$, ** $0.01 > P > 0.001$, *** $P < 0.001$; $n = 8$.

significantly higher (Fig. 4B). These data indicate that the reduced C4H, C3H1, and F5H1 activities are not the result of reduced gene expression. More likely, the reduced C4H, C3H1, and F5H1 activities are caused by the compromised electron-donating capacities of CPRs in the *atr2* mutants. Notably, our results do not exclude the possibility that C4H, C3H1, and F5H1 activities were reduced via a posttranscriptional mechanism (e.g. on the protein activity level, where particular accumulating phenolics would act as inhibitors of the CYP450s).

Mutation of *ATR2* Has Modest Effects on Plant Growth

Fully grown *atr2-2* plants were only slightly smaller at final height than their wild-type counterparts, whereas *atr2-1* was similar in size (Fig. 5). However, the reduction in dry weight of the main inflorescence stem in both *atr2* mutants showed that normal plant development was affected (Fig. 5). A reduction in biomass has been observed previously in Arabidopsis plants with reduced lignin levels (e.g. in *c4h*, *c3h1*, *cse*, and *ccr1* mutants; Jones et al., 2001; Franke et al., 2002;

Do et al., 2007; Schillmiller et al., 2009; Van Acker et al., 2013; Vanholme et al., 2013c). Several hypotheses have been put forward to explain the growth penalty of low-lignin plants, of which some were proven (Bonawitz and Chapple, 2013). For example, a reduced amount of lignin results in weaker vessel cell walls and, in extreme cases, in vessel collapse, which in turn affects the transport of water and solutes (Turner and Somerville, 1997; Yang et al., 2013). Strengthening the cell walls of *c4h* mutants by vessel-specific expression of C4H could restore the growth phenotype, even though overall lignin levels were still reduced compared with the wild type (Yang et al., 2013). It was also recently discovered that the transcriptional coregulatory complex, Mediator, is a key component for mediating the plant's response commonly associated with lignin biosynthesis reduction (stunted growth and less lignin). Mutation of genes encoding components of the Mediator complex resulted in a partial reversion of the growth penalty of *c3h1* mutants (Bonawitz et al., 2014). We have not observed any deformation of cell walls in *atr2* mutants (Fig. 5), but as *ATR2* affects the functioning of C3H1, it is possible that the *atr2* mutation triggers the Mediator-regulated transcriptional cascade, resulting in reduced biomass.

Because Arabidopsis has 244 genes encoding CYP450s, it is striking that *atr2* mutants did not display stronger morphological defects. *ATR1* and *ATR2* are the only known genes in Arabidopsis for which the corresponding proteins have genuine CPR activity, meaning that they have the ability to transfer electrons from NADPH to CYP450s. It has been reported that *ATR1* is constitutively expressed whereas *ATR2* is mainly induced by wounding and light (Mizutani and Ohta, 1998). We further note that *ATR2*, but not *ATR1*, expression is inducible in seedlings treated with isoxaben, a cellulose inhibitor that induces several processes, including ectopic lignification (Fig. 3). The differences in *ATR1* and *ATR2* expression were also apparent after reevaluation of published data (Fig. 2). That *ATR1* appears to be constitutively expressed suggests it to be partially redundant with *ATR2* function, and this may explain the absence of strong morphological defects in *atr2* mutants. However, we cannot rule out the possibility that the truncated *ATR2* transcripts detected in both *atr2* mutant alleles (Fig. 4) still provide leaky *ATR2* activity and that a full knockout would have revealed stronger phenotypes, as was the case, for example, for the allelic *c4h* mutant series (Schillmiller et al., 2009). Taken together, our results support the hypothesis that *ATR1* provides CPR activity constitutively, fulfilling the need for electrons in most cell types during development, and that *ATR2* is induced in cases where additional CPR activity is needed that cannot be fulfilled by *ATR1* alone (e.g. during lignification).

ATR2 Homologs as Targets to Improve Saccharification in Crops

The presence of lignin in the cell wall physically hinders the enzymatic hydrolysis of cellulose into Glc (Alvira et al., 2010). The saccharification efficiency can

be improved by applying a pretreatment to render the cellulose more accessible to cellulases. Different pretreatments have specific effects on the cell wall material (Mosier et al., 2005; Carvalho et al., 2008; Alvira et al., 2010). We have tested alkaline (NaOH) pretreatment prior to the saccharification of cell wall material of wild-type plants and *atr2* mutants and compared it with the saccharification efficiencies of cell walls without pretreatment. Although no improvement in saccharification efficiency was observed without pretreatment, a 26% to 27% increase in cellulose-to-Glc conversion was observed with alkaline-pretreated *atr2* mutants stems (Fig. 7). The improved saccharification of *atr2* mutant biomass may be explained by the reduced lignin content but also by the alterations of the lignin structure. Enrichment in *p*-coumaryl alcohol results in shorter lignin polymers because the H units tend to stop the polymerization process (Russell et al., 2006; Ziebell et al., 2010; Vanholme et al., 2012a, 2013a). This is supported by the identity of the oligolignols that accumulated in the *atr2* mutant: they all had a terminal H unit, in most cases β -O-4 linked to a G unit (Table II). Shorter lignins will be more easily extracted during the alkaline pretreatment.

Besides ATR2, other members of the electron transport chain also might become targets for cell wall engineering. It was apparent from Vanholme et al. (2012b) that there is at least one CB5 coexpressed with lignin biosynthetic genes in a similar way to ATR2. Another CB5 has been shown by tandem affinity purification to interact with C4H in Arabidopsis cell cultures grown under standard conditions and under conditions inducing differentiation to tracheary elements (Bassard et al., 2012). Functional analysis of these CB5s will shed further light on their putative role as electron donors of the CYP450s involved in lignification.

In conclusion, ATR or other components of the CYP450 electron chain such as CB5s may become interesting candidates for the genetic engineering of lignocellulosic crops if the yield penalty can be overcome (e.g. by vessel-specific complementation or by engineering the Mediator complex; Yang et al., 2013; Bonawitz et al., 2014).

MATERIALS AND METHODS

Plant Material

Arabidopsis (*Arabidopsis thaliana*) T-DNA insertion lines of ATR2 (At4g30210), SALK_152766 and SALK_026053, were ordered from the Nottingham Arabidopsis Stock Centre and called *atr2-1* and *atr2-2*, respectively. Homozygous lines were screened by PCR using two primer combinations to distinguish heterozygous lines from homozygous ones: for the presence of the *atr2-2* allele, 5'-CTCGTTTGGAGGAGATCCG-3' (primer 1) and 5'-GCCCTTTGACGTTGGAGTC-3' (primer 2), and for the absence of the *atr2-2* allele, primer 1 and 5'-CATCTC-CATATCTACATCCAG-3' (primer 3); for the presence of the *atr2-1* allele, primer 2 and 5'-GTGGAACACACTCCCTTATG-3' (primer 4), and for the absence of the *atr2-1* allele, primer 4 and 5'-CACACCAATCAGCAGCTCAC-3' (primer 6).

Isoxaben Treatment

Seeds were germinated in 25 mL of liquid Murashige and Skoog medium (0.5× Murashige and Skoog medium + 1% [w/v] Glc, pH 5.7), and seedlings

were grown in a 16-/8-h light/dark cycle at 21°C with gentle agitation (130 rpm). After 3 d, isoxaben dissolved in dimethyl sulfoxide (DMSO) was added (80 nM final concentration), and the same amount of DMSO was also added to the control. Twenty seedlings per repeat were harvested at 0, 6, 24, and 72 h after isoxaben addition.

qRT-PCR Analysis of Stems and Seedlings

To determine the transcript levels in the *atr2* mutants, the basal 2 cm of the inflorescence stems was sampled from plants with a height of about 24 cm. Total RNA of seedlings and stem material was prepared using the RNeasy Plant Mini Kit (Qiagen). The RNA from the seedlings was treated with RQ1 RNase-Free DNase (Promega), and the RNA from stems was treated with DNA-free (Ambion) to remove traces of genomic DNA. Subsequent complementary DNA preparation was done using the iScript cDNA Synthesis Kit (Bio-Rad). Relative expression levels were determined with the LightCycler 480 Real-Time SYBR Green PCR System (Escoubas et al., 1995). For qRT-PCR of seedlings, *CYCLIN-DEPENDENT KINASE A1* (At3g48750) was used as the reference gene, whereas for qRT-PCR of stem material, the gene encoding S-adenosyl-L-Met-dependent methyltransferases superfamily protein (At2g32170) was used. To measure ATR2 expression in the experiment performed on seedlings, primers 13 and 14 were combined, while for the experiment performed with stem material, primer combinations 15/16 (primer set 2) and 17/18 (primer set 1) were used. Primers are listed in Supplemental Table S3.

Plant Growth and Harvest

Plants were randomized and germinated in short-day conditions (21°C, 9 h of light, 120 $\mu\text{E s}^{-1} \text{m}^{-2}$; 18°C, 15 h of darkness) on soil in pots 5.3 cm wide × 5.5 cm high. Plants were moved to long-day conditions (16-/8-h light/dark cycle) after 8 weeks to allow bolting. For metabolomics, plants were grown until the stem reached 22 to 26 cm in height. The bottom 10 cm of the main stem was then harvested, frozen in liquid nitrogen, and deprived of all side branches. For saccharification and cell wall compositional analyses, plants were grown until full senescence. The bottom 15 cm of the main stem was harvested and analyzed. For microscopy, stems of fully grown but not yet senesced plants were used.

Microscopy

The bottom 2 cm of inflorescence stems was embedded in 7% (w/v) agarose. Using a vibratome from Campden Instruments, slices 100 μm thick were made and stained as described by Rohde et al. (2004) with slight modifications: Wiesner staining was performed with a drop of a mixture of 1 g of phloroglucinol (Sigma-Aldrich) in 100 mL of 95% (v/v) ethanol and 16 mL of 37% (v/v) HCl. For Mäule staining, the samples were incubated for 5 min in 1% (w/v) KMnO_4 , and after rinsing with water, the samples were incubated in 37% (v/v) HCl, after which a drop of NH_4OH was added to the sample. Images were taken with an Axioskop 2 (Zeiss) microscope. The photographs showing lignin and chlorophyll autofluorescence were taken with an LSM 710 (Zeiss) microscope. The fluorescence signal for lignin was 410 nm with filter 420 to 635 nm, and the fluorescence signal for chlorophyll was 470 nm with filter 650 to 755 nm.

Metabolomics

Approximately 50 mg of frozen ground stem tissue was extracted at room temperature by shaking with 500 μL of methanol. After centrifugation, the supernatant was transferred to new tubes and lyophilized. The pellet was resuspended in equal volumes of cyclohexane and ultrapure water (approximately 50 μL each) such that the portion of initially used stem tissue was 1 mg μL^{-1} aqueous phase. After vortexing, samples were centrifuged, and the aqueous phase was transferred to vials for injection of 15 μL onto the ultra-HPLC-MS system. $n = 8$ for the wild type and *atr2-2*, and $n = 7$ for *atr2-1*. Ultra-HPLC-MS analysis was performed as described by Vanholme et al. (2013c).

Metabolomic Data Processing

From the resulting chromatograms, 1,067 deisotoped peaks (compound ions) were integrated and aligned via Progenesis Q1, each compound ion having an m/z and a retention time. Statistics (ANOVA with posthoc Student's

t test) and principal component analysis (Pareto scaling, with all 1,067 peaks included) was performed in Progenesis Q1 extension EZinfo. Statistical analyses were performed on arcsinh-transformed ion intensities. The following thresholds for significance were used: *P* value for ANOVA < 0.01, and differentially significant *P* < 0.01 in both *atr2-1* and *atr2-2*.

Saccharification

Saccharification was performed according to Van Acker et al. (2013). For alkali pretreatment, 1 mL of NaOH (0.25%, w/v) was added to the dry CWR, and the samples were shaken at 70°C for 3 h. After centrifugation, the pellet was washed three times with ultrapure water, followed by an overnight extraction with 1 mL of 70% ethanol at 55°C and shaking at 750 rpm. After centrifugation, the pellet was washed an additional three times with 1 mL of 70% ethanol and once with 1 mL of acetone.

Lignin Amount via Acetyl Bromide

The determination of the lignin amount was done for eight biological replicates according to Van Acker et al. (2013).

Cellulose Amount

Cellulose was measured on 3 mg of dry stems with sulfuric acid and anthrone in accordance with Foster et al. (2010). Briefly, hemicellulose was removed by adding 1 mL of 2 M trifluoroacetic acid and followed by incubation for 1 h at 99°C while shaking at 750 rpm. After centrifugation at 10,000 rpm, the pellet was washed three times with 0.5 mL of water and twice with 0.5 mL of acetone before being dried and weighed. Next, 1 mL of Updegraff reagent was added to the pellet followed by incubation at 100°C for 30 min. After centrifugation at 10,000 rpm for 15 min, the supernatant was discarded. The pellet was washed three times with 1 mL of acetone. The pellet was left to air dry overnight, and the next day, 175 μ L of 72% (v/v) H₂SO₄ was added. The samples were incubated for 30 min at room temperature, whereupon the samples were vortexed with an additional incubation time of 15 min at room temperature. A total of 825 μ L of water was added, and the samples were vortexed again. The samples were centrifuged at 10,000 rpm for 10 min. Ten microliters was diluted with 90 μ L of water on a 96-well plate. To each sample, 200 μ L of freshly prepared anthrone reagent (2 mg anthrone mL⁻¹ pure H₂SO₄) was added. After incubation at 80°C for 30 min, the samples were cooled on the bench and the absorption was measured at 625 nm.

Lignin Composition via Thioacidolysis

This was performed in accordance with Robinson and Mansfield (2009).

NMR Sample Preparation

The gel samples of whole-plant cell walls for NMR experiments were prepared as described previously (Kim and Ralph, 2010). The dried *Arabidopsis* stems were preground for 1 min in a Retsch MM400 mixer mill at 30 Hz using zirconium dioxide (ZrO₂) vessels (10 mL) containing ZrO₂ ball bearings (2 $\times$ 10 mm). The preground cell walls were extracted with distilled water (ultrasonication, 1 h, three times) and 80% ethanol (ultrasonication, 1 h, three times). Isolated cell walls were dried and ball milled using a Retsch PM100 planetary ball mill at 600 rpm using ZrO₂ vessels (50 mL) containing ZrO₂ ball bearings (10 $\times$ 10 mm). Each sample (100 mg) was ground for 25 min (interval, 5 min; break, 5 min). The cell walls were collected directly in the NMR tubes and formed gels in DMSO-*d*₆:pyridine-*d*₅ (4:1).

NMR Experiments

NMR experiments on the whole-plant cell wall gel samples were performed as described previously (Kim et al., 2008; Kim and Ralph, 2010). NMR spectra were acquired on a Bruker Biospin Avance 500-MHz spectrometer fitted with a cryogenically cooled 5-mm TCI (for triple resonance cryoprobe optimized for ¹H and ¹³C observation) gradient probe with inverse geometry (proton coils closest to the sample). The central DMSO solvent peak was used as an internal reference (δ_C 39.5 ppm, δ_H 2.49 ppm). The ¹³C-¹H correlation experiment was an adiabatic HSQC experiment (Bruker standard pulse sequence hsqcetgpsisp.2;

phase-sensitive gradient-edited two-dimensional HSQC using adiabatic pulses for inversion and refocusing; Kupče and Freeman, 2007). HSQC experiments were carried out using the following parameters: acquired from 10 to 0 ppm in F2 (¹H) with 1,000 data points (acquisition time, 100 ms) and from 200 to 0 ppm in F1 (¹³C) with 400 increments (F1 acquisition time, 8 ms) of 72 scans with a 500-ms interscan delay; the *d*₂₄ (for maximum duration of the chemical shift selective filter with constant time) delay was set to 0.89 ms (1/8J, J: 145 Hz). The total acquisition time was 5 h. Processing used typical matched Gaussian apodization in F2 and squared cosine-bell and one level of linear prediction (32 coefficients) in F1. Volume integration of contours in HSQC plots from data that were not linear predicted used Bruker's TopSpin 3.1 software.

Supplemental Data

The following materials are available in the online version of this article.

Supplemental Figure S1. Phenolic profiling.

Supplemental Figure S2. MS/MS spectra and the reasoning for the tentative structural identification of 33 metabolites of which the MS/MS spectra were not published before.

Supplemental Figure S3. Pathways dependent on CYP450s with metabolites that were detected via ultra-HPLC-MS.

Supplemental Figure S4. Two-dimensional HSQC NMR spectra of the lignin side chain and polysaccharide region of *atr2-1* and *atr2-2* compared with the wild type.

Supplemental Table S1. The 1,067 peaks detected in wild-type plants and *atr2-1* and *atr2-2* mutants.

Supplemental Table S2. The 238 peaks significantly different in abundance in *atr2-1* and *atr2-2* mutants as compared with wild-type plants.

Supplemental Table S3. Overview of primers used for qRT-PCR

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