

Silencing *CHALCONE SYNTHASE* in Maize Impedes the Incorporation of Tricin into Lignin and Increases Lignin Content^{1[OPEN]}

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Lignin is a phenolic heteropolymer that is deposited in secondary-thickened cell walls, where it provides mechanical strength. A recent structural characterization of cell walls from monocot species showed that the flavone triclin is part of the native lignin polymer, where it is hypothesized to initiate lignin chains. In this study, we investigated the consequences of altered triclin levels on lignin structure and cell wall recalcitrance by phenolic profiling, nuclear magnetic resonance, and saccharification assays of the naturally silenced maize (*Zea mays*) *C2-Idf* (*inhibitor diffuse*) mutant, defective in the *CHALCONE SYNTHASE Colorless2* (*C2*) gene. We show that the *C2-Idf* mutant produces highly reduced levels of apigenin- and triclin-related flavonoids, resulting in a strongly reduced incorporation of triclin into the lignin polymer. Moreover, the lignin was enriched in β - β and β -5 units, lending support to the contention that triclin acts to initiate lignin chains and that, in the absence of triclin, more monolignol dimerization reactions occur. In addition, the *C2-Idf* mutation resulted in strikingly higher Klason lignin levels in the leaves. As a consequence, the leaves of *C2-Idf* mutants had significantly reduced saccharification efficiencies compared with those of control plants. These findings are instructive for lignin engineering strategies to improve biomass processing and biochemical production.

Lignocellulosic biomass is a renewable natural resource for the carbon-neutral production of biofuels and biochemicals (Vanholme et al., 2013a; Wilkerson et al., 2014; Welker et al., 2015) that is readily available from agricultural crop residues, inedible plant tissues, or dedicated biomass crops (Abramson et al., 2010). Plant secondary cell walls, which make up the bulk of lignocellulosic biomass, are composed mainly of cellulose, hemicelluloses, and lignin. The cell wall polysaccharide fraction, which constitutes roughly 75% of the total lignocellulosic mass, can be converted into fermentable sugars (Wilke et al., 1981; Vanholme et al., 2013a; Marriott et al., 2016). However, the compact structure and complex chemical composition of the plant cell wall negatively affect the enzymatic digestibility of the biomass, a feature known as biomass recalcitrance (Zhao et al., 2012). Several factors limit the enzymatic degradation, such as carbohydrate-lignin cross-linking and, especially, the presence of the aromatic polymer lignin (Chen and Dixon, 2007; Grabber et al., 2008; Van Acker et al., 2013; Vermerris and Abril, 2015; Wang et al., 2015a). Lignin negatively affects the

conversion of cell wall polysaccharides by immobilizing hydrolytic enzymes and by blocking enzyme access to the polysaccharides (Chundawat et al., 2011; Vanholme et al., 2012). Therefore, decreasing lignin content and changing lignin composition by genetic engineering or breeding are promising strategies to improve biomass-processing properties.

Lignin is a phenolic heteropolymer that is deposited in secondary-thickened cell walls to provide strength and rigidity to specialized cell types (Boerjan et al., 2003; Vanholme et al., 2010a; Cesarino et al., 2012). Lignin is derived from the oxidative radical-radical coupling of three main hydroxycinnamyl alcohol monomers, the monolignols *p*-coumaryl, coniferyl, and sinapyl alcohol, that differ in their degree of aromatic ring methoxylation and couple (as their radicals) in a primarily end-wise manner with the growing polymer radical. Once incorporated into the polymer, these monolignols produce *p*-hydroxyphenyl (H), guaiacyl (G), and syringyl (S) units (Boerjan et al., 2003). In grasses, a large fraction of the coniferyl and sinapyl

alcohol monomers are γ -*O*-acylated with acetate or *p*-coumarate, resulting in acetylated and *p*-coumaroylated units after their polymerization into the lignin polymer (Ralph, 2010).

Besides the canonical monolignols and their acylated analogs, some intermediates of the monolignol biosynthetic pathway, such as *p*-hydroxycinnamaldehydes and *p*-hydroxycinnamates, also may be incorporated into native lignins (Ralph et al., 1994, 2004; Ralph, 1996; Lu and Ralph, 2002; Morreel et al., 2004). The remarkable ability of lignin to incorporate a variety of monomers highlights the inherent malleability of lignification, confirms the original combinatorial chemical coupling theory of lignification (Ralph et al., 2008a), and suggests that genetic engineering of novel lignins is a promising strategy to tailor plants with improved processing properties (Vanholme et al., 2012; Wilkerson et al., 2014; Mottiar et al., 2016).

Recent structural characterization of cell walls from monocot species using NMR showed that the flavone triclin [5,7-dihydroxy-2-(4-hydroxy-3,5-dimethoxyphenyl)-4*H*-chromen-4-one] is part of the native lignin polymer in wheat (*Triticum aestivum*; del Río et al., 2012; Zeng et al., 2013), coconut (*Cocos nucifera*) coir (Rencoret et al., 2013), bamboo (*Phyllostachys pubescens*; Wen et al., 2013), maize (*Zea mays*; Lan et al., 2015), and sugarcane (*Saccharum officinarum*; del Río et al., 2015). Triclin is the first flavonoid that is recognized to be an authentic monomer involved in lignification (del Río et al., 2012; Lan et al., 2015). Moreover, triclin is already well recognized as a valuable health-promoting compound due to its antioxidant, antiaging, anticancer, and cardioprotective potential (Zhou and Ibrahim, 2010; Li et al., 2016). Recent findings show that triclin incorporates into the lignin polymer of maize plants via 4'-*O*- β cross-coupling with normal and acylated monolignols, acting as an initiation site for lignin

polymerization because it can only start a lignin chain (Lan et al., 2015, 2016a). Moreover, the initiation role of triclin has helped explain a long-standing enigma of how the lignin chain is initiated in monocots. In dicots and gymnosperms, the lignin polymer is initiated by monolignol dimerization, giving rise to β -ether, phenylcoumaran, and resinol dimers via β -*O*-4, β -5, and β - β coupling, respectively. In monocots, there is an additional dimeric unit, the tetrahydrofuran structure derived from the β - β coupling of sinapyl *p*-coumarate (Lan et al., 2015, 2016a). Nevertheless, in monocots, lignin only has a relatively low frequency of β -5 and β - β interunit linkages, which could most easily be explained by the alternative chain initiation, involving either triclin (Lan et al., 2015) or (di)ferulates (Ralph et al., 1995; Grabber et al., 2000; Ralph, 2010) or both. However, this causal relationship has not yet been unambiguously proven.

The flux toward flavonoid biosynthesis and, thus, to the triclin monomer is controlled by chalcone synthase (CHS; Fig. 1). This enzyme converts *p*-coumaroyl-CoA into naringenin chalcone, which is then isomerized by chalcone isomerase (CHI) to the flavanone naringenin, a common precursor for several flavonoid classes (Shih et al., 2008; Dixon and Pasinetti, 2010). The biochemical route from naringenin toward triclin in grasses has only recently been fully resolved (Lam et al., 2014, 2015). In rice (*Oryza sativa*), naringenin is desaturated by flavone synthase to form apigenin, after which an extra hydroxyl group is added by flavonoid 3'-hydroxylase to produce luteolin. This hydroxyl group is then methylated by flavonoid *O*-methyltransferase (FOMT) to form chrysoeriol. Another hydroxyl group is added to the same ring by flavonoid 5'-hydroxylase, and the resulting selgin is then again methylated by FOMT to form triclin (Lam et al., 2015). In maize, two genes have been described to encode CHS: *Colorless2* (*C2*) and *Whitepollen1* (*Whp1*; Coe et al., 1981; Franken et al., 1991). The coding regions of *C2* and *Whp1* share 94% sequence identity, but the genes have only partially overlapping expression patterns and are independently regulated (Coe et al., 1981; Franken et al., 1991). The *C2* gene is expressed in many parts of the plant, including the pericarp, the aleurone layer of the endosperm, tassels, and vegetative organs such as ear husks and leaf sheaths, whereas *Whp1* is expressed only in pollen and the aleurone layer of the kernel (Coe et al., 1981; Franken et al., 1991). Based on these expression studies, the *C2* gene is the best candidate for having a role in flavonoid biosynthesis in vegetative tissues, such as stems and leaves of maize. As *C2* directs the flux toward the biosynthesis of all flavonoids, disruption of *C2* would cause the depletion of flavonoids, including triclin, in stems and leaves of maize. Plants carrying the *colorless dominant inhibitory c2* mutation, called *C2-Idf* (*inhibitor diffuse*), have a duplication of the *C2* gene that causes gene silencing (Della Vedova et al., 2005). As a consequence, no *C2*-specific mRNA can be detected in the *C2-Idf* mutants (Franken et al., 1991), and these plants do not produce anthocyanins (Coe et al., 1988).

In this study, we examined the consequences of reduced triclin levels on lignin structure and amount, on biomass processing efficiency, and on the carbon flux into the phenylpropanoid pathway using the maize

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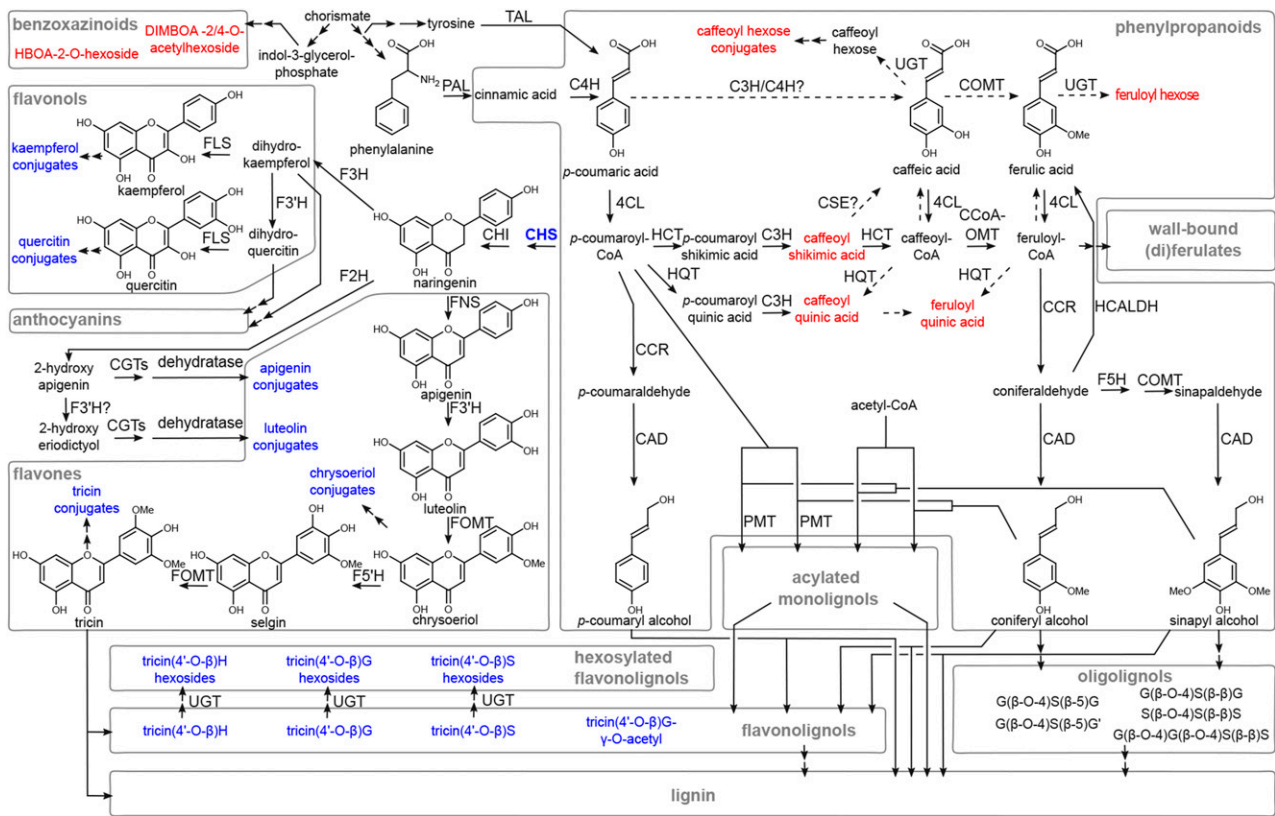


Figure 1. Metabolic map of the phenolic and benzoxazinoid pathways in maize. Solid arrows represent enzymatic conversions for which experimental evidence is available, and dashed arrows represent suggested conversions. Two successive arrows represent two or more metabolic conversions. Metabolic shifts in *C2-Idf* mutants compared with *C2* control plants (Tables II–V) are indicated in color, where red represents a significant increase and blue represents a significant decrease in abundance. Only those metabolic sinks that have been measured in the *C2-Idf* mutants are shown. Reduced biosynthesis of anthocyanins in *C2-Idf* mutants has been reported by Coe et al. (1988). The position of the CHS (*C2*) enzyme is emphasized in a blue boldface font. CAD, Cinnamyl alcohol dehydrogenase; CCoAOMT, caffeoyl-CoA *O*-methyltransferase; CCR, cinnamoyl-CoA reductase; CGT, C-glycosyltransferase; C3H, *p*-coumarate 3-hydroxylase; C4H, cinnamate-4-hydroxylase; CHI, chalcone isomerase; 4CL, 4-coumarate:CoA ligase; COMT, caffeic acid *O*-methyltransferase; CSE, caffeoyl shikimate esterase; F2H, flavanone 2-hydroxylase; F3H, naringenin 3-dioxygenase; F3'H, flavonoid 3'-hydroxylase; F5'H, flavonoid 5'-hydroxylase; FLS, flavonol synthase; FNS, flavone synthase; FOMT, flavonoid *O*-methyltransferase; HCALDH, hydroxycinnamaldehyde dehydrogenase; HCT, *p*-hydroxycinnamoyl-CoA:quinic/shikimate *p*-hydroxycinnamoyltransferase; HQT, hydroxycinnamoyl-CoA:quinic hydroxycinnamoyltransferase; PAL, Phe ammonia lyase; PMT, *p*-coumaroyl-CoA:monolignol transferase; TAL, Tyr ammonia lyase; UGT, UDP-glucosyltransferase. The relation between TAL and PAL activity has been reported by Rösler et al. (1997) for maize and by Cass et al. (2015) for brachypodium. Enzymes involved in the general phenylpropanoid (PAL, TAL, C4H, 4CL, HCT, C3H, and CCoAOMT) and monolignol-specific (CCR, F5H, COMT, and CAD) pathways are described by Guillaumie et al. (2007). Homologs for the Arabidopsis HCALDH (Nair et al., 2004) are present in maize (Končítiková et al., 2015; Missihoun et al., 2016). PMT has been shown in rice (Withers et al., 2012), brachypodium (Petrik et al., 2014), and maize (Marita et al., 2014). The flavone biosynthesis pathway is shown according to Zhou et al. (2008), Cortés-Cruz et al. (2003), and Lam et al. (2014, 2015). The enzymes involved in the metabolic route via F2H toward apigenin and luteolin conjugates has been described in rice, wheat, and buckwheat (*Fagopyrum esculentum*; Brazier-Hicks et al., 2009; Du et al., 2010). The biosynthetic routes via HCT and HQT have been described in switchgrass (Escamilla-Treviño et al., 2014) and suggested to be present in maize (Cortés-Cruz et al., 2003). In addition, HCT of sorghum (*Sorghum bicolor*) has been studied via in vitro activity assays (Walker et al., 2013). The FOMT involved in the conversion of luteolin and selgin into chrysoeriol and tricetin, respectively, in maize is described by Zhou et al. (2008) and is identical to the COMT involved in the conversion of 5-hydroxyconiferinaldehyde to sinapaldehyde described by Collazo et al. (1992), Vignols et al. (1995), and Piquemal et al. (2002). Question marks indicate biosynthetic steps that are questionable; the reaction from *p*-coumaric acid toward caffeic acid via a C3H/C4H complex has so far only been shown by in vitro tests using poplar enzymes (Chen et al., 2011). The activity of CSE has been described in switchgrass (Escamilla-Treviño et al., 2014; Ha et al., 2016), but the absence of close homologs with the reported Arabidopsis CSE (Vanholme et al., 2013b; Ha et al., 2016) makes it questionable whether this reaction is present in maize. For nomenclature of the oligolignols, see Supplemental Table S2.

C2-Idf mutant allele (Franken et al., 1991). The *C2-Idf* allele was in a *B; Pl; R-scm2* W22 background (see “Materials and Methods”). The seeds and vegetative tissues of the *B; Pl; R-scm2* W22 background (*C2* control plants) are purple pigmented, whereas seeds and vegetative tissues of the homozygous *C2-Idf* plants lack the purple pigment and are yellow and green, respectively. Our findings are instructive for cell wall engineering strategies aimed at improving saccharification and biochemical production.

RESULTS

Phenotypic Characterization of the Maize *C2-Idf* Mutant

To determine the residual *C2* mRNA level in the *C2-Idf* mutant, a real-time quantitative PCR (RT-qPCR) analysis was conducted. *C2* expression was measured in the ninth internode of *C2-Idf* and control plants at the developmental stage where the plants had 14 unfolded leaves (i.e. growth stage V14). The expression level of *C2* in stems of *C2-Idf* plants was reduced to 5% of the value in control plants (Fig. 2A), confirming the findings of Franken et al. (1991) that *C2* expression is highly reduced in the *C2-Idf* mutants compared with the *C2* control plants. Seeds and leaf sheaths of the homozygous *C2-Idf* plants lacked the purple anthocyanin pigmentation that is typically observed in *C2* control plants (Fig. 2, B and C).

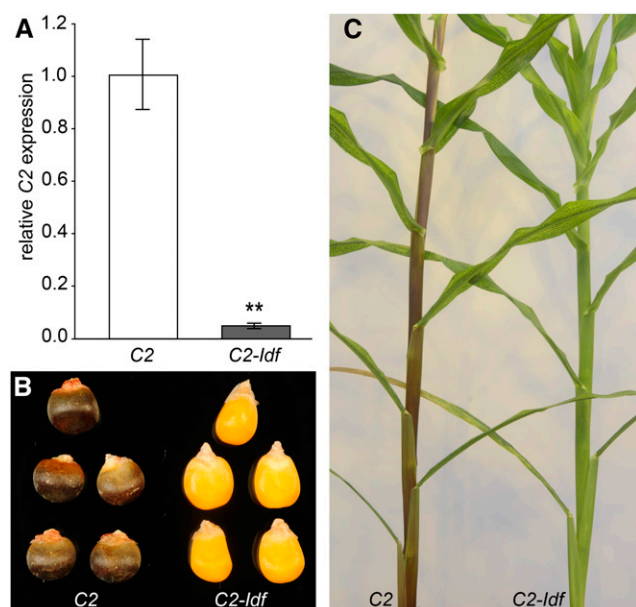


Figure 2. Expression levels of *C2* (*CHS*) and visible *C2-Idf* mutant phenotypes. A, Expression levels of *C2* (*CHS*) in *C2* control and *C2-Idf* mutant internodes were determined by RT-qPCR and are shown relative to the highest *C2* expression value within the replicates. Error bars represent SE of three biological replicates. **, $P < 0.01$. B, Photograph of representative *C2* control and *C2-Idf* mutant seeds. C, Photograph of representative plants at the V14 stage, showing the vegetative tissues.

Next, we investigated the effect of *CHS* down-regulation on plant growth and development. Total plant height, with or without tassel, did not show any significant difference between *C2* control and *C2-Idf* plants (Table I). Also, the length of the ear leaf was not altered significantly. However, the total dry weight of fully senesced *C2-Idf* plants was increased significantly (+18%). The increase could be attributed to the increase in dry weight of the leaves (+27%), as the dry weight of stems was not altered significantly (Table I). To validate the growth phenotype observed in *C2-Idf* maize plants, we grew another set of plants. In agreement with the previous results, the dry weight of fully senesced *C2-Idf* plants was increased significantly compared with the dry weight of *C2* control plants, but the difference was larger (+32%; Supplemental Table S1). However, in this second growth experiment, the increase in dry weight was attributable to a significant increase in dry weight of both stems (+34%) and leaves (+25%; Supplemental Table S1).

CHS-Deficient Plants Have an Altered Phenolic Metabolism

Phenolic profiling of mutants in the lignin biosynthetic pathway has provided extensive insight into which branches of the phenylpropanoid biosynthesis the carbon flux is redirected (Vanholme et al., 2012). To investigate whether flavonoid levels were reduced in the *C2-Idf* mutant, and to reveal which pathways absorb the carbon flux, we performed ultra-high-performance liquid chromatography-mass spectrometry (UHPLC-MS)-based untargeted metabolic profiling (Vanholme et al., 2013b; Sundin et al., 2014) of the ninth internode and the ninth leaf from *C2* control and *C2-Idf* plants. A total of 8,832 peaks could be detected, aligned, and integrated. Only those peaks that were present in all replicates ($n = 7$) of at least one of the four sample types (two genotypes and two tissues) and that were above a 500-count cutoff, regarded as the noise level, were selected. This selection retained 2,005 and 4,073 peaks in internode and leaf samples, respectively, for further analysis. To pinpoint the metabolites for which the abundance was most affected due to the mutation, a selection with more stringent filters was applied based on significance level (Student's *t* test $P < 0.001$), peak intensity (at least 5,000 counts in either genotype), and fold change between *C2-Idf* and *C2* control samples (at least a 10-fold difference in abundance).

In internodes, the 55 peaks with a significantly lower intensity in *C2-Idf* plants compared with the *C2* control could be assigned to 49 compounds (because of in-source fragmentation and adduct formation, a single compound can give rise to multiple mass-to-charge ratio [m/z] peaks), of which 34 could be structurally characterized (Table II). The 28 peaks with a higher intensity could be assigned to 15 compounds, of which nine compounds could be structurally characterized (Table III). The 34 identified compounds that had

Table 1. Growth parameters of *C2* control and *C2-Idf* mutant maize plants

Plant and ear height, leaf length, and dry biomass measurements were performed on senesced greenhouse-grown plants. Leaf 7, The seventh leaf that appeared after germination; ear height, the height of the implantation of the ear on the stem; n.s., not significantly different. *C2* and *C2-Idf* values are means \pm SE ($n = 7$). Underlined values are significantly different compared with the control: *, $0.05 \geq P > 0.01$; and ***, $P < 0.001$.

Growth Parameter	<i>C2</i>	<i>C2-Idf</i>	Percentage Difference	Student's <i>t</i> Test
Plant height with tassel (cm)	268.6 \pm 16.6	259.9 \pm 10.3	-3.2	n.s.
Plant height without tassel (cm)	225.7 \pm 12.0	217.1 \pm 10.6	-3.8	n.s.
Ear height (cm)	123.4 \pm 6.45	121.1 \pm 6.7	-1.9	n.s.
Leaf 7 length (cm)	91.3 \pm 4.9	89.4 \pm 2.1	-2.1	n.s.
Leaf dry weight (g)	34.9 \pm 4.5	<u>44.2 \pm 6.2</u>	26.6	***
Stem dry weight (g)	56.5 \pm 10.45	64.3 \pm 9.2	13.8	n.s.
Total (leaves + stems) dry weight (g)	91.4 \pm 14.3	<u>107.6 \pm 150.1</u>	17.7	*

reduced abundance in *C2-Idf* internodes contained apigenin, tricetin, or kaempferol as a substructure and could be classified into four metabolic classes: flavones, flavonols, hexosylated flavonolignols, and flavonolignols (Table II). Here, we define flavonolignols as 4'-*O*- β -linked coupling products of a flavone and one or more phenylpropanoids. This term is in analogy with the existing term flavonolignan that is used for the low-molecular-mass compounds composed of flavonoid and lignan moieties (Begum et al., 2010) and the term oligolignol that is used for oligomers of canonical monolignols (Morreel et al., 2004, 2010a, 2010b). The fact that some flavonolignols were found without further decoration with sugars and their lack of optical activity suggest that these might be considered as lignin oligomers (Lan et al., 2015, 2016a). Alternatively, they were in the process of being further decorated to become part of the hexosylated flavonolignol pool. The identified flavones and flavonols that were reduced in abundance in *C2-Idf* internodes were decorated with glycerol, pentose, deoxypentose, hexose, hexuronic acid, quinic acid, phenylpropanoids, benzenoids, or a combination of these. Seven of the nine identified compounds that accumulated in *C2-Idf* plants were phenylpropanoids: four had a caffeate moiety and three had a ferulate moiety (Table III). In addition, two benzoxazinoids had an increased abundance in *C2-Idf* internodes: 2-hydroxy-2*H*-1,4-benzoxazin-3(4*H*)-one-2-*O*-hexoside and 2,4-dihydroxy-7-methoxy-2*H*-1,4-benzoxazin-3(4*H*)-one-2/4-*O*-acetylhexoside (Table III).

The 79 peaks with a lower intensity in leaves of *C2-Idf* plants compared with the *C2* control could be assigned to 66 compounds, of which 39 could be structurally characterized (Table IV). The 36 peaks with a higher intensity in leaves could be attributed to 34 compounds, of which nine could be (partially) structurally characterized (Table V). The same classes of compounds that had a reduced abundance in the internode of *C2-Idf* plants also were severely reduced in abundance in their leaves (Table IV). In addition to apigenin- and tricetin-containing flavones, three luteolin- and one chrysoeriol-containing flavones also were present in *C2* control leaves, but their abundances dropped below 10% in leaves of *C2-Idf* plants. In the metabolic classes of flavonolignols and hexosylated flavonolignols,

coupling products with H units were present, in addition to coupling products with G and S units (Table IV). The nine (partially) identified compounds that accumulated in leaves of *C2-Idf* plants all contained a caffeate moiety (Table V). The collective metabolic shifts in *C2-Idf* leaves and stems are visualized in Figure 1.

To investigate whether the decrease in tricetin levels would lead to alternative initiation of the lignin polymer by homodimerization of canonical monolignols, a targeted analysis was carried out for oligolignols coupled β -*O*-4, β -5, or β - β in the phenolic profile of internodes of *C2-Idf* plants. A total of five trilignols and one tetralignol could be identified based on their retention times, *m/z*, and MS/MS spectra (Supplemental Table S2). However, no significant differences were observed between their abundances in *C2* and *C2-Idf* samples for both leaves and internodes.

C2-Idf Plants Are Affected in Lignin Content and Composition

Next, we investigated the impact of *CHS* down-regulation, and thus the lack of tricetin, on the content and composition of cell wall components. The cell wall residue (CWR), cellulose and lignin contents, and lignin composition were determined in senesced stems and leaves of greenhouse-grown *C2-Idf* mutant and *C2* control plants. No differences were found in the CWR per unit of dry weight in leaves and stems (Table V). Measurement of the crystalline cellulose content, expressed as a percentage of the CWR, did not show a significant difference between *C2-Idf* mutant and *C2* control plants in stems, whereas in leaves the crystalline cellulose content was reduced significantly (-9%; Table VI).

The ash content was significantly higher in *C2-Idf* mutant stems (+213%) compared with that of the *C2* control (Table VI). In contrast, the ash content was significantly lower in *C2-Idf* mutant leaves (-16%; Table VI). However, in both stems and leaves, the ash content remained relatively low (i.e. less than 0.1% and less than 1% of the CWR, respectively). The lignin amount in *C2-Idf* and *C2* control plants was determined

Table II. List of compounds that are decreased in abundance in *C2-Idf* internodes

For each compound, its unique number (No.), mass-to-charge ratio (m/z), retention time (RT), peak area (as mean \pm sd), and ratio of the peak area in *C2* and *C2-Idf* are given. inf, Infinite. Tandem mass spectrometry (MS/MS)-based structural characterization is given in Supplemental File S1.

No.	m/z	RT (min)	Name	Peak Area, <i>C2</i>	Peak Area, <i>C2-Idf</i>	Ratio, <i>C2/C2-Idf</i>
Flavones						
1	533.1310	8.60	Apigenin-6C,8C-dipentose	68,464 \pm 24,794	2,551 \pm 1,641	27
2	547.1454	10.00	Apigenin-6C-deoxyhexoside-8C-pentose 1	11,323 \pm 3,720	11 \pm 23	987
3	547.1457	10.63	Apigenin-6C-deoxyhexoside-8C-pentose 2	116,748 \pm 37,294	1,482 \pm 730	79
4	561.1615	11.99	Apigenin-6C,8C-dideoxyhexoside 1	6,708 \pm 1,755	43 \pm 41	156
5	561.1601	12.69	Apigenin-6C,8C-dideoxyhexoside 2	34,529 \pm 8,066	100 \pm 97	346
6	563.1405	7.08	Apigenin-6C-hexoside-8C-pentose	270,510 \pm 91,998	2,207 \pm 1,168	123
7	577.1553	8.36	Apigenin-6C-hexoside-8C-deoxyhexoside 1	17,831 \pm 5,471	17 \pm 29	1,046
8	577.1557	8.88	Apigenin-6C-hexoside-8C-deoxyhexoside 2	75,223 \pm 23,574	162 \pm 167	464
9	577.1556	9.04	Apigenin-6C-hexoside-8C-deoxyhexoside 3	62,832 \pm 19,239	103 \pm 155	613
10	593.1493	5.95	Apigenin-6C,8C-dihexoside	236,212 \pm 79,208	1,328 \pm 498	178
11	491.1213	9.94	Tricin- <i>O</i> -hexoside 1	125,244 \pm 47,891	425 \pm 380	295
12	491.1208	11.40	Tricin- <i>O</i> -hexoside 2	45,980 \pm 14,565	5 \pm 10	9,082
13	403.1057	15.49	Tricin + glycerol	12,669 \pm 4,586	0 \pm 0	inf
14	637.1389	10.87	Tricin + hexuronic acid + pentose	132,255 \pm 47,827	1,577 \pm 1,258	84
15	787.1686	12.34	Tricin + hexuronic acid + pentose + vanillic acid	54,489 \pm 16,951	868 \pm 475	63
16	817.1781	11.70	Tricin + hexuronic acid + pentose + syringic acid	8,932 \pm 2,848	29 \pm 31	310
17	813.1832	13.41	Tricin + hexuronic acid + pentose + ferulic acid	10,840 \pm 2,690	54 \pm 55	199
18	843.1947	12.78	Tricin + hexuronic acid + pentose + sinapic acid	17,337 \pm 4,384	405 \pm 234	43
19	1,009.2540	13.66	Tricin + hexuronic acid + pentose + G(8- <i>O</i> -4)ferulic acid 1	18,550 \pm 4,836	71 \pm 83	262
20	1,009.2537	13.87	Tricin + hexuronic acid + pentose + G(8- <i>O</i> -4)ferulic acid 2	22,561 \pm 6,697	107 \pm 152	212
21	1,039.2637	13.71	Tricin + hexuronic acid + pentose + S(8- <i>O</i> -4)ferulic acid	7,129 \pm 1,928	2 \pm 5	4,106
22	801.1826	13.91	Tricin + hexuronic acid + pentose + 182 D	12,375 \pm 5,013	7 \pm 13	1,881
23	827.1988	15.24	Tricin + hexuronic acid + pentose + 208 D	26,744 \pm 5,908	376 \pm 237	71
24	887.2195	11.60	Tricin + hexuronic acid + pentose + 250 D	21,962 \pm 5,950	135 \pm 50	163
25	667.1479	9.59	Tricin + hexuronic acid + hexose 1	14,943 \pm 4,672	55 \pm 83	272
Flavonols						
26	447.0959	10.14	Kaempferol- <i>O</i> -hexoside	28,991 \pm 16,515	5 \pm 8	6,295
27	593.1499	8.85	Kaempferol- <i>O</i> -hexoside + deoxyhexose	7,781 \pm 2,787	0 \pm 0	inf
Hexosylated flavonolignols						
28	687.1894	12.17	Tricin(4'- <i>O</i> - β)G-5/7- <i>O</i> -hexoside 1	37,436 \pm 13,500	51 \pm 52	731
29	687.1891	12.86	Tricin(4'- <i>O</i> - β)G-5/7- <i>O</i> -hexoside 2	27,043 \pm 12,014	0 \pm 0	inf
30	687.1900	13.59	Tricin(4'- <i>O</i> - β)G-5/7- <i>O</i> -hexoside 3	8,291 \pm 4,223	94 \pm 57	88
31	687.1895	14.18	Tricin(4'- <i>O</i> - β)G- $\gamma/\alpha/4$ - <i>O</i> -hexoside 1	26,948 \pm 11,401	3 \pm 8	8,896
32	717.2028	11.13	Tricin(4'- <i>O</i> - β)S-5/7- <i>O</i> -hexoside	12,748 \pm 4,345	740 \pm 431	17
Flavonolignols						
33	721.2093	16.18	Tricin(4'- <i>O</i> - β)G(4- <i>O</i> - β)G 1	9,700 \pm 2,703	0 \pm 0	inf
34	721.2097	16.88	Tricin(4'- <i>O</i> - β)G(4- <i>O</i> - β)G 2	7,333 \pm 4,487	0 \pm 0	inf

using the Klason method. The acid-soluble lignin of stems and leaves was not affected in the *C2-Idf* mutant, and also the acid-insoluble (Klason) and total lignin of stems did not differ significantly (Table VI). However, the acid-insoluble lignin and total lignin content of leaves of *C2-Idf* plants were significantly higher by +34% and +27% compared with the values in the *C2* control, respectively (Table VI).

Next, analytical thioacidolysis was used to determine possible shifts in lignin composition of stems and leaves from the *C2-Idf* mutant. Thioacidolysis followed by gas chromatography-mass spectrometry provides an estimate of the amount of monolignols linked only by β -*O*-4 bonds. Stem lignin was richer in thioacidolysis-released S units than in G units ($S/G = 3.3$), whereas in leaves, thioacidolysis-released S and G units were in the same range ($S/G = 0.94$). In stems, the relative

abundances of thioacidolysis-released G and S units did not differ between *C2-Idf* plants and the *C2* controls. In contrast, the relative abundance of thioacidolysis-released H units was 24% lower in *C2-Idf* stems compared with the *C2* controls, although the overall frequency of H units was very low (Table VI). When expressed relative to the Klason lignin content, the amount of thioacidolysis-released H units was lower by 11%. In leaves, no significant differences in the relative abundances of H, G, and S units were observed between *C2-Idf* and *C2* control plants (Table VI). However, the amount of thioacidolysis-released G units, S units, and total H + G + S units per Klason lignin were significantly higher (+34%, +38%, and +36%, respectively) in leaves of *C2-Idf* plants. No difference in ferulate content was detected (Ralph et al., 2008b; Table VI).

Table III. List of compounds that are increased in abundance in *C2-Idf* internodes

For each compound, its unique number (No.), *m/z*, retention time (RT), peak area (as mean \pm sd), and ratio of the peak area in *C2-Idf* and *C2* are given. *, Compound detected as an in-source fragment. MS/MS-based structural characterization is given in Supplemental File S1.

No.	<i>m/z</i>	RT (min)	Name	Peak Area, <i>C2</i>	Peak Area, <i>C2-Idf</i>	Ratio, <i>C2-Idf/C2</i>
35	353.0908	2.84	3- <i>O</i> -Caffeoyl quinic acid	62 \pm 104	28,607 \pm 22,141	461
36	353.0808	4.07	5- <i>O</i> -Caffeoyl quinic acid	4,775 \pm 1,626	72,877 \pm 48,103	15
37	335.0800	6.00	Caffeoyl shikimic acid 1	4,478 \pm 2,717	112,871 \pm 86,101	25
38	335.0792	7.50	Caffeoyl shikimic acid 2	170 \pm 319	5,571 \pm 6,418	33
39	355.1037	5.24	Feruloyl hexose	1,133 \pm 247	41,868 \pm 16,125	37
40	193.0503	4.58	3- <i>O</i> -Feruloyl quinic acid*	3,854 \pm 824	43,075 \pm 15,987	11
41	367.1060	6.59	4- <i>O</i> -Feruloyl quinic acid	1,871 \pm 745	52,468 \pm 25,642	28
42	326.0910	4.41	2-Hydroxy-2 <i>H</i> -1, 4-benzoxazin-3(4 <i>H</i>)-one-2- <i>O</i> -hexoside	5 \pm 14	8,934 \pm 3,412	1,740
43	414.1062	7.23	2,4-Dihydroxy-7-methoxy-2 <i>H</i> -1, 4-benzoxazin-3(4 <i>H</i>)-one-2/4- <i>O</i> -acetylhexoside	24 \pm 30	42,132 \pm 1,4371	1,746

Tricin Is Barely Detectable in *C2-Idf* Mutants

In order to evaluate how the down-regulation of *C2* alters lignin structure, 2D-NMR was performed on enzyme lignins from stems and leaves of *C2-Idf* and *C2* control plants. Differences in lignin monomer composition can be visualized from the aromatic regions, whereas the distributions of dimeric units represented by their interunit linkages are deduced from the oxygenated aliphatic regions of the two-dimensional ^1H - ^{13}C correlation heteronuclear single-quantum coherence (HSQC) spectra (Fig. 3; Supplemental Table S3). The colored contours in Figure 3 reflect the different units in the lignin polymer, with the red contours highlighting the triclin units. The spectra from control samples display the typical signals from the commonly observed H, G, and S units, derived from the polymerization of the canonical monolignols (including their acylated analogs). The signals for H units were observed at low levels compared with those corresponding to G and S units. In addition, four distinctive signals corresponding to triclin appeared in the aromatic region of the HSQC spectra. Interestingly, when the lignin monomeric composition was compared in stems and leaves of the control line, large differences could be observed. Whereas the stem lignin appeared to be rich in S units, with minor amounts of triclin (with an integral of 1.9% based on the total of the S and G components; i.e. S + S' + G integrals), leaf lignin had a lower S/G ratio and higher levels of triclin (27.6%; Fig. 3; Supplemental Table S3). Note that end groups (like triclin) are significantly and nonlinearly overestimated in such spectra but provide a qualitative view of the altered levels (Mansfield et al., 2012). Leaf lignin also was richer in H units (10.1%) than stem lignin (1%). It should be noted that attributing the entire integrals from contours in the H region has always been in doubt, as other components appear to contaminate this H peak; it is for this reason that we do not sum S + G + H, as the value becomes too unreliable: all comparisons are made based on the total of the S and G components being set at 100%. Most importantly for this study, upon the strong *CHS* down-regulation found in *C2-Idf* plants, triclin units were undetectable from both stem and leaf lignins

(Fig. 3; Supplemental Table S3) in HSQC spectra. An accurate quantitative analytical method, thioacidolysis of cell wall material followed by liquid chromatography-mass spectrometry (LC-MS) analysis (Lan et al., 2016b), was applied to determine the absolute value of triclin. For the *C2* control samples, the content of triclin was 7.44 mg g⁻¹ total lignin for the stem sample and 18.26 mg g⁻¹ for the leaf (Table VI). For the *C2-Idf* samples, no triclin could be detected in the stem, whereas 0.51 mg g⁻¹ triclin was found in the leaf samples (Table VI). It is also worth pointing out that the percentage of tetrahydrofuran C' units (β - β interunit linkage) increased from 4.7% to 6% in the stems of *C2-Idf* maize and, more strikingly, that in leaves, both β - β interunit linkage types C and C' increased (from 0.3% to 1.2% and from 0% to 0.7%, respectively; Supplemental Table S3).

Lignin M_r Is Mainly Not Changed in *C2-Idf* Plants

Gel permeation chromatography (GPC) analysis was performed to determine the M_r of the acetylated lignin isolated from *C2-Idf* and *C2* control plants. The weight-average (M_w) and number-average (M_n) M_r were estimated from the GPC curves (relative values related to polystyrene standards), and the polydispersity index was calculated. The GPC scans of all samples exhibited three peaks, the first of which was out of the range of the polystyrene standard curve and represented an abnormally large M_r for lignin samples. Therefore, we regarded it as an unknown contaminant (Supplemental Fig. S1). The second peak represented the major lignin polymer, and the third originated from lignin oligolignols containing trilignols, tetralignols, and pentalignols with M_r values at around 800 to 1,000. The M_r values of the main peak of the stem lignin samples (second peak) were 8,300 (M_n) and 24,200 (M_w) for the *C2-Idf*, which was not significantly different from the 8,500 (M_n) and 25,000 (M_w) measured for the *C2* control (Supplemental Fig. S1). The lignin polymer from *C2-Idf* leaves showed 8,400 (M_n), which was not significantly different from the M_r of the lignin from *C2* control leaves with 8,000 (M_n), and 22,900 (M_w), which

Table IV. List of compounds that are decreased in abundance in *C2-Idf* leaves

For each compound, its unique number (No.), *m/z*, retention time (RT), peak area (as mean \pm sd), and ratio of the peak area in *C2* and *C2-Idf* are given. inf, Infinite. MS/MS-based structural characterization is given in Supplemental File S1.

No.	<i>m/z</i>	RT (min)	Name	Peak Area, <i>C2</i>	Peak Area, <i>C2-Idf</i>	Ratio, <i>C2/C2-Idf</i>
Flavones						
2	547.1454	10.00	Apigenin-6C-deoxyhexoside-8C-pentoside	1,770 \pm 3,555	1,054 \pm 268	16
3	547.1457	10.63	Apigenin-6C-deoxyhexose-8C-pentoside	43,830 \pm 10,877	2,118 \pm 452	21
5	561.1601	12.69	Apigenin-6C,8C-dideoxyhexoside 2	8,953 \pm 2,355	538 \pm 309	17
6	563.1405	7.08	Apigenin-6C-hexoside-8C-pentoside	289,600 \pm 64,693	5,024 \pm 1,453	58
7	577.1553	8.36	Apigenin-6C-hexoside-8C-deoxyhexoside 1	31,880 \pm 6,590	293 \pm 133	109
8	577.1557	8.88	Apigenin-6C-hexoside-8C-deoxyhexoside 2	45,078 \pm 9,786	587 \pm 238	77
9	577.1556	9.04	Apigenin-6C-hexoside-8C-deoxyhexoside 3	39,304 \pm 8,518	217 \pm 56	181
10	593.1493	5.95	Apigenin-6C,8C-dihexoside	194,461 \pm 42,585	4,538 \pm 1,082	43
44	593.1502	6.77	Apigenin-C-hexoside + hexose	7,521 \pm 1,510	31 \pm 27	244
45	447.0955	7.02	Luteolin-C-hexoside	30,524 \pm 9,242	1,589 \pm 182	19
46	579.1343	6.15	Luteolin-6C-hexoside-8C-pentoside	13,356 \pm 4,131	2 \pm 4	8,203
47	593.1507	7.13	Luteolin-C-hexoside + deoxyhexose	14,903 \pm 2,980	182 \pm 92	82
48	651.1179	8.50	Chrysoeriol + hexuronic acid + hexuronic acid	13,558 \pm 2,317	1,100 \pm 529	12
11	491.1213	9.94	Tricin-O-hexoside 1	16,812 \pm 4,302	823 \pm 131	20
12	491.1208	11.40	Tricin-O-hexoside 2	17,680 \pm 4,715	495 \pm 153	36
13	403.1057	15.49	Tricin + glycerol	8,511 \pm 2,242	5 \pm 14	1,621
25	667.1479	9.59	Tricin + hexuronic acid + hexose 1	11,573 \pm 3,063	951 \pm 423	12
49	667.1498	9.36	Tricin + hexuronic acid + hexose 2	33,817 \pm 6,894	2,334 \pm 661	14
14	637.1389	10.87	Tricin + hexuronic acid + pentose	20,469 \pm 5,449	1,185 \pm 496	17
17	813.1832	13.41	Tricin + hexuronic acid + pentose + ferulic acid	15,008 \pm 3,821	1,450 \pm 537	10
Flavonols						
27	593.1499	8.85	Kaempferol-O-hexoside + deoxyhexose	136,928 \pm 31,412	199 \pm 41	688
49	609.1452	7.74	Quercetin-O-hexoside + deoxyhexoside	20,081 \pm 3,690	525 \pm 196	38
Hexosylated flavonolignols						
50	657.1798	14.74	Tricin(4'-O- β)H- γ / α /4-O-hexoside	19,127 \pm 4,975	47 \pm 28	408
28	687.1894	12.17	Tricin(4'-O- β)G-5/7-O-hexoside 1	19,765 \pm 5,627	0 \pm 0	inf
29	687.1891	12.86	Tricin(4'-O- β)G-5/7-O-hexoside 2	18,526 \pm 5,906	3 \pm 8	5,993
30	687.1900	13.59	Tricin(4'-O- β)G-5/7-O-hexoside 3	17,549 \pm 5,108	4 \pm 12	3,965
31	687.1895	14.18	Tricin(4'-O- β)G- γ / α /4-O-hexoside 1	68,867 \pm 17,163	11 \pm 15	6,275
51	687.1901	14.80	Tricin(4'-O- β)G- γ / α /4-O-hexoside 2	46,488 \pm 10,726	58 \pm 18	806
Flavonolignols						
52	495.1309	17.65	Tricin(4'-O- β)H 1	11,193 \pm 2,943	4 \pm 4	2,653
53	495.1310	18.36	Tricin(4'-O- β)H 2	12,388 \pm 3,166	3 \pm 4	4,177
54	525.1408	18.01	Tricin(4'-O- β)G 1	93,393 \pm 24,017	17 \pm 17	5,614
55	525.1411	18.77	Tricin(4'-O- β)G 2	75,740 \pm 17,902	2 \pm 5	42,988
56	539.1562	21.74	Tricin(4'-O- β)G- α -O-methyl	6,914 \pm 3,117	10 \pm 27	675
57	567.1497	22.13	Tricin(4'-O- β)G- γ -O-acetyl	9,341 \pm 4,185	2 \pm 6	4,472
33	721.2093	16.18	Tricin(4'-O- β)G(4-O- β)G 1	14,826 \pm 5,233	10 \pm 12	1,558
34	721.2097	16.88	Tricin(4'-O- β)G(4-O- β)G 2	20,322 \pm 6,742	5 \pm 14	3,873
58	721.2097	16.43	Tricin(4'-O- β)G(4-O- β)G 3	9,059 \pm 3,083	0 \pm 0	inf
59	721.2096	17.07	Tricin(4'-O- β)G(4-O- β)G 4	10,052 \pm 3,231	5 \pm 8	2,131
60	721.2096	17.34	Tricin(4'-O- β)G(4-O- β)G 5	22,808 \pm 7,483	1 \pm 3	20,696

appeared to be moderately (8%) higher than that from the *C2* control leaves at 21,100 (Mw; $P = 0.03$; Supplemental Fig. S1).

C2 Down-Regulation Impacts the Saccharification Efficiency of Leaves But Not Stems

Because triclin units are incorporated into native maize lignin, we further evaluated whether the absence of this monomer, as well as its consequences on lignin structure, affect the saccharification efficiency in stems and leaves of *C2-Idf* plants. Therefore, senesced stems and leaves of *C2-Idf* and *C2* control plants were saccharified

for 48 h, using pretreatments with either acid (1 M HCl, 80°C, 2 h) or alkali (62 mM NaOH, 90°C, 3 h), or without pretreatment, after which enzymatic glucose (Glc) release was measured. Subsequently, the cellulose-to-Glc conversion was calculated based on the measured cellulose content and the obtained saccharification yield.

When the stem material was analyzed, no significant differences were found in the saccharification yields expressed on CWR between the *C2-Idf* plants and the *C2* control (Fig. 4, A and B; Supplemental Table S4). Consistent with the fact that the cellulose/CWR did not differ either (Table VI), the cellulose-to-Glc conversion also was found to not differ significantly (Fig. 4B). In

Table V. List of compounds that are increased in abundance in *C2-Idf* leaves

For each compound, its unique number (No.), *m/z*, retention time (RT), peak area (as mean \pm SD), and ratio of the peak area in *C2-Idf* and *C2* are given. *, Compound detected as a trimer. MS/MS-based structural characterization is given in Supplemental File S1.

No.	<i>m/z</i>	RT (min)	Name	Peak Area, <i>C2</i>	Peak Area, <i>C2-Idf</i>	Ratio, <i>C2-Idf/C2</i>
35	1,061.2660	2.84	3- <i>O</i> -Caffeoyl quinic acid*	104 \pm 256	5,436 \pm 2,952	52
61	429.1067	4.84	Caffeoyl hexose glyceric acid	2,387 \pm 480	53,779 \pm 17,531	23
62	445.1372	3.97	Caffeoyl hexose + 104 D	1,658 \pm 496	24,959 \pm 4,946	15
63	457.1377	7.52	Caffeoyl hexose + 116 D 1	956 \pm 408	20,515 \pm 5,574	21
64	457.1376	8.66	Caffeoyl hexose + 116 D 2	2,168 \pm 604	21,900 \pm 5,091	10
65	459.1520	4.83	Caffeoyl hexose + 118 D	1,514 \pm 623	33,623 \pm 4,508	22
66	475.1467	3.72	Caffeoyl hexose + 134 D	3,756 \pm 1,396	72,128 \pm 14,663	19
67	485.1679	11.05	Caffeoyl hexose + 144 D	3,247 \pm 1,219	46,892 \pm 6,757	14
68	537.1647	7.01	Caffeoyl hexose + 196 D	2,036 \pm 701	35,294 \pm 6,598	17

contrast, the saccharification yield was reduced significantly in leaves of *C2-Idf* plants compared with that of the *C2* control under all conditions tested (Fig. 4C; Supplemental Table S4). The differences in Glc release were already observed after 2 h of hydrolysis without pretreatment or following acidic pretreatment (Fig. 4C). Following alkaline pretreatment, significant differences were observed from 4 h of hydrolysis onward (Fig. 4C). The cellulose-to-Glc conversion was calculated for all time points in the hydrolysis. At 48 h of saccharification without pretreatment, the cellulose-to-Glc conversion was reduced significantly, by 39%, in leaves of *C2-Idf* plants compared with the *C2* control, whereas no difference was observed with acidic or

alkaline pretreatment at this time point (Fig. 4D; Supplemental Table S4).

DISCUSSION

The flavone tricrin was recently reported to be part of the lignin polymer in the cell walls of grasses, and it was suggested to act as an initiation site for lignification (del Río et al., 2012; Lan et al., 2015, 2016a), making it a potential target for bioenergy crop improvement. The biosynthesis of tricrin has been described to be accomplished via CHS and CHI activity (Shih et al., 2008; Dixon and Pasinetti, 2010). Based on expression

Table VI. Cell wall composition of senesced greenhouse-grown *C2-Idf* mutant and *C2* control stem and leaf biomass

Stem and leaf values are given as means \pm SD. Lignin monomer composition was determined by thioacidolysis using gas chromatography-mass spectrometry quantitation ($n = 7$ biological replicates). The accurate tricrin level was determined by thioacidolysis using LC-MS quantitation ($n = 6$ biological/2 technical replicates). Acid-insoluble (Klason), soluble, and total lignin were determined by the Klason method ($n = 7$ biological/2 technical replicates). Acid-insoluble and total lignin were corrected for the ash content. Ash content was determined on $n = 4$ biological replicates for *C2* control leaf and *C2-Idf* leaf and stem and $n = 3$ for *C2* control stem. FA, Ferulic acid; FA (EtSH), FA EtSH addition product. Boldface and underlined values indicate significantly increased and decreased values, respectively, compared with those of the control plants. n.s., Not significant; *, $0.05 \geq P > 0.01$; **, $0.01 \geq P > 0.001$; and ***, $P \leq 0.001$.

Cell Wall Parameter	<i>C2</i> Stem	<i>C2-Idf</i> Stem	Percentage Difference	Student's <i>t</i> Test	<i>C2</i> Leaf	<i>C2-Idf</i> Leaf	Percentage Difference	Student's <i>t</i> Test
CWR/dry weight (weight %)	60.72 \pm 6.26	55.76 \pm 4.69	-8.2	n.s.	65.38 \pm 2.79	64.08 \pm 2.5	-2.0	n.s.
Cellulose/CWR (weight %)	33.39 \pm 2.13	33.66 \pm 2.04	-0.8	n.s.	22.62 \pm 1.89	<u>20.65 \pm 0.99</u>	<u>-8.7</u>	*
Ash/CWR (‰)	0.28 \pm 0.25	0.88 \pm 0.24	210	*	8.77 \pm 0.42	<u>7.40 \pm 0.78</u>	<u>-16</u>	*
Acid-insoluble lignin/CWR (%)	18.99 \pm 0.71	18.73 \pm 0.82	-1.4	n.s.	13.75 \pm 1.15	<u>18.47 \pm 0.74</u>	<u>34</u>	***
Acid-soluble lignin/CWR (%)	3.31 \pm 0.23	3.28 \pm 0.15	-0.9	n.s.	4.70 \pm 1.48	5.05 \pm 0.34	7.4	n.s.
Total lignin/CWR (%)	22.30 \pm 0.82	22.01 \pm 0.53	-1.3	n.s.	18.45 \pm 0.93	<u>23.5 \pm 0.76</u>	<u>27</u>	***
H/(H + G + S) (mol %)	1.17 \pm 0.12	<u>0.89 \pm 0.13</u>	<u>-24</u>	***	2.89 \pm 0.24	2.66 \pm 0.85	-8.0	n.s.
G/(H + G + S) (mol %)	23.24 \pm 2.63	25.63 \pm 0.93	-10	n.s.	50.21 \pm 1.41	49.61 \pm 1.33	-1.2	n.s.
S/(H + G + S) (mol %)	75.57 \pm 2.62	73.47 \pm 1.02	-2.8	n.s.	46.89 \pm 1.58	47.72 \pm 1.49	1.8	n.s.
H/total lignin (μ mol mg ⁻¹)	0.025 \pm 0.007	<u>0.022 \pm 0.003</u>	<u>-12</u>	*	0.024 \pm 0.004	0.030 \pm 0.012	25	n.s.
G/total lignin (μ mol mg ⁻¹)	0.50 \pm 0.16	0.64 \pm 0.05	28	n.s.	0.41 \pm 0.06	0.55 \pm 0.08	34	**
S/total lignin (μ mol mg ⁻¹)	1.63 \pm 0.44	1.84 \pm 0.12	13	n.s.	0.38 \pm 0.06	0.53 \pm 0.09	39	**
(H + G + S)/total lignin (μ mol mg ⁻¹)	2.16 \pm 0.59	2.54 \pm 0.17	18	n.s.	0.81 \pm 0.12	1.11 \pm 0.17	37	**
S/G	3.30 \pm 0.54	2.87 \pm 0.14	-13	n.s.	0.94 \pm 0.04	0.96 \pm 0.05	2.0	n.s.
FA/total lignin (μ mol mg ⁻¹)	0.010 \pm 0.001	0.011 \pm 0.001	10	n.s.	0.008 \pm 0.002	0.008 \pm 0.001	4.8	n.s.
FA (EtSH)/total lignin (μ mol mg ⁻¹)	0.014 \pm 0.004	0.017 \pm 0.004	21	n.s.	0.012 \pm 0.002	0.011 \pm 0.002	-8.3	n.s.
Tricrin/(H + G + S) (mol%)	1.05 \pm 0.29	<u>0 \pm 0</u>	<u>-100</u>	***	6.78 \pm 1.42	<u>0.14 \pm 0.04</u>	<u>-98</u>	***
Tricrin/total lignin (mg g ⁻¹)	7.44 \pm 0.85	<u>0 \pm 0</u>	<u>-100</u>	***	18.26 \pm 2.66	<u>0.51 \pm 0.13</u>	<u>-98</u>	***

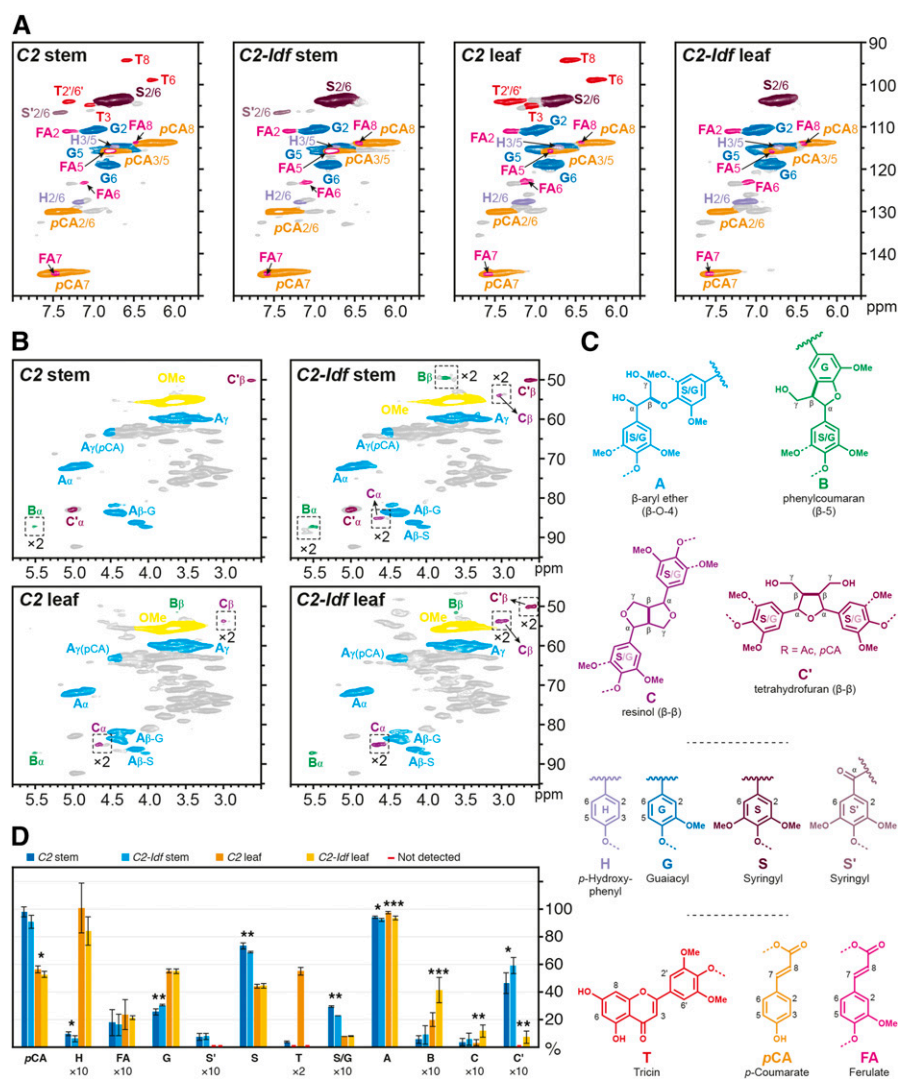


Figure 3. HSQC spectra of the enzymatic lignins from stem and leaf of *C2-ldf* mutant and *C2* control plants. A and B, Spectra of the aromatic regions and the side chain regions, respectively. C, The colors of the substructures shown match those of the corresponding signals in the HSQC spectra. The signal intensities in the framed areas indicated by $\times 2$ are at twice the intensity for convenience. D, The bar graph shows the quantification of the lignin units with their characteristic interunit linkages, with some of them, for convenience, being shown as 2- or 10-fold of their original value, as indicated by $\times 2$ and $\times 10$. For statistics, the leaf and stem material of *C2-ldf* was compared with leaf and stem of the *C2* control, respectively. *, $0.05 < P \leq 0.01$; **, $0.01 < P \leq 0.001$; and ***, $P < 0.001$. For values corresponding to the data points in this graph, see Supplemental Table S3.

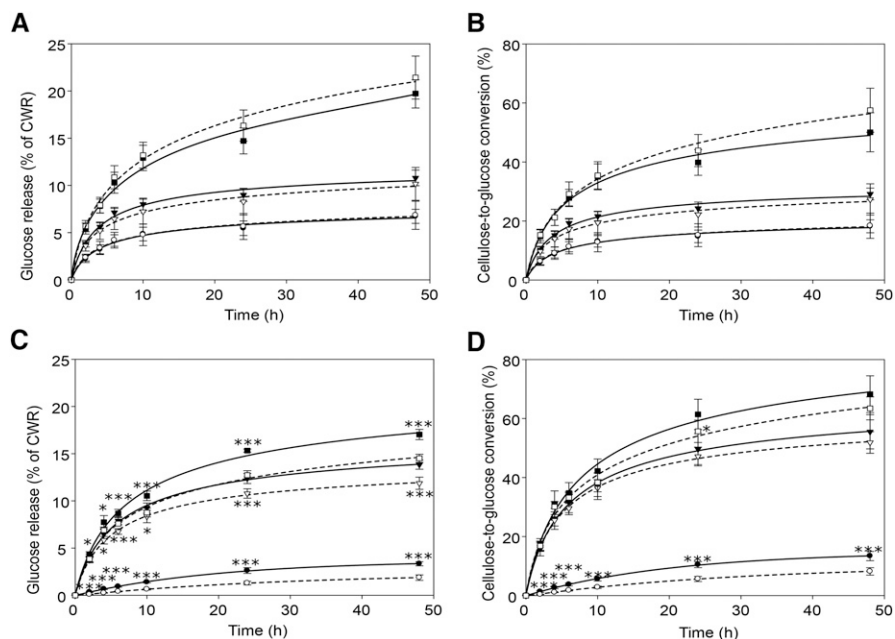
analysis and mutant studies of the two *CHS* homologs in maize (i.e. *C2* and *Whp1*; Coe et al., 1981; Franken et al., 1991), we deduced that *C2* is the most likely gene encoding the flux-determining *CHS* isozyme in lignifying tissues. To investigate whether altering tricetin levels in lignin affects lignin structure, we have studied *C2-ldf* maize plants that have a natural mutation leading to highly reduced *CHS* mRNA levels in stems and leaves.

The RT-qPCR-based expression study showed very low residual expression (5%) of the *C2* gene in internodes of *C2-ldf*, corroborating previously reported northern- and western-blot analyses in which no residual *C2* expression or *C2* protein could be detected in the mutant's aleurone tissue (Franken et al., 1991). Via phenolic profiling of both internodes and leaves of *C2-ldf* mutant and *C2* control plants, we found that disruption of *CHS* leads to a severe reduction in the levels of flavones, flavonols, flavonolignols, and hexosylated flavonolignols. The low residual amount of tricetin, and flavonoids in general, detected by phenolic profiling

suggests that some residual protein is made from the remaining 5% *C2* transcript. The majority of the flavones with a reduced abundance in *C2-ldf* stems and leaves contained an apigenin or tricetin substructure. Additionally, we found luteolin and chrysoeriol derivatives as differential flavones. Our data thus support the recent finding that chrysoeriol, instead of tricetin, is the intermediate in tricetin biosynthesis in grasses (Lam et al., 2015).

Compounds that were increased significantly in abundance in *C2-ldf* plants were mainly caffeate and ferulate derived, esterified by quinic acid, shikimic acid, or hexoses. The accumulation of these compounds implicates detoxification routes for phenylpropanoid pathway intermediates that accumulate upon the perturbation of *CHS*. Remarkably, whereas the identified highly accumulating compounds in *C2-ldf* leaves were all derived from caffeic acid, mostly hexose conjugates that are further derivatized with small M_r adducts, no caffeoyl hexosides appeared on the list of 15 top-accumulating compounds in internodes. Instead, the

Figure 4. Saccharification efficiency of senesced stem (A and B) and leaf (C and D) biomass from *C2* control (solid lines) and *C2-Idf* mutant (dashed lines) plants. The biomass was not pretreated (circles) or was pretreated with either 1 M HCl (triangles) or 62 mM NaOH (squares). The saccharification efficiency is expressed as Glc yield per CWR (A and C) and cellulose-to-Glc conversion (B and D). The numbers corresponding to the data points in these graphs are listed in Supplemental Table S4. Error bars represent SD values of seven biological repeats. *, $0.05 > P \geq 0.01$; and ***, $P < 0.001$.



most accumulating phenylpropanoids in internodes were mainly quinic and shikimic derivatives of caffeic and ferulic acid. Apparently, the carbon fluxes upon the down-regulation of *CHS* are different in leaves and internodes.

In addition, the phenolic profiling revealed that, in *C2* control stems and leaves, triclin was coupled 4'-*O*- β to *p*-coumaryl, coniferyl, and sinapyl alcohols in dimeric and trimeric structures. We recently reported all these flavonolignols (which were previously termed triclin oligolignols) to be present in maize internodes (Lan et al., 2016a). Also, when specifically searching for oligolignols without triclin, we found trilignols and tetralignols coupled via all three types of bonds (β -*O*-4, β - β , and β -5), as expected for combinatorial coupling (Morreel et al., 2010a; Ralph, 2010). This shows that not all monolignols couple in muro to (di)ferulate or triclin nucleation sites but that they also couple mutually to initiate lignin chains. The presence of hexosylated flavonolignols suggests that the aglycone dimers are made in the cytoplasm prior to being glycosylated. Indeed, it was shown in *Arabidopsis* (*Arabidopsis thaliana*) leaf protoplasts that monolignols are not only coupled in the cell wall, as commonly accepted, but also inside the cell, where the monolignol radicals couple combinatorially to produce a plethora of dimers and small lignin oligomers that are subsequently glycosylated and stored in the vacuole (Dima et al., 2015). Our data indicate that triclin also couples through radical coupling in the cytoplasm, as glycosylated derivatives were detected for various triclin-monolignol coupling products.

NMR data (Fig. 3; Supplemental Table S3) showed that phenylcoumaran (β -5; **B**), resinol (β - β ; **C**), and tetrahydrofuran (also β - β , but from acylated monolignols; **C'**) units were elevated in leaves from the *C2-Idf*

mutant. However, in stems, the increase was only significant for the tetrahydrofuran units **C'**. The increased occurrence of β - β units is likely the direct consequence of the lack of triclin in the lignin of these mutants. Triclin can only start a lignin chain and can only do so via 4'-*O*- β coupling with a monolignol or acylated monolignol (Lan et al., 2015, 2016a). In the absence of triclin, the canonical monomers (or their acylated counterparts) will initiate the polymerization more frequently by dimer formation, via β - β , β -5, and β -*O*-4 coupling, if at least one of the monomers is coniferyl alcohol (or its conjugates), or only via β - β coupling, if the monomer is sinapyl alcohol (or its conjugates). Whereas β -*O*-4 and β -5 units **A** and **B** also are made during lignin chain elongation, the β - β units **C** and **C'** can only be part of a starting structure and, in dicots, overwhelmingly derive from sinapyl alcohol and, in grasses, from either sinapyl alcohol or its *p*-coumarate conjugate (Lan et al., 2015, 2016a). Therefore, the increased frequencies of β - β units **C** and **C'**, along with the reduced signals from β -*O*-4 units **A**, are signs of increased monolignol dimerization in the absence of triclin. This finding is in agreement with the fact that the level of β - β units had an opposite trend to the level of triclin units in a series of wheat straw lignin fractions that differed in M_r (Lan et al., 2016b). On the other hand, we cannot exclude the possibility that the shifts in coupling frequencies are a consequence of the increased flux toward monolignols or acylated monolignols in *C2-Idf* mutants, as determined by the increased Klason lignin content in these plants, rather than the absence of triclin per se.

Based on our NMR data of the *C2* control, the lignin from maize leaves contained a higher fraction of triclin units than the lignin of maize stem (with relative integrals of 28% versus 2% indicating a major difference). However, as noted above, end units are nonlinearly

overestimated in HSQC spectra, so analytical thioacidolysis was used to provide accurate quantitation of the level of incorporation (Lan et al., 2016b). Also upon accurate quantitation, the tricetin content was higher in leaves (about 18 mg g⁻¹) than in stems (about 7 mg g⁻¹). The higher levels of tricetin in leaves may be attributable to the biological function of tricetin in protecting plants against insect attack (Bing et al., 2007) and protecting DNA against radiation damage (Stapleton and Walbot, 1994; van de Staaij et al., 2002). In line with the higher tricetin content in leaves, the consequences of the *C2-Idf* mutation were more pronounced in leaves than in stems; the relative fraction of condensed β -5, β - β , and β - β' units was more elevated in *C2-Idf* leaves.

Total lignin content was 27.5% higher in *C2-Idf* leaves compared with that in *C2* leaves. It is likely that the increased lignin content is simply the result of the redirection of the carbon flux toward monolignols upon the blockage of flavonoid biosynthesis. Such a redirection of the flux would have a particularly strong effect in the *C2* background that has a high flux toward flavonoids. Recently, it was reported that stems of *C3H1*-RNAi (for RNA interference) maize plants (which produce less lignin) have increased levels of tricetin, consistent with a redirection of the carbon flux to the synthesis of flavonoids instead of the canonical monolignols (Fornalé et al., 2015). In the leaves of maize *C2-Idf* mutants, the reverse takes place: a substantial redirection of the carbon flux from the biosynthesis of flavonoids to the biosynthesis of the canonical monolignols. Although this scenario seems obvious, it is not generally observed. For instance, the redirection of the flux was observed in leaves but not in stems, possibly because lignifying cells of leaves contain more flavonoids (as leaves are more exposed to environmental stresses) than lignifying cells in the stem, which are largely protected from environmental stresses by the leaf sheaths. Alternatively, the flux was (partially) redirected to lignin in leaves but to other metabolic sinks in stems (e.g. to caffeoyl and feruloyl shikimates), as revealed via phenolic profiling. Furthermore, the *Arabidopsis chs* loss-of-function mutant showed no alterations in lignin content and composition (Li et al., 2010; Vanholme et al., 2010b), and *CHS* suppression in flax (*Linum usitatissimum*) even led to lignin reduction (Zuk et al., 2016). In addition, although an increase in lignin content was observed in fruits of strawberry (*Fragaria* × *ananassa*) silenced for *CHS* via agroinfiltration (Ring et al., 2013), this increase was later assigned to the agroinfiltration method and not to the silencing of *CHS*, because injection of *Agrobacterium tumefaciens* carrying the control vector already increased lignin content (Yeh et al., 2014). Taken together, these combined observations suggest different metabolic plasticity in different plant species and even in different organs upon modified *CHS* expression.

We also observed a significant decrease in cellulose content as a percentage of CWR in leaves of *C2-Idf* plants that might logically be a direct consequence of the relative increase in lignin content. A series of studies

using chemical inhibitors, mutants, or transgenic plants have suggested that cellulose and noncellulosic components of secondary cell walls might be regulated in a compensatory fashion, but the underlying mechanisms remain to be understood. Transgenic poplars (*Populus* spp.) deficient in different steps of the lignin biosynthetic pathway have been reported to have less lignin and relatively more cellulose in their cell walls (Jouanin et al., 2000; Li et al., 2003a; Leplé et al., 2007). However, such an inverse relation between lignin and cellulose was not observed in a set of *Arabidopsis* mutants altered in lignin biosynthesis, in which a reduction in lignin content seemed to be compensated for by a relative increase in hemicellulose content (Vanholme et al., 2012; Van Acker et al., 2013). In grasses, whose cell wall composition and structure differ significantly from those of dicot plants, the same relation (i.e. more hemicellulose in low-lignin plants) has been reported. Stems of low-lignin transgenic switchgrass (*Panicum virgatum*) down-regulated for *COMT* had increased amounts of xylan, the predominant hemicellulose component in switchgrass, both when plants were cultivated in the greenhouse (Fu et al., 2011) and in the field (Baxter et al., 2014). In brachypodium (*Brachypodium distachyon*), the reduced lignin content of a mutant for the lignin-specific laccase *BdLAC5* was compensated for by a slight increase in the content of hemicellulose, whereas the amount of cellulose remained unchanged (Wang et al., 2015b). The *brown midrib* mutants isolated in maize show a characteristic brown coloration of the leaf midrib that is associated with reduced lignin levels and altered lignin composition (Sattler et al., 2010). Compositional analyses of different maize *brown midrib* mutants showed no differences in cellulose content but increased levels of hemicellulose in both stalk and stover (Lechtenberg et al., 1972; El-Tekriti et al., 1976; Cone and Engels, 1993). Also, the inverse relation between lignin and cellulose amount has been described in grasses. Down-regulation of *CCoAOMT* in maize led to reduced lignin content and increased cellulose levels, with no significant differences in hemicellulose levels (Li et al., 2013). The *brittle culm* (*bc*) mutants in rice show reduced tissue mechanical strength due to significant reductions in cellulose content, which is often accompanied by increased levels of lignin, as in the case of *bc1*, *bc10*, and *bc12*, each of which is affected in a different aspect of cellulose deposition (Li et al., 2003b; Zhou et al., 2009; Zhang et al., 2010). Similarly, the reduced cellulose content in mature stems of the spontaneous maize mutant *brittle stalk2* (*bk2*), disrupted in a gene orthologous to the *BC1* gene in rice, goes along with a marked enrichment in lignin deposition, resulting in brittleness of all aerial plant organs (Sindhu et al., 2007). In agreement with its corresponding mutant in rice, *bk2* and wild-type plants were indistinguishable in all aspects of plant growth and development (Ching et al., 2006; Sindhu et al., 2007). In brachypodium, the dwarf *spaghetti1* (*spa1*) mutant presents a unique phenotype combining brittleness with increased elasticity of the internodes (Timpano et al., 2015). The cell walls of *spa1*,

whose causative mutation remains elusive, contain less crystalline cellulose and higher levels of lignin and hemicellulose. However, these compensatory effects are not general. For example, the T-DNA insertion mutant *flexible culm1* in rice, which is deficient in a CAD, shows reduced secondary cell wall thickness and decreased levels of both lignin and cellulose (Li et al., 2009).

Additionally, our results showed that *C2-Idf* plants have an increase in vegetative dry weight, which is a desirable trait for lignocellulosic crops (Feltus and Vandenbrink, 2012). The reason for this increase is not clear. In previous work, the dry weight and stem height measurements of an Arabidopsis *chs* mutant did not differ from those of control plants (Li et al., 2010; Vanholme et al., 2010b). In a different study using the same *chs* mutant, reduced growth was observed versus control plants when grown under adverse environmental growth conditions, such as high light intensity and nitrogen starvation (Misyura et al., 2013), indicating that the growth phenotype is dependent on the environment. *C2-Idf* maize plants might allocate more carbon to biomass production by saving on plant defense compounds (among which flavonoids are made via CHS). Alternatively, some of the differentially abundant compounds might have growth-stimulating properties. However, although the donor line with the *C2-Idf* mutation was backcrossed seven times to the W22 background, we cannot exclude the possibility that residual linkage drag is at the basis of the increased dry weight.

We have shown that stems of *C2-Idf* and *C2* control plants have a similar saccharification efficiency, whereas the leaves of *C2-Idf* plants show a significant decrease in Glc release with and without pretreatment compared with leaves of *C2* control plants. However, the reduction in Glc release in *C2-Idf* leaves was less apparent when the saccharification efficiency was expressed as cellulose-to-Glc conversion. This is because *C2-Idf* leaves also had a lower cellulose content per CWR. Therefore, at least part of the lower Glc release is due to the reduced cellulose content in the leaves of *C2-Idf* plants. It has been established that saccharification efficiency is negatively correlated with lignin amount in both monocot and dicot species (Chen and Dixon, 2007; Sattler et al., 2010; Xu et al., 2011; Bouvier d'Yvoire et al., 2013; Jung et al., 2013; Van Acker et al., 2013; Vanholme et al., 2013a) and that a lower degree of polymerization of lignin eases its extraction from the biomass (Eudes et al., 2012; Vanholme et al., 2012; Shen et al., 2013). The alkaline pretreatment hydrolyzes ester bonds, thereby releasing (di)ferulates that acylate the hemicellulose (and of which a part is also coupled to lignin) and *p*-coumarates that additionally acylate the lignin backbone (Ralph, 2010). Consequently, the alkaline pretreatment results in partial extraction of the lignin from the biomass. The acid pretreatment used, on the other hand, hydrolyzes hemicellulosic glycosidic bonds (Gómez et al., 2014). As (di)ferulates link hemicelluloses with lignin, this acid

pretreatment also will solubilize part of lignin. Thus, both alkaline and acid pretreatments temper the influence of the higher lignin content on the saccharification efficiency. Because the cellulose-to-Glc conversion was not different between *C2-Idf* and *C2* control leaf samples upon acidic or alkaline pretreatment, whereas it was different when no pretreatment was applied, we contend that the 27% increase in total lignin content in leaves of *C2-Idf* plants is the main cause of the reduced cellulose-to-Glc conversion when no pretreatment was used prior to saccharification (Fig. 4D). The saccharification data on stems did not show any significant difference in Glc release, regardless of the pretreatment, in agreement with the observation that lignin content was increased only in leaves. In addition, these results indicate that the shifts in lignin composition in *C2-Idf* stems were too subtle to influence the saccharification efficiency.

Recent findings highlight the metabolic malleability of plant lignification, indicating that lignin can be engineered to dramatically reduce its adverse impact for industrial purposes (Vanholme et al., 2012; Mottiar et al., 2016). For example, the incorporation of coniferyl ferulate into poplar and gramineous feedstock shows (slightly) reduced lignification and allows more efficient delignification and enzymatic hydrolysis of cell walls. The cell walls lignified with coniferyl ferulate were more readily hydrolyzed with fibrolytic enzymes both with and without alkaline pretreatment (Grabber et al., 2008; Wilkerson et al., 2014). As a second example, incorporation of acylated monolignols (mainly coniferyl and sinapyl *p*-coumarate) into the lignin of Arabidopsis resulted in an increased solubilization of these lignins in alkaline medium (Sibout et al., 2016). Our data point to an overall positive relation between the abundance of triclin and saccharification efficiency in leaves. However, this relation is an indirect consequence of the lower lignin amount in *C2* relative to *C2-Idf*. To test whether triclin and the resulting reduced frequency of β - β , β - β' , and β -5 interunit linkages and the increased frequency of β -O-4 interunit linkages have a direct effect on cell wall properties, plants with higher amounts of triclin in their lignin than *C2* plants are needed. The incorporation of triclin into dicot lignins that normally do not produce triclin, or its overproduction in grasses through genetic engineering, may be a promising strategy to test the direct influence of triclin levels in the lignin on lignocellulose recalcitrance. The production of triclin in species that normally do not produce this metabolite has already been achieved in the dicot Arabidopsis (Lam et al., 2015), but its incorporation into lignins and the effect on saccharification have not yet been investigated.

Given the growing consensus that lignin valorization is essential for the environmental sustainability and economics of the lignocellulosic biorefinery (Ragauskas et al., 2014; Rinaldi et al., 2016), high-lignin biomass crops may prove particularly useful if significant value can be extracted from the lignin component. Because flavonoids protect plants from biotic and abiotic

stresses (Stapleton and Walbot, 1994; Falcone Ferreyra et al., 2012), it is unlikely that high-lignin and low-flavonoid *C2-Idf* or *chs* knockout mutants would flourish under field conditions. However, RNA interference-based genetic engineering approaches could be used to mimic *C2-Idf* or *chs* knockout in lignifying cells only (Smith et al., 2013). Leaving the carbon flux toward flavonoids in nonlignifying cell types unaltered, such plants should not necessarily be more sensitive to stresses than their wild-type counterparts. CHS is likely the most elegant step to engineer low-flavonoid and high-lignin characteristics, because CHS is the branch point toward flavonoids; the substrate that accumulates upon CHS deprivation (*p*-coumaroyl-CoA) can readily be used by HCT in the general phenylpropanoid- and monolignol-specific pathways to produce lignin. The strategy to down-regulate *CHS* only in cells that produce monolignols is challenging, because the exact site of monolignol production is not yet known. Depending on whether lignifying cells produce the majority of the monolignols themselves or whether monolignols are produced by neighboring parenchyma cells and then transported toward the cell wall of lignifying cells (Pesquet et al., 2013; Smith et al., 2013), different cell-specific promoters are needed for the silencing approach. In the first case, secondary cell wall-specific promoters (such as promoters of secondary cell wall cellulose biosynthesis or lignin-specific genes; Petrik et al., 2016) could be used to drive the expression of silencing constructs. The thermal conversion of lignocellulosics to renewable gas and bio-oils has been heavily researched, and the benefits for industrial implementation have become clear (Yaman, 2004; Saidi et al., 2014). Feedstocks with high lignin levels might be a promising source of aromatic/phenolic compounds to produce bio-based products from lignin, as proposed in a lignin-first biorefinery approach (Van den Bosch et al., 2015; Rinaldi et al., 2016) and as demonstrated recently with high-level monomer production via hydrogenolysis (Shuai et al., 2016).

MATERIALS AND METHODS

Plant Material

All analyses were carried out with homozygous *C2-Idf* mutant and *C2* control maize (*Zea mays*) plants. Seed stocks of these lines were obtained from the James Birchler laboratory (University of Missouri). The *C2-Idf* allele was in a *B; Pl; R-scm2* W22 background, where *B* stands for the *B-I* allele conditioning pigmentation (Goff et al., 1990), *Pl* stands for the *Pl-Rhoades* (*Pl-Rh*) allele (the *Pl-Rh* allele is highly expressed and produces strong pigmentation in most tissues of the maize plant; Hollick et al., 1995), and *R-scm2* stands for an *R* allele that conditions anthocyanin pigmentation in the embryo and endosperm of the kernel (Styles et al., 1973). The *C2; B; Pl; R-scm2* W22 (*C2* control) was generated by crossing a full-color *C2; b; pl; R-scm2* W22 inbred stock with a *C2; B; Pl; r-g* W22 inbred stock. The two W22 stocks originally trace to materials from the University of Wisconsin. From the cross of these two W22 lines, selection for pigment in the kernel as well as strong pigment in the plant was conducted at each generation of selfing until a true breeding stock with pigment in kernels and plants was selected. This stock carries a normal allele of the *C2* gene. *C2-Idf; B; Pl; R-scm2* W22 (*C2-Idf*) was generated as follows: the *C2-Idf* allele (Maize Genetics Cooperation Stock Center ear 95-1318-3⊗) was backcrossed to the *b; pl; R-scm2* W22 stock for seven generations to introgress it into W22 and then selfed

to produce a true breeding line *C2-Idf; b; pl; R-scm2* W22. Next, this *C2-Idf; b; pl; R-scm2* W22 stock was crossed with the *B; Pl; r-g* W22 stock mentioned above and then self-pollinated over several generations with selection for a true breeding line that was *C2-Idf; B; Pl; R-scm2* W22. The *C2-Idf* mutant and *C2* maize plants were grown in a greenhouse that was kept at 26°C/22°C and 16-h/8-h (day/night) rhythm using a combination of high-pressure sodium vapor lamps (RNP-T/LR/600W/S/230/E40; Radium) and metal halide lamps with quartz burners (HRI-BT/600W/D230/E40; Radium). For expression analysis, six biological replicates of each genotype were grown until the V14 stage. Internode 9 was dissected with removal of the leaf sheath and collected in liquid nitrogen. The samples were then kept at -70°C and subsequently milled with a Grindomix GM200 using precooled stainless-steel jars (Retsch). These samples were pooled two-by-two to obtain three biological replicates. For metabolomics analysis, seven biological replicates of each genotype were grown until the V14 stage, and internode and leaf 9 were harvested and kept at -70°C and subsequently milled as above. Another set of seven biological replicates of each genotype was grown to maturity for growth phenotyping and all cell wall analysis (lignin, cellulose, NMR, GPC, and saccharification). The classification of the growth stages was according to Ritchie et al. (1996), Nafziger (2009), and O'Keeffe (2009). The leaves and internodes were counted from the bottom, as is common for maize. We kept a record of leaf development so as not to miss out on leaves that shriveled up during plant development. No distinction was made between juvenile and adult leaves. Whole leaves were sampled (i.e. including leaf blade, midrib, ligule, and leaf sheath).

Growth Analysis

The final plant height was measured from soil level using a foldable measuring stick. After senescence, stems and leaves from whole plants, excluding the top and bottom 15 cm, were harvested separately for each individual plant and weighed separately. Each sample was milled separately using a cutting mill (Fristch) fitted with a 0.5-mm sieve and used for cell wall characterization and saccharification assays. From the total of seven replicates, six samples were used for NMR analysis (see below).

Gene Expression Analysis

For expression analyses of the *CHS* (*C2*) gene, primers at the 3' end of *CHS* were designed using Primer3 (<http://bioinfo.ut.ee/primer3-0.4.0/primer3/>) with standard settings. The primers were 5'-CAGAAGCGGATCAAGGAGTG-3' and 5'-GGTACATCATGAGGCGGTTTC-3'. The RNA was extracted using an RNeasy kit (Qiagen), and DNase treatment was performed using DNA-free (Ambion, Life Technologies). Extracted RNA was quantified using a Nano-drop ND-1000 spectrophotometer (Thermo Scientific), and 400 ng of RNA was used for cDNA synthesis using the First Strand cDNA Synthesis Kit (Thermo Fisher Scientific). A 10× diluted cDNA sample was used for RT-qPCR using a sensiFAST SYBR No-ROX kit (Bioline) on a LightCycler 480 (Roche). Samples were run in technical triplicates on the LC480 with the following protocol: one activation cycle of 10 min at 95°C; 45 amplification cycles of 10 s at 95°C, 10 s at 60°C, and 10 s at 72°C; and one melting curve cycle measuring from 65°C to 95°C. Fluorescence values were exported from the LightCycler program, whereupon Ct values, normalization factors, and primer efficiencies were calculated according to Ramakers et al. (2003) using *ZmEF1a* and *Zm18S* as reference genes. Primers were 5'-AGTCCGTTGAGATGCACCATG-3' and 5'-CACATACCCACGCTTCAGATCC-3' for *ZmEF1a* and 5'-ACCTTACCAGCCCTTGACATATG-3' and 5'-GACTTGACCAAAACATCTCACGAC-3' for *Zm18S*, as described by Voorend et al. (2016).

Metabolic Profiling

Ground material (250 mg) was extracted after shaking with 7.5 mL of methanol (HPLC grade) at 70°C for 10 min. After centrifugation, methanol was evaporated and samples were suspended in 300 μL of water:cyclohexane (2:1) for extraction. A 15-μL sample of the aqueous phase was subjected to UHPLC-MS on a Waters Acquity UHPLC system (Waters) connected to a Synapt HDMS Q-TOF mass spectrometer (Micromass), basically as described previously (Vanholme et al., 2013b). In brief, chromatographic separation was performed on a Waters Acquity BEH C18 column (2.1 mm × 150 mm, 1.7 μm) with a gradient elution, with the mobile phase composed of water containing 1% acetonitrile and 0.1% formic acid (A) and acetonitrile containing 1% water and 0.1% formic acid (B; all v/v). During the gradient elution, a flow rate of 350 μL

min⁻¹ was applied, with initialization at 0 min, 5% B; 30 min, 50% B; and 33 min, 100% B. Negative mode mass spectrometry setting, chromatogram integration, and alignment via Progenesis QI software (Waters) were as described previously (Vanholme et al., 2013b). The integrated peak area of the *m/z* values (peaks) was normalized with the function Normalize to External Standard, with dry weight (mg) of the pellet remaining after methanol extraction used as the external standard. To select peaks for which metabolites were significantly increased or decreased in *C2-Idf* compared with control plants, the following criteria were used: (1) present in all samples; (2) average normalized abundance higher than 5,000 counts; (3) 10-fold increased/decreased peak area in *C2-Idf* versus control; and (4) Student's *t* test $P < 0.001$. Annotation of compounds matching these criteria was based on accurate *m/z* (± 0.02 D), isotope distribution, and MS/MS similarities. For MS/MS acquisition, a pooled sample of all seven biological replicates per sample type was run on the UHPLC-MS device in both negative and positive ionization mode. The obtained deconvoluted spectra were structurally elucidated based on their similarity with commercially available standards and previously identified metabolites that have already been described in the literature (Supplemental File S1). For targeted analysis of the oligolignols, compounds were identified via *m/z* (± 0.02 D), retention time, and MS/MS similarity.

Statistical Analysis

Statistical analyses consisted of Student's *t* tests for comparisons of *C2-Idf* mutant and *C2* control samples for lignin, cellulose, CWR, and saccharification assays. These analyses were carried out in the software package SPSS Statistics 22 (IBM). For metabolic profiling, the normalized (by dry weight) peak area was first transformed (arcsinh) and then subjected to Student's *t* test (two-tailed distribution based on *F* test) performed in Microsoft Excel Professional Plus 2013.

Purified CWR Analysis

Aliquots (10 mg) of milled stem or leaf material were subjected to a sequential extraction to obtain a purified CWR: water (98°C), ethanol (76°C), chloroform (59°C), and acetone (54°C), 30 min each. The remaining CWR was dried under vacuum.

Lignin and Ash Analysis

Klason lignin was determined using a modified Klason lignin analysis method as described previously (Cullis et al., 2004). Briefly, ground samples were solvent extracted (water [3 × 40 mL], 80% ethanol [3 × 40 mL], and acetone [1 × 40 mL]) and freeze dried for 2 d. Once dried, 0.2 g of dried and extractive-free samples was weighed into test tubes and treated with 3 mL of 72% sulfuric acid for 2 h, with stirring every 10 min. The samples were then diluted to 4% sulfuric acid with 112 mL of nanopure water. The solutions were transferred to flasks and autoclaved at 121°C for 60 min. The samples were filtered through medium-coarseness crucibles (Pyrex) and washed with 200 mL of nanopure water. The retentate was dried at 105°C and then weighed to determine acid-insoluble residue content. Some of the filtrate was collected for acid-soluble lignin analysis. Acid-soluble lignin was determined by measuring the absorbance of the filtrate by a UV spectrophotometer at 205 nm and calculated using an extinction coefficient of 110 L g⁻¹ cm⁻¹. Ash content was determined largely as described by the National Renewable Energy Laboratory in its laboratory analytical procedure "Determination of Ash in Biomass" (<http://www.nrel.gov/docs/gen/fy08/42622.pdf>). In short, acid-insoluble residue was prepared from about 0.3 g of CWR as described above and filtered through a filter paper (Sartorius) using a Buchner filter system (Merck Millipore). Next, the filter paper with the retentate was transferred to a porcelain crucible and placed in a furnace (LT 15/13; Nabertherm) using the following settings: 105°C for 12 min, to 250°C at 10°C min⁻¹, 250°C for 30 min, to 575°C at 20°C min⁻¹, 575°C for 180 min, and finally drop to 105°C. The ash content was determined by weighing the crucible and ash on an analytical balance (XPE105; Mettler-Toledo) and correcting for the empty crucible and the ash content of an empty filter paper. The acid-insoluble lignin (Klason lignin) was then calculating by subtracting the ash content from the acid-insoluble residue. The lignin composition was investigated via thioacidolysis similar to Rolando et al. (1992), but on 10 mg of CWR and a 1-mL reaction mixture (0.025 mL of boron trifluoride etherate + 0.1 mL of ethanethiol + 0.875 mL of dioxane).

Cellulose Analysis

Aliquots (5 mg) of milled stem or leaf material were subjected to a sequential extraction to obtain a purified CWR, as described above. Cellulose was measured on 5 mg of dry stems with sulfuric acid and anthrone according to Foster et al. (2010). The CWR was incubated with 2 M trifluoroacetic acid and 20 μL of inositol (5 mg mL⁻¹) for 2 h at 99°C while shaking (750 rpm). After incubation, the remaining pellet was washed three times with water and twice with acetone and dried under vacuum. H₂SO₄ (175 μL of 72% [v/v]) was added to the dried pellet. The samples were vortexed 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 μL 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 μL of water on a 96-well plate. To each sample, 200 μL of freshly prepared anthrone reagent (2 mg anthrone mL⁻¹ 72% 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.

Tricin Content Analysis

Tricin was determined by analytical thioacidolysis essentially as published (Lapierre et al., 1986) but using LC-MS detection as described (Lan et al., 2016b).

Preparation of Enzyme Lignin for NMR and GPC

Isolated maize stem and leaf samples (1 g, ground to 60–80 mesh) were successively extracted with water and 80% ethanol (5 × 45 mL for each) under ultrasonication to remove the soluble extractives. The extractive-free materials (approximately 500 mg) were ball milled for 1 h (5-min interval break every 5 min). The ball-milled material was then treated with crude cellulases (*Trichoderma viride*; Calbiochem) at 35°C (3 × 48 h) to saccharify most of the carbohydrates, producing a lignin-enriched fraction containing essentially all of the original lignin. The enzyme lignins from maize stem and leaf were obtained as brown powders after lyophilization.

GPC Analysis

The obtained enzyme lignin was first acetylated (10 mg) in 0.5 mL of pyridine:acetate anhydride (2:1, v/v; 10 mL) at room temperature for 12 h. Acetylated maize lignin was obtained after coevaporation of the solvent with ethanol under reduced pressure at 45°C until pyridine and acetic anhydride were completely removed. The acetylated product was dissolved in *N,N*-dimethylformamide (0.1 M lithium bromide) for GPC acquisition. Analytical GPC was performed on a Shimadzu LC20 device with a photodiode array detector (SPD-M20A). Lithium bromide (0.1 M) in *N,N*-dimethylformamide was used as an eluent at 40°C with a flow rate of 0.3 mL min⁻¹. Polystyrene (low molecular) Standard ReadyCal Set M(p) 250–70,000 (P/N 76552; Fluka) was used to calibrate and calculate the *M_r* distribution of the lignin samples. Separation was performed using a series of three columns, first a TSKgel guard column with α-stationary phase (6 mm × 4 cm, 13 μm; TOSOH Bioscience), then a TSKgel Alpha-2500 column (7.8 mm × 30 cm, 13 μm; TOSOH Bioscience), followed by a TSKgel Alpha-M column (7.8 mm × 30 cm, 13 μm; TOSOH Bioscience).

NMR Characterization of Lignin Samples

Enzyme lignins (20–30 mg) were dissolved in 1 mL of dimethyl sulfoxide-*d*₆:pyridine-*d*₅ (4:1, v/v) in an NMR tube. NMR spectra were recorded on a Bruker Biospin (Billerica) AVANCE 700-MHz spectrometer fitted with a cryogenically cooled 5-mm ¹H-optimized resonance (¹H/³¹P/¹³C/¹⁵N; QCI) gradient probe with inverse geometry (proton coils closest to the sample). Bruker's Topspin 3.5 (Mac) software was used to process spectra. The central dimethyl sulfoxide-*d*₆ solvent peaks were used as internal references ($\delta_{\text{C}}\text{-}\delta_{\text{H}}$, 39.51:2.49). The HSQC experiment used a Bruker standard pulse sequence, hsqcqp2psp2, with the following parameters: acquired from 10 to 0 ppm in F2 (¹H) with 200-ms acquisition time, 200 to 0 ppm in F1 (¹³C) with 8-ms acquisition time of 32 scans; the *d*₂₄ delay was set to 0.86 ms (1/8J, J = 145 Hz); and the relaxation delay was 1 s. The total acquisition time for a sample was approximately 6 h. Processing used typical matched Gaussian apodization (GB = 0.001, LB = -0.3) in F2 and squared cosine-bell apodization in F1. The quantitation of aromatic and side chain distributions was performed based on the volume integration of contours in HSQC spectra. For aromatics, the percentages of H units, ferulate (FA), *p*-coumarate (*p*CA), and triclin (T) were calculated based on the total of G, S, and oxidized syringyl (S') units (i.e. on the total prototypical G + S aromatic lignin units). For the aliphatic region, the

percentages of cinnamaldehyde (X2) and benzaldehyde (X3) end groups were calculated based on the sum of the normal units: β -aryl ether (β -O-4; A), phenylcoumaran (β -5; B), resinol (β - β ; C), and tetrahydrofuran (β - β ; C'). The correlations from G2, S2/6, S'2/6 (logically halved for S and S'), H2/6 (halved), FA2, pCA2/6 (halved), T6 in the aromatic region, and A α , B α , C β , C' β , X2 γ , and X3 α in the aliphatic region were used for integral calculations. Note that these are uncorrected integrals and are only comparatively accurate, not absolute quantifications; end units, in particular (and including tricin), are overquantitated compared with polymer internal units in these spectra (Mansfield et al., 2012).

Saccharification Assay

Aliquots (10 mg) of dry stem or leaf material were used. The biomass was pre-treated with 1 mL of either 1 M HCl (80°C, 2 h) or 62 mM NaOH (90°C, 3 h) while shaking (850 rpm). The supernatant was removed, and the pellet containing pre-treated material was washed three times with water to obtain a neutral pH. Subsequently, the saccharification assay was performed as described by Van Acker et al. (2013, 2016). The enzyme mix added contained cellulase from *Trichoderma reesei* ATCC 26921 and cellobiase from *Aspergillus niger* (Sigma-Aldrich) in a 5:3 ratio. Both enzymes were first desalted over an Econo-Pac 10DG column (Bio-Rad) stacked with Bio-Gel P-6DG gel (Bio-Rad) according to the manufacturer's guidelines. The desalted cellobiase was 350-fold diluted prior to mixing with desalted cellulase. The enzyme mix was further diluted 10-fold, and the activity of the diluted enzyme mix was measured with a filter paper assay (Xiao et al., 2004). To each sample, the enzyme mix with an activity of 0.014 filter paper units, dissolved in acetic acid buffer (pH 4.8), was added. Aliquots (20 μ L) were taken after 0, 2, 6, 10, 24, and 48 h of incubation at 50°C and 10-fold diluted with acetic acid buffer (pH 4.8). The concentration of Glc was measured as described by Van Acker et al. (2013, 2016).

Supplemental Data

The following supplemental materials are available.

Supplemental Figure S1. GPC comparison of lignin from *C2-ldf* mutant and *C2* control stem and leaf.

Supplemental Table S1. Growth parameters of *C2-ldf* mutant and *C2* control plants.

Supplemental Table S2. Targeted search for oligolignols isolated from stem and leaf of *C2-ldf* and *C2* control plants.

Supplemental Table S3. Relative quantification of lignin units and linkages based on volume integrals from the HSQC spectra of stem and leaf of *C2-ldf* mutant and *C2* control plants.

Supplemental Table S4. Saccharification efficiency of senesced stem and leaf biomass from *C2-ldf* mutant and *C2* control plants.

Supplemental File S1. MS/MS spectra of the characterized metabolites.

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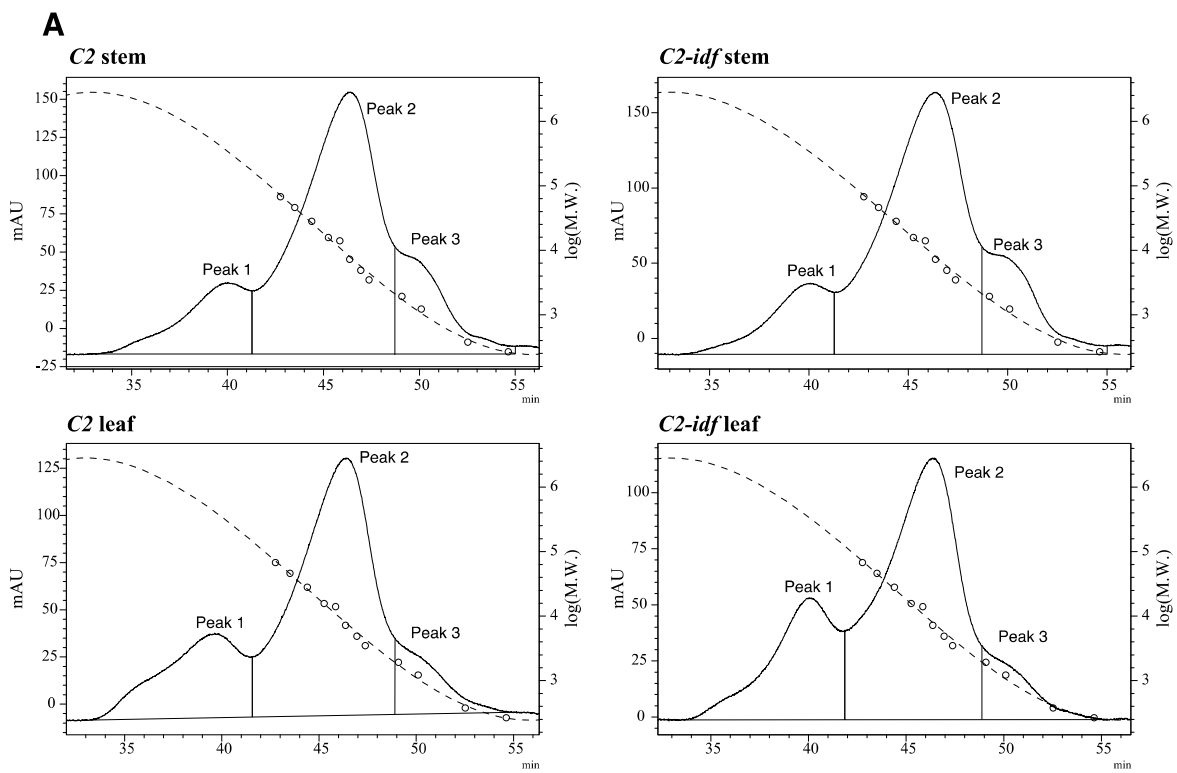
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5 **Supplemental Figure S1. GPC chromatographic comparison of lignin from *C2-Idf* mutant and *C2***
6 **control stem and leaf. (A)** The dashed line shows the standard curve ($R^2=0.9962$) of polystyrene with
7 12 data points. The molecular weight distribution curves were each divided into three peaks and the
8 Mn, Mw, and polydispersity index (Mw/Mn) of each peak were analyzed and shown in the table (B).
9 Peak 1 represent an abnormal large molecular weight for lignin samples and was therefore regarded
10 as an unknown contaminant, the peak 2 represented the major lignin polymer and peak 3 originated
11 from lignin oligolignols with molecular weight at around 800-1000. Statistics were performed on
12 peak 2. The Mn, Mw and Mw/Mn values of stem and the Mn value of leaf were not significantly
13 different between the *C2-Idf* mutant and *C2* control (Student t-test, p-value>0.05). The Mw and
14 Mw/Mn values of leaf were significantly different between the *C2-Idf* mutant and *C2* control (8%
15 increase in *C2-Idf* as compared to *C2*, Student t-test, p-value=0.03 for Mw, and 4% increase, Student
16 t-test, p-value=0.04 for Mn/Mw). Average \pm standard deviation are given for 4 to 6 biological
17 replicates.



B

<i>C2</i> stem			<i>C2-Idf</i> stem				
	Mn	Mw	Mw/Mn		Mn	Mw	Mw/Mn
Peak 1	421200 \pm 37000	720000 \pm 66500	1.71 \pm 0.03	Peak 1	477100 \pm 55300	812300 \pm 112500	1.70 \pm 0.08
Peak 2	8500 \pm 300	25000 \pm 1400	2.95 \pm 0.06	Peak 2	8300 \pm 300	24200 \pm 1400	2.92 \pm 0.06
Peak 3	810 \pm 50	1100 \pm 30	1.34 \pm 0.04	Peak 3	900 \pm 80	1100 \pm 50	1.29 \pm 0.06
<i>C2</i> leaf			<i>C2-Idf</i> leaf				
	Mn	Mw	Mw/Mn		Mn	Mw	Mw/Mn
Peak 1	507100 \pm 23700	891700 \pm 42200	1.76 \pm 0.05	Peak 1	405300 \pm 76700	710200 \pm 144600	1.75 \pm 0.05
Peak 2	8000 \pm 300	21100 \pm 1000	2.62 \pm 0.05	Peak 2	8400 \pm 300	22900 \pm 1100	2.71 \pm 0.07
Peak 3	800 \pm 20	1100 \pm 10	1.33 \pm 0.02	Peak 3	800 \pm 80	1000 \pm 60	1.38 \pm 0.06

19 **Supplemental Table S1. Growth parameters of *C2-ldf* mutant and *C2* control plants.** Plant height and dry biomass measurements were performed on
 20 senesced greenhouse-grown plants. DW: dry weight; SEM: standard error of the mean with n=12; n.s.: not significantly different; underlined values:
 21 significantly different compared to the control; *:0.05≥p>0.01, **: p<0.01.

22

Growth parameter	<i>C2</i> (mean ± SEM)	<i>C2-ldf</i> (mean ± SEM)	% difference	t-test
Plant height with tassel (cm)	296.33 ± 17.78	292.01 ± 19.01	-1.4	n.s.
Leaf DW (g)	36.11 ± 6.71	<u>45.13</u> ± <u>6.20</u>	25	*
Stem DW (g)	90.79 ± 31.45	<u>121.80</u> ± <u>28.08</u>	34.16	**
Total DW (g) (leaves + stems)	126.90 ± 37.76	<u>166.94</u> ± <u>33.94</u>	31.55	**

23

24 **Supplemental Table S2. Targeted search for oligolignols isolated from stem and leaf of *C2-ldf* and *C2* control plants.** b.d.: below detection limit; SD:
 25 standard deviation, p-values are calculated via Student t-tests statistics. Nomenclature of the oligolignols is explained in Morreel et al., (2004; 2010a;
 26 2010b), the prime (') signifies an aldehyde moiety.

27

nr	Name	[M-H] ⁺	RT (min)	<i>C2</i> internode (Mean ± SD)	<i>C2-ldf</i> internode (Mean ± SD)	p-value	Ratio <i>C2-ldf</i> / <i>C2</i>	<i>C2</i> leaf (Mean ± SD)	<i>C2-ldf</i> leaf (Mean ± SD)	p- value	Ratio <i>C2-ldf</i> / <i>C2</i>
1	G(8-O-4)S(8-5)G 1	583.2180	14.81	29243 ± 7109	39767 ± 20579	0.267	1.36	2539 ± 934	2162 ± 896	0.442	0.85
2	G(8-O-4)S(8-5)G 2	583.2180	15.61	13912 ± 3862	14941 ± 9379	0.903	1.07	1411 ± 648	887 ± 640	0.125	0.63
3	G(8-O-4)S(8-8)G	583.2194	17.04	2494 ± 653	4590 ± 3247	0.110	1.84	b.d.	b.d.	-	-
4	G(8-O-4)S(8-5)G'	581.2034	16.93	2172 ± 4397	1062 ± 826	0.611	0.49	2011 ± 546	1889 ± 1202	0.372	0.94
5	S(8-O-4)S(8-8)S	643.2390	16.54	713 ± 398	984 ± 727	0.465	1.38	3233 ± 2903	2791 ± 742	0.645	0.86
6	G(8-O-4)G(8-O-4)S(8-8)S	809.3000	18.09	2153 ± 1182	4997 ± 3693	0.083	2.32	1110 ± 327	1415 ± 646	0.439	1.27

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29

30 **Supplemental Table S3. Relative quantification of lignin units and linkages based on volume integrals from the HSQC spectra of stem and leaf of C2-Idf**
 31 **mutant and C2 control plants.** The lignin units were expressed as a percentage of the sum of G (guaiacyl), S' (benzylic-oxidized S units, or syringaldehyde- or
 32 sinapaldehyde-derived), and S (syringyl) units. The linkage types were expressed as percentage of the sum of interunit linkage types A-C (including C'). Units
 33 relatively quantified (as volume integrals only) include: cinnamaldehyde end-groups X2; benzaldehyde end-groups X3, phenylcoumaran (β -5) units B, β -aryl
 34 ether (β -O-4) units A; resinol (β - β) units C; and β - β' tetrahydrofuran units C'. n= 6 biological replicates. **Bold or underlined values indicate significantly**
 35 **increased or decreased values, respectively, as compared to those of the control plants.**, pCA: p-coumarate, H: p-hydroxyphenyl unit, FA: ferulate, T: triclin
 36 unit. SD: standard deviation, inf: infinite, n.d.: not detected, -: not applicable, n.s.: not significant, *:0.05<p≤0.01, **: 0.01<p≤0.001, ***: p<0.001.

feature	C2 stem (mean ± SD)	C2-Idf stem (mean ± SD)	% difference	t-test	C2 leaf (mean ± SD)	C2-Idf leaf (mean ± SD)	% difference	t- test
pCA%	98.2 ± 3.7	91.2 ± 4.8	-9.1	n.s	56.5 ± 2.7	<u>52.8 ± 2.3</u>	<u>-6.5</u>	*
H%	1.0 ± 0.1	<u>0.6 ± 0.2</u>	<u>-43.3</u>	*	10.1 ± 1.8	8.5 ± 1.0	-16.4	n.s.
FA%	1.8 ± 0.9	1.6 ± 0.8	-9.5	n.s.	2.4 ± 1.1	2.2 ± 0.2	-9.3	n.s.
G%	25.6 ± 2.3	30.4 ± 0.5	18.7	**	55.6 ± 1.4	55.3 ± 1.7	-0.6	n.s
S'%	0.8 ± 0.3	0.8 ± 0.2	4.0	n.s.	n.d.	n.d.	-	-
S%	73.6 ± 2.2	<u>68.8 ± 0.6</u>	<u>-6.6</u>	**	44.4 ± 1.4	44.7 ± 1.7	0.7	n.s
T%	1.9 ± 0.3	n.d.	<u>inf down</u>	***	27.6 ± 1.4	<u>n.d.</u>	<u>inf down</u>	***
Total (G+S+S')	100	100			100	100		
S/G	3.0 ± 0.5	<u>2.3 ± 0.1</u>	<u>-20.7</u>	**	0.8 ± 0.1	0.8 ± 0.1	1.3	n.s.
X2 (cinnamaldehyde end-group)%	0.9 ± 0.4	1.0 ± 0.3	13.5	n.s.	n.d.	0.4 ± 0.3	inf up	**
X3 (benzaldehyde end-group)%	- ± -	0.1 ± 0.1	66.6	n.s.	n.d.	n.d.	-	-
B (β -5)%	0.6 ± 0.3	0.9 ± 0.6	57.9	n.s.	2.0 ± 0.5	4.2 ± 0.9	109.6	***
A (β -O-4)%	94.4 ± 0.9	<u>92.6 ± 1.2</u>	<u>-2.0</u>	*	97.7 ± 0.7	<u>93.9 ± 1.5</u>	<u>-3.9</u>	***
C (β - β , resinol)%	0.3 ± 0.3	0.6 ± 0.6	67.6	n.s.	0.3 ± 0.3	1.2 ± 0.5	310.3	**
C' (β - β , tetrahydrofuran)%	4.7 ± 0.7	6.0 ± 0.6	27.9	*	n.d.	0.7 ± 0.5	inf up	***
Total (A+B+C+C')	100	100			100	100		

37

38

39 **Supplemental Table S4. Saccharification efficiency of senesced stem and leaf biomass from *C2-Idf* mutant and *C2* control plants.** The numbers listed in this
40 table are visualized in the graphs of Figure 4.

41 See separate file, Supplemental Table S4.xlsx

42

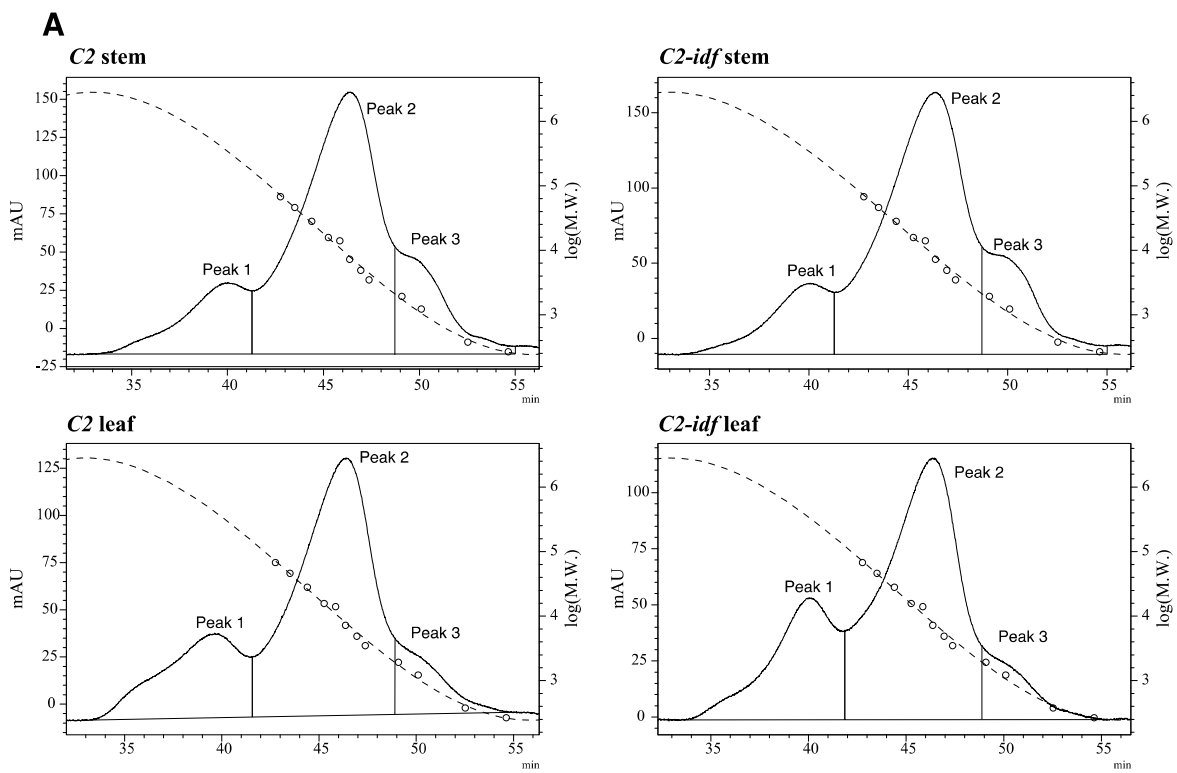
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Leaf DW (g)	36.11	± 6.71	<u>45.13</u>	± <u>6.20</u>	25	*
Stem DW (g)	90.79	± 31.45	<u>121.80</u>	± <u>28.08</u>	34.16	**
Total DW (g) (leaves + stems)	126.90	± 37.76	<u>166.94</u>	± <u>33.94</u>	31.55	**

23

24 **Supplemental Table S2. Targeted search for oligolignols isolated from stem and leaf of C2-Idf and C2 control plants.** b.d.: below detection limit; SD:
 25 standard deviation, p-values are calculated via Student t-tests statistics. Nomenclature of the oligolignols is explained in Morreel et al., (2004; 2010a;
 26 2010b), the prime (') signifies an aldehyde moiety.

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nr	Name	[M-H] ⁺	RT (min)	C2 internode (Mean ± SD)	C2-Idf internode (Mean ± SD)	p-value	Ratio C2-Idf/C2	C2 leaf (Mean ± SD)	C2-Idf leaf (Mean ± SD)	p- value	Ratio C2- Idf/C2
1	G(8-O-4)S(8-5)G 1	583.2180	14.81	29243 ± 7109	39767 ± 20579	0.267	1.36	2539 ± 934	2162 ± 896	0.442	0.85
2	G(8-O-4)S(8-5)G 2	583.2180	15.61	13912 ± 3862	14941 ± 9379	0.903	1.07	1411 ± 648	887 ± 640	0.125	0.63
3	G(8-O-4)S(8-8)G	583.2194	17.04	2494 ± 653	4590 ± 3247	0.110	1.84	b.d.	b.d.	-	-
4	G(8-O-4)S(8-5)G'	581.2034	16.93	2172 ± 4397	1062 ± 826	0.611	0.49	2011 ± 546	1889 ± 1202	0.372	0.94
5	S(8-O-4)S(8-8)S	643.2390	16.54	713 ± 398	984 ± 727	0.465	1.38	3233 ± 2903	2791 ± 742	0.645	0.86
6	G(8-O-4)G(8-O-4)S(8-8)S	809.3000	18.09	2153 ± 1182	4997 ± 3693	0.083	2.32	1110 ± 327	1415 ± 646	0.439	1.27

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30 **Supplemental Table S3. Relative quantification of lignin units and linkages based on volume integrals from the HSQC spectra of stem and leaf of C2-Idf**
 31 **mutant and C2 control plants.** The lignin units were expressed as a percentage of the sum of G (guaiacyl), S' (benzylic-oxidized S units, or syringaldehyde- or
 32 sinapaldehyde-derived), and S (syringyl) units. The linkage types were expressed as percentage of the sum of interunit linkage types A-C (including C'). Units
 33 relatively quantified (as volume integrals only) include: cinnamaldehyde end-groups X2; benzaldehyde end-groups X3, phenylcoumaran (β -5) units B, β -aryl
 34 ether (β -O-4) units A; resinol (β - β) units C; and β - β' tetrahydrofuran units C'. n= 6 biological replicates. **Bold or underlined values indicate significantly**
 35 **increased or decreased values, respectively, as compared to those of the control plants.**, pCA: p-coumarate, H: p-hydroxyphenyl unit, FA: ferulate, T: triclin
 36 unit. SD: standard deviation, inf: infinite, n.d.: not detected, -: not applicable, n.s.: not significant, *:0.05<p≤0.01, **: 0.01<p≤0.001, ***: p<0.001.

feature	C2 stem (mean ± SD)	C2-Idf stem (mean ± SD)	% difference	t-test	C2 leaf (mean ± SD)	C2-Idf leaf (mean ± SD)	% difference	t- test
pCA%	98.2 ± 3.7	91.2 ± 4.8	-9.1	n.s	56.5 ± 2.7	<u>52.8 ± 2.3</u>	<u>-6.5</u>	*
H%	1.0 ± 0.1	<u>0.6 ± 0.2</u>	<u>-43.3</u>	*	10.1 ± 1.8	8.5 ± 1.0	-16.4	n.s.
FA%	1.8 ± 0.9	1.6 ± 0.8	-9.5	n.s.	2.4 ± 1.1	2.2 ± 0.2	-9.3	n.s.
G%	25.6 ± 2.3	30.4 ± 0.5	18.7	**	55.6 ± 1.4	55.3 ± 1.7	-0.6	n.s
S'%	0.8 ± 0.3	0.8 ± 0.2	4.0	n.s.	n.d.	n.d.	-	-
S%	73.6 ± 2.2	<u>68.8 ± 0.6</u>	<u>-6.6</u>	**	44.4 ± 1.4	44.7 ± 1.7	0.7	n.s
T%	1.9 ± 0.3	n.d.	<u>inf down</u>	***	27.6 ± 1.4	<u>n.d.</u>	<u>inf down</u>	***
Total (G+S+S')	100	100			100	100		
S/G	3.0 ± 0.5	<u>2.3 ± 0.1</u>	<u>-20.7</u>	**	0.8 ± 0.1	0.8 ± 0.1	1.3	n.s.
X2 (cinnamaldehyde end-group)%	0.9 ± 0.4	1.0 ± 0.3	13.5	n.s.	n.d.	0.4 ± 0.3	inf up	**
X3 (benzaldehyde end- group)%	- ± -	0.1 ± 0.1	66.6	n.s.	n.d.	n.d.	-	-
B (β -5)%	0.6 ± 0.3	0.9 ± 0.6	57.9	n.s.	2.0 ± 0.5	4.2 ± 0.9	109.6	***
A (β -O-4)%	94.4 ± 0.9	<u>92.6 ± 1.2</u>	<u>-2.0</u>	*	97.7 ± 0.7	<u>93.9 ± 1.5</u>	<u>-3.9</u>	***
C (β - β , resinol)%	0.3 ± 0.3	0.6 ± 0.6	67.6	n.s.	0.3 ± 0.3	1.2 ± 0.5	310.3	**
C' (β - β , tetrahydrofuran)%	4.7 ± 0.7	6.0 ± 0.6	27.9	*	n.d.	0.7 ± 0.5	inf up	***
Total (A+B+C+C')	100	100			100	100		

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39 **Supplemental Table S4. Saccharification efficiency of senesced stem and leaf biomass from *C2-Idf* mutant and *C2* control plants.** The numbers listed in this
40 table are visualized in the graphs of Figure 4.

41 See separate file, Supplemental Table S4.xlsx

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