RESEARCH PAPER

Cell-wall properties contributing to improved deconstruction by alkaline pre-treatment and enzymatic hydrolysis in diverse maize (Zea mays L.) lines

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Received 12 September 2014; Revised 23 December 2014; Accepted 2 January 2015

Abstract

A maize (Zea mays L. subsp. mays) diversity panel consisting of 26 maize lines exhibiting a wide range of cell-wall properties and responses to hydrolysis by cellulolytic enzymes was employed to investigate the relationship between cell-wall properties, cell-wall responses to mild NaOH pre-treatment, and enzymatic hydrolysis yields. Enzymatic hydrolysis of the cellulose in the untreated maize was found to be positively correlated with the water retention value, which is a measure of cell-wall susceptibility to swelling. It was also positively correlated with the lignin syringyl/guaiacyl ratio and negatively correlated with the initial cell-wall lignin, xylan, acetate, and p-coumaric acid (pCA) content, as well as pCA released from the cell wall by pre-treatment. The hydrolysis yield following pre-treatment exhibited statistically significant negative correlations to the lignin content after pre-treatment and positive correlations to the solubilized ferulic acid and pCA. Several unanticipated results were observed, including a positive correlation between initial lignin and acetate content, lack of correlation between acetate content and initial xylan content, and negative correlation between each of these three variables to the hydrolysis yields for untreated maize. Another surprising result was that pCA release was negatively correlated with hydrolysis yields for untreated maize and, along with ferulic acid release, was positively correlated with the pre-treated maize hydrolysis yields. This indicates that these properties that may negatively contribute to the recalcitrance in untreated cell walls may positively contribute to their deconstruction by alkaline pre-treatment.

Key words: Biofuels, cell-wall recalcitrance, enzymatic hydrolysis, maize, plant cell-wall characterization, pre-treatment.

Introduction

Structural polymers within plant cell walls (i.e. cellulose, hemicelluloses, and lignins) offer potential for long-term sustainable production of renewable fuels, chemicals, polymers, and materials that are currently produced from petrochemicals. Many of the promising conversion pathways for these products are based on cascades of biochemical and/or catalytic reactions starting with the sugars derived from cellulose (glucose) and hemicelluloses (primarily xylose in angiosperms).
The recalcitrance of plant cell walls to biological degradation, deconstruction, or conversion is considered to be the most crucial factor to overcome in order to develop successful bioprocessing technologies for lignocellulose conversion to renewable fuels and chemicals (Himmel et al., 2007). As such, in order to generate high sugar yields from the cellulose and hemicellulose within plant cell walls, pre-treatment is required in combination with the subsequent polysaccharide hydrolysis by either enzymes or an acid catalyst (Wyman et al., 2011).

The grasses or graminaceous monocots represent many important agricultural residues such as corn (maize) stover (Zea mays L. subsp. mays), wheat straw (Triticum spp.), rice straw (Oryza sativa L.), and sugarcane bagasse (Saccharum spp.), as well as many promising perennial bioenergy feedstocks such as switchgrass (Panicum virgatum L.) and Miscanthus spp. (Vermerris, 2011). Outcomes of pre-treatment are obviously dependent on the combination of pretreatment chemistry and the plant cell-wall properties. The cell walls of graminaceous monocots are known to contain alkali-labile ferulate ester cross-links within the hemicellulose and lignin (Scalbert et al., 1985; Hatfield et al., 1999; Iiyama and Lam, 2001), as well as high phenolic hydroxyl contents in their lignins, resulting in increased alkali solubility (Lapierre et al., 1989), with these properties rendering the cell wall highly susceptible to delignification by alkaline pre-treatments (Li et al., 2012; Liu et al., 2014). As a consequence, mild alkali pre-treatment of grasses such as maize has shown substantial promise, as these can be employed for both fractionating biomass and generating a pre-treated biomass that is highly amenable to enzymatic hydrolysis (Chen et al., 2013; Karp et al., 2014; Liu et al., 2014).

Corn stover (i.e. the stems and leaves) is considered the most promising agricultural residue in the USA for extensive deployment of technologies for biomass conversion to biofuels and biochemicals (US Department of Energy, 2011). A substantial body of literature exists on the correlation of cell-wall properties in maize with in vitro digestibility by rumen microbiota (Jung and Buxtono, 1994; Lundvall et al., 1994; Argillier et al., 2000; Jung et al., 2000; Méchin et al., 2000; Fontaine et al., 2003; Lorenz et al., 2010; Zhang et al., 2011) and in vivo ruminant digestibility (Jung et al., 1997), as well as hydrolysis by cellulolytic enzymes (Oakley et al., 2013), with trends typically identifying strong negative correlations between the lignin content and compositions with cell-wall biological degradability. Other important initial cell-wall properties that have been correlated with digestibility or hydrolysis include the lignin syringyl/guaiacyl (S/G) ratio, lignin β-O-4 content, etherified ferulic acid (FA) content, and esterified p-coumaric acid (pCA) content (Jung and Buxtono, 1994; Méchin et al., 2000; Iiyama and Lam, 2001). Strategies for engineering maize lines for its cellulosic biofuel or silage/feed value have been reviewed recently (Barrière et al., 2003; Jung et al., 2012) with strategies that include increasing the non-starch biomass yield and the total carbohydrate content (Vermerris et al., 2007; de Leon et al., 2013) and redirection of carbon to non-cellulosic sugars in the cell wall (e.g. β-glucan, starch; Slewinski, 2012; Chuck et al., 2011; Chuck, 2013; Pauly et al., 2014), as well as reducing the overall recalcitrance through decreasing the lignin content (He et al., 2003), altering the lignin monomer content (Piquemal et al., 2002), and alteration of feruloylation (Jung and Phillips, 2010; Barros-Rios et al., 2012).

Cell-wall structural differences are typically neglected in cell-wall property correlation studies, although it is known that substantial differences in digestibility or sugar hydrolysis yields are found between cell types in untreated (Wilson et al., 1993; Wilson and Hatfield, 1997; Hansey et al., 2010) or pre-treated (Zeng et al., 2012) grasses. As an example, Sarath et al. (2011) found that while cell-wall lignin content in diverse switchgrass lines was negatively correlated with hydrolysis yields following dilute acid pre-treatment, more than 50% of the variability could be attributed to differences in cell-wall architecture. The water retention value (WRV) has been used in our recent work as a predictor of enzymatic hydrolysis yields for corn stover and switchgrass subjected to alkali, alkaline oxidative, and liquid hot water pre-treatments (Williams and Hodge, 2014), and is proposed to act as an indirect measurement of a number of cell-wall properties (e.g. overall hydrophilicity, porosity, and polysaccharide accessibility) that are manifested in the cell wall’s ability to swell in water. This measurement may be able to incorporate structural differences between different cell-wall types such as the highly hygroscopic and digestible pith parenchyma cell walls versus the more hydrophobic and less-digestible cell walls within vascular bundles and the epidermis.

There is substantially less published literature on the correlation between plant cell-wall properties and their response to pre-treatment and hydrolysis. Recently, approaches for high-throughput screening of pre-treatment and enzymatic hydrolysis have been developed for the purpose of screening large sets of cell-wall material for the purpose of identifying potential reduced recalcitrance phenotypes both with and without a pre-treatment (Decker et al., 2009; Santoro et al., 2010). Understanding the cell-wall traits associated with reduced recalcitrance phenotypes or phenotypes that exhibit specific responses to pre-treatment are an important component of screening. However, typically only composition analysis is performed across the sample sets, limiting the understanding of the cell-wall properties, traits, or phenotypes that are also associated with the cell wall’s response to enzymatic hydrolysis. Notably, recent work has employed high-throughput screening of panels of promising bioenergy grasses that include diverse cultivars of wheat (Triticum aestivum L.) (Lindedam et al., 2012), Miscanthus spp. (Zhang et al., 2012), and maize (Penning et al., 2014) subjected to hydrothermal pre-treatment, and a sorghum (Sorghum bicolor (L.) Moench) diversity panel subjected to aqueous ammonia pre-treatment (Vandenbrink et al., 2010). Importantly, from these cited studies, while they were able to identify substantial differences in cell-wall responses to pre-treatment, it was difficult to draw conclusions about the cell-wall properties, traits, or phenotypes that are also associated with improved hydrolysis yields or sugar release following pre-treatment. Relative to other high-throughput screening studies of poplar, where lignin content as well as the S/G ratio could be clearly be
identified as a strong contributor to hydrolysis yields following pre-treatment (Studer et al., 2011), lignin was not correlated with hydrolysis yields in these studies, indicating that other unquantified cell-wall properties may be responsible for the observed variability in the data sets.

While at least one study has investigated the alteration of cell-wall properties and in vitro digestibility in a diverse set of orchard grass (Dactylis glomerata L.) subjected to acid chloride delignification (Darcy and Belyea, 1980), there is scant literature on the relationship between plant cell-wall properties and hydrolysis yields for diverse maize lines subjected to delignifying pre-treatments such as mild NaOH pre-treatment. Thus, the goal of this work was to understand better how maize cell-wall properties impact initial recalcitrance as well as NaOH pre-treatment. Specifically, for this work 12 cell-wall properties were selected that may influence overall cell-wall recalcitrance including WRV, xylan and lignin contents before and after pre-treatment, initial cell-wall acetate content, pCA and FA content before pre-treatment, solubilized pCA and FA during pre-treatment, and the S/G ratio of the lignin in the untreated cell-wall. These properties were quantified for a diversity panel of 26 maize lines that were previously identified as exhibiting substantial phenotypic diversity in glucose release following mild NaOH pre-treatment (Muttoni et al., 2012). These were subjected to mild NaOH pre-treatment and the glucose yields determined for the different lines before and after the pre-treatment. Subsequently, correlations between the cell-wall properties and hydrolysis yields were investigated.

Materials and methods

Maize diversity panel

The maize diversity set was grown in Arlington, WI, USA, in 2012 and the complete list of maize lines with accession number or resources is presented in Supplementary Table S1 at JXB online. The maize was harvested at grain physiological maturity with a Case IH® 2144 axial-flow combine, which allows harvesting grain and biomass of samples including untreated and alkaline oxidative pre-treated samples, 2 g of biomass was freshly pre-treated with NaOH, filtered using a fabricated 200-mesh Buchner funnel, and rinsed with 500 ml of deionized water. Subsequently, a plug of filtered, wet biomass was analysed as described in our previous work. Measurements were taken in triplicate.

NaOH pre-treatment, composition analysis, and enzymatic hydrolysis

Biomass samples were subjected to a mild alkaline pre-treatment whereby 2 g of biomass was added to 20 ml of an aqueous 8 g l⁻¹ NaOH solution in a 50 ml centrifuge tube corresponding to an alkali loading of 0.08 g NaOH g⁻¹ of biomass. These tubes were placed in a static water bath at 80 °C for 1 h. Following pre-treatment, the liquid was removed by filtration and the residual biomass was washed with deionized water until neutral. The pre-treatment mass yield was determined by quantifying the difference between the original and air-dried pre-treated materials gravimetrically. Composition analysis was carried out according to the NREL/TP 510–42618 to determine the structural carbohydrates, acetate, and Klasson lignin of the untreated and pre-treated maize lines using a high-performance liquid chromatography (HPLC) machine equipped with an Aminex HPX-87 H (Bio-Rad, Hercules, CA, USA) column. The enzymatic hydrolysis was performed at pH 5.0, 50 °C, and 180 rpm shaking, with a 30 mg g⁻¹ of protein loading of Cellic CTec2 (Novozymes A/S, Bagsværd, Denmark) on glucan (37.7 filter paper units g⁻¹ glucan) for 6 or 72 h. The yields were determined as the amount of glucose (as glucan) released following enzymatic hydrolysis divided by the glucan content of the untreated or pre-treated samples as determined by composition analysis. All data for cell-wall composition, pre-treatment mass yields, and hydrolysis yields are available in Supplementary Table S3 at JXB online. The composition analysis and enzymatic hydrolysis were performed in duplicate, with the data representing the mean and error bars representing the data range.

Quantification of p-hydroxycinnamic acids

The p-hydroxycinnamic acids were determined by first treating 0.50 g of biomass with 25 ml of 3 M NaOH in sealed pressure tubes at 120 °C for 1 h in an autoclave to release both esterified pCA and FA and etherified FA. After cooling to room temperature, 250 μl of 10 mg ml⁻¹ o-coumaric acid in methanol was added as an internal standard for each sample. The mixture was transferred to 1.5 ml centrifuge tubes and centrifuged at 13 000 rpm for 10 min. The pH of the supernatant was adjusted to 2.0 using concentrated HCl and the samples were then stored overnight at 4 °C. These samples were subsequently analysed by HPLC (Agilent 1100 Series) equipped with a C18 column (Discovery, 5 μm particle size, 5 cm length by 2.1 mm internal diameter; Sigma-Aldrich). Standards containing FA, pCA, and o-coumaric acid were also analysed and, together with the internal standard, were used to determine the concentration of the p-hydroxycinnamic acids in the samples. Concentrations were converted to a mass per original mass sample basis.

Quantification of WRV

The WRV of untreated samples was determined as described in our previous work (Williams and Hodge, 2014). However, to avoid problems associated with sample drying for NaOH-pre-treated samples, 2 g of biomass was freshly pre-treated with NaOH, filtered using a fabricated 200-mesh Buchner funnel, and rinsed with 500 ml of deionized water. Subsequently, a plug of filtered, wet biomass was analysed as described in our previous work. Measurements were taken in triplicate.

Determination of the S/G ratio

The S/G ratio was predicted by the combination of principal component analysis and partial least squares regression based on their pyrolysis molecular beam mass spectrometry (py-MBMS) spectra provided by Robert Sykes (National Renewable Energy Laboratory). The parameters of the prediction model were generated previously by correlating thioacidolysis S/G and py-MBMS spectra of a set of samples including untreated and alkaline oxidative pre-treated hybrid corn stover, brown midrib stover mutants bml1 and bml3, switchgrass, and Miscanthus.

Data analysis

Hierarchical cluster analysis according to Euclidian distance was performed in MATLAB (Mathworks, Natick, MA, USA) on the complete matrix of the Pearson’s correlation coefficients for all properties and yield combinations. Correlations were identified as significant if the P value associated with the proportionality coefficient was less than or equal to 0.05. Missing data points for some of the properties and yields were handled by only determining correlations and statistical parameters between sets of samples containing properties and yields that were being correlated.

Results and discussion

Cell-wall properties and hydrolysis yields

Twelve cell-wall properties or traits that may be indicators of cell-wall recalcitrance were quantified across the maize
Within the correlation map (Fig. 2), several multi-property clusters of positive correlations were observed along the top left to bottom right diagonal, while one multi-property cluster of negative correlations stood out in both the bottom left and top right corners. This indicated that there a number of properties that are correlated across diverse maize lines and may be responsible for differences in the cell wall’s response to enzymatic hydrolysis as well as the response to pre-treatment. A number of strong positive correlations between related properties or yields were observable along the top left to bottom right diagonal, namely the initial 6 and final 72h hydrolysis yields, initial pCA content and pCA solubilization, initial FA content and FA solubilization, and initial and final WRV. These specific results are not surprising as they may be responsible for differences in the cell wall’s response to enzymatic hydrolysis as well as the response to pre-treatment.

**Table 1.** Variability within the data set for the 12 properties across the 27 maize lines

|    | Initial WRV | Final WRV | Initial xylan | Final xylan | Initial lignin | Final lignin | Initial pCA | Final pCA | Initial acetate | Final acetate | Initial FA | Final FA | S/G ratio | pCA release | FA release |
|----|-------------|-----------|---------------|-------------|----------------|--------------|-------------|-----------|----------------|--------------|-----------|----------|-----------|-------------|------------|
| X1 | 1.7         | 2.2       | 0.16          | 0.20        | 0.13           | 0.20         | 0.27        | 0.20      | 0.20           | 0.20         | 0.30      | 0.02     | 0.02      | 0.02       | 0.02       |
| X2 | 2.0         | 2.7       | 0.20          | 0.27        | 0.17           | 0.21        | 0.21        | 0.21      | 0.21           | 0.21         | 0.21      | 0.02     | 0.02      | 0.02       | 0.02       |
| X3 | 2.4         | 3.3       | 0.26          | 0.30        | 0.20           | 0.26        | 0.26        | 0.26      | 0.26           | 0.26         | 0.26      | 0.02     | 0.02      | 0.02       | 0.02       |
| Min| 0.17        | 0.30      | 0.16          | 0.20        | 0.068          | 0.068       | 0.14        | 0.14      | 0.02           | 0.02         | 0.02      | 0.02     | 0.02      | 0.02       | 0.02       |
| Mean| 1.7         | 2.7       | 0.16          | 0.20        | 0.20           | 0.20        | 0.27        | 0.27      | 0.20           | 0.20         | 0.30      | 0.02     | 0.02      | 0.02       | 0.02       |
| Max| 2.4         | 3.3       | 0.26          | 0.30        | 0.26           | 0.26        | 0.30        | 0.30      | 0.26           | 0.26         | 0.30      | 0.02     | 0.02      | 0.02       | 0.02       |

**Correlations between cell-wall properties**

In order to visualize better the relationships between the variables, a correlation map was developed and organized using hierarchical clustering according to the Euclidian distance between sets of the Pearson correlation coefficients ($R$) (Fig. 2). The magnitude, scale, and significance of the correlations are presented in Supplementary Table S2 at JXB online. Within the correlation map (Fig. 2), several multi-property clusters of positive correlations were observed along the top left to bottom right diagonal, while one multi-property cluster of negative correlations stood out in both the bottom left and top right corners. This indicated that there a number of properties that are correlated across diverse maize lines and may be responsible for differences in the cell wall’s response to enzymatic hydrolysis as well as the response to pre-treatment.

Glucose hydrolysis yields ranged between 37.8 and 55.5% for untreated biomass and between 58.6 and 82.5% for NaOH-pre-treated biomass after 72h of hydrolysis (Fig. 1), clearly exhibiting a diverse range of digestibility phenotypes as well as diverse responses to mild NaOH pre-treatment. The 6h hydrolysis yields are intended to represent the initial hydrolysis rates, while the 72h hydrolysis yields represent the extent of hydrolysis as only minimal additional sugar release is observed beyond 72h (data not shown). It should be noted that these data were plotted for glucose yield rather than glucose release, so that results were not biased towards cell walls with higher glucan content.

**Additional Notes:**

- WRV is proposed to act as a proxy variable for initial FA content and FA solubilization, and initial and final hydrolysis yields, initial and final pCA release, initial acetyl content, pCA content, and FA content, as well as corresponding pCA and FA release. The S/G ratio as determined by py-MBMS coupled to partial least squares models also showed a diverse range, although within the range reported for maize (Morrison et al., 1998).

- Wide ranges were observed for initial cell-wall acetyl content, pCA content, and FA content, as well as corresponding pCA and FA release. The S/G ratio as determined by py-MBMS coupled to partial least squares models also showed a diverse range, although within the range reported for maize (Morrison et al., 1998).

- Initial WRV is proposed to act as a proxy variable for initial FA content and FA solubilization, and initial and final hydrolysis yields, initial and final pCA release, initial acetyl content, pCA content, and FA content, as well as corresponding pCA and FA release. The S/G ratio as determined by py-MBMS coupled to partial least squares models also showed a diverse range, although within the range reported for maize (Morrison et al., 1998).
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The acetate content showed a diverse range (26.7–39.2 mg g\(^{-1}\)) there was no correlation with the cell-wall xylan content (Fig. 3A), indicating either that the xylans exhibited either a wide range of \(O\)-acetylation—calculated to be 0.25–0.60 moles of acetyl per mole of xylopyranosyl, which is within the reported range for maize (Selig et al., 2009)—or that the acetyl groups are substituted on moieties other than xylan (e.g. glucomannans, pectins, or lignins). While uncorrelated to the initial xylan content (Fig. 3A), this diverse range of acetate contents could be observed to exhibit correlations with a number of other cell-wall properties, including strong, statistically significant positive correlations with the final xylan content (Fig. 3A), and the initial content and release of \(p\)CA (Fig. 3C), while showing negative correlations to the initial lignin content (Fig. 3B) and a strong positive correlation to both the final lignin content (Fig. 3B) and the initial WRV (Fig. 3D).

One of the primary outcomes of alkaline pre-treatment of grasses is the saponification of ester bonds in the plant cell-wall biopolymers resulting in solubilization of acetate, FA, \(p\)CA, xylans, and lignin fragments. The acetate results can be interpreted as potentially relating to the lignin release. While it may be expected that high-acetate contents would consume more alkali and presumably hinder lignin removal, the opposite is the case. As observed in Fig. 3B, the initial cell-wall acetate content was positively correlated to initial lignin contents and negatively correlated to final lignin content, indicating that high-acetate cell walls are likely to be more recalcitrant yet respond better to alkaline pre-treatment with respect to lignin removal. Additionally, it was observed that initial \(p\)CA content and \(p\)CA released were positively correlated with initial acetate content (Fig. 3C). This may be a consequence of the correlation of between acetate and lignin since \(p\)CA is known to be acylated to syringyl lignins and may only represent the correlation to lignin content (Jung and Allen, 1995). The correlation to final xylan content may be a consequence of high acetate corresponding to higher lignin removal, which

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**Fig. 1.** Range of hydrolysis yields obtained for untreated and NaOH-pre-treated maize for 6 h (A) and 72 h (B) hydrolysis yields. Error bars represent data range for duplicate samples. Due to missing data, some samples do not appear. (This figure is available in colour at JXB online.)

**Fig. 2.** Correlation map of the Pearson’s correlation coefficients for the 12 cell-wall properties and four hydrolysis yields (red text) across the 27 maize lines as organized by hierarchical cluster analysis. Clusters of properties and yields exhibiting strong correlations are highlighted. ‘Initial’ indicates the property in the original untreated biomass sample, while ‘final’ indicates the property following pre-treatment.
would enrich the xylan content of the pre-treated biomass. Overall, it is not clear whether this cluster of co-varying properties is an indication of differences in the ‘average’ cell-wall property or whether these may indicate differences in the abundance of, for example, high-lignin, high-\(p\)-\(p\)-coumaroylated (\(p\)-\(p\)-CA), and high-acetate tissues. Recently, work has been targeted at altering the expression of genes associated with \(O\)-\(p\)-acetylation as a strategy for altering cell-wall recalcitrance (Xiong et al., 2013), although it is currently not clear how only alteration in xylan \(O\)-\(p\)-acetylation will impact cell-wall recalcitrance.

Interestingly, WRVs did not show a significant correlation with any of the cell-wall biopolymer content (i.e. lignin and xylan) while it did exhibit significant correlations (albeit weak) with substitutions on these biopolymers (i.e. \(p\)-\(p\)-CA and acetate; Fig. 3D). Specifically, the initial cell-wall \(p\)-\(p\)-CA content was inversely correlated to the initial WRV, which may provide evidence that highly \(p\)-coumaroylated cell walls may be more hydrophobic (and higher in lignin content), potentially imparting increased recalcitrance to degradation by microbial pathogens or rumen microbiota. The (weak) negative correlation between initial cell-wall acetate content and initial WRV (Fig. 3D) could potentially be due to increasing hydrophobicity of exposed xylans (i.e. the glycan will have an exposed ethyl group rather than a hydroxyl group). However, the hypothesis that highly acetylated cell walls are more porous due to a decreased level of H-bonding between xylans and cellulose does not fit this data since increasing acetyl content corresponded to decreasing WRVs (Fig. 3D). As initial lignin content was positivity correlated with the cell-wall acetate content (Fig. 3B), the decreasing initial WRV with increasing acetate could be a consequence of the increasing lignin content decreasing the cell-wall’s capacity to absorb water.

**Correlations between cell-wall properties and hydrolysis yields**

Besides between-property correlations, correlations between cell-wall properties and hydrolysis yields are important for understanding property contributions to cell-wall recalcitrance. All significant \((P \leq 0.05)\) correlations between cell-wall properties and hydrolysis yields are plotted in Fig. 4. Notably, this plots only shows correlations for untreated 6 h yields and...
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NaOH-pre-treated 72h yields. The untreated 72h yield did not exhibit significant correlations with any properties other than the untreated 6h yields (Fig. 2 and Table 1). However, the correlations that were strongest for the untreated 6h hydrolysis yields were not as strong for 72h digestibility, exhibiting similar correlations with P values between 0.05 and 0.15. The differences between the hydrolysis yields obtained at different time points (6 vs 72h) may indicate that the initial cell-wall composition impacts the hydrolysis rate more strongly than the hydrolysis extent. The 6h hydrolysis yields for NaOH-pre-treated corn stover were not found to exhibit strong correlations with any other properties and, furthermore, were often lower than the untreated 6h hydrolysis yields (Fig. 1B). This contradictory result may be due to the drying of the pre-treated material necessitated by the analysis that resulted in its stronger resistance to rehydration, which may have introduced more variability in the data for the initial glucose release by hydrolysis but presumably not the extent of hydrolysis.

A number of important trends can be identified from the data in Fig. 4. The first is the identification that WRV was positively correlated with the untreated 6h hydrolysis yield (Fig. 4A). As discussed in the introduction, WRV was included as a parameter that may offer the potential for consolidating
a number of other, unquantified cell-wall properties that may be able to be correlated with hydrolysis yields. Our previous work used only two types of biomass (corn stover and switchgrass) subjected to varying levels of delignification by alkaline oxidative pre-treatment and found strong correlations between WRV and hydrolysis yields (Williams and Hodge, 2014). The current work used a wide range of (similar) biomass that was subjected to a single pre-treatment condition, which may explain why WRVs following pre-treatment were not able to predict differences in hydrolysis yields following pre-treatment.

The initial cell-wall acetate content demonstrated a negative correlation with the untreated 6 h hydrolysis yield although not with the pre-treated 72 h yield (Fig. 4B). As discussed previously, the acetate content is strongly correlated to many other properties and potentially this cluster of related properties may be able to explain differences in hydrolysis yields rather than a single property. For example, initial acetate content was strongly correlated with initial lignin content (Fig. 3B), which may be the property most strongly responsible for the response in hydrolysis yields. Previously, non-linear models have been developed relating decreases in acetate and lignin to increases in hydrolysis yields for lime-pre-treated corn stover (Kim and Holtzapple, 2006) and hybrid poplar (O’Dwyer et al., 2008), although these correlated simultaneous acetate and lignin removal by alkali with improved hydrolysis yields rather than initial variability in these properties, as was done in the present work.

Lignin is thought to impact saccharification yields by physical occlusion of polysaccharides (Brunecky et al., 2009), providing resistance to swelling (O’Neil and York, 2003), and by non-specific binding to cellulases (Palonen et al., 2004). Studies have generally shown a strong negative correlation between lignin content and hydrolysis yields for a wide range of untreated grasses (Jung and Vogel, 1986; Jung et al., 1997; Zhang et al., 2011) with strong correlations following a delignifying pre-treatment (Sewalt et al., 1997; Kim and Holtzapple, 2006; Li et al., 2012). As expected, the untreated lignin content was negatively correlated with the untreated 6 h hydrolysis yield (Fig. 4C) and the lignin content following pre-treatment was negatively correlated with the pre-treated 72 h hydrolysis yield (Fig. 4D). However, the lignin contents prior to and following pre-treatment did not correlate with each other (Fig. 2 and Table 1), indicating that lignin removal for mild NaOH pre-treatment is not necessarily dependent on the initial lignin content but is dependent on many other cell-wall properties. This is significant in that many strategies for reduced cell-wall recalcitrance have targeted lignin levels (Simmons et al., 2010), although it may be sufficient that the lignin is easily removable by the pre-treatment to decrease the final cell-wall recalcitrance rather than the initial cell-wall recalcitrance.

As we have already identified in this work, the lignin removal is more important than the initial lignin content for mild NaOH pre-treatment and consequently, lignin properties that may contribute to improved lignin removal are important. Another interesting lignin-related finding was that the initial S/G ratio was positively correlated with the untreated 6 h hydrolysis yield (Fig. 4E), while the trend apparently reversed for the pre-treated hydrolysis yield (although this was not a statistically significant correlation). The correlation between S/G ratio in untreated biomass and in vitro ruminant digestibility for grasses has been somewhat contradictory in the literature (Méchin et al., 2000, 2005; Zhang et al., 2011) although a general trend is that an increasing S/G ratio may be linked to increasing digestibility.

The initial xylan content was found to be negatively correlated with the untreated 6 h hydrolysis yield (Fig 4F). The fact that both initial lignin and xylan were negatively correlated with the untreated hydrolysis yields yet these two properties were uncorrelated with each (Fig. 2) may indicate that both of these components play an important role in the limited access of cellulolytic enzymes to cellulose. The enzyme cocktail utilized for this study was not supplemented with xylanase, which may be a factor contributing to this result.

Ferulate and diferulate esters are known to be attached to the primary hydroxyl at the C5 position of α-L-arabinofuranosyl residues in xylans and be involved in cross-linking of xylans to lignin by incorporation of the aromatic moiety through ether linkages into growing lignin polymers or by cross-coupling (Hatfield et al., 1999). This cross-linking of cell-wall polymers by ferulates is generally accepted to play an important role in cell-wall recalcitrance (Grabber et al., 1998). Negative correlations have been found in the literature for total cell-wall ferulate content and in vitro ruminant digestibility in maize (Jung et al., 2011), as well as in other diverse grasses (Iiyama and Lam, 2001), and decreasing cell-wall ferulate content has been investigated as a strategy for improving the forage quality of maize (Jung and Phillips, 2010). However, while strong, statistically significant negative correlations were identified between etherified ferulates and in vitro digestibility, esterified ferulate content was found to be positively correlated with in vitro digestibility in smooth bromegrass (Bromus inermis Leyss subsp. inermis), cocksfoot (D. glomerata L.), and reed canary grass (Phalaris arundinacea L.). (Casler and Jung, 2006; Casler et al., 2008), and other recent work has validated a strategy for increasing esterified diferulate content in maize to improve its digestibility (Barros-Rios et al., 2012). The present work did not distinguish between etherified and esterified ferulates, although the method for ferulate content was performed under harsh enough conditions that this term probably comprises both etherified and esterified as increasing the alkali loading and temperature during saponification did not result in additional ferulate release (data not shown). The ferulate release during pre-treatment may be more representative of esterified ferulate due to the mild conditions utilized for pre-treatment. Nonetheless, ferulate content and the ferulate released were found to be strongly positively correlated to each other (Fig. 2 and Table 1) and, importantly, the ferulate released showed a strong correlation with the 72 h hydrolysis yields following NaOH pre-treatment (Fig. 4G), although not with any of the other hydrolysis conditions. This can be understood, as cell walls that are highly cross-linked are more susceptible to alkaline pre-treatment. Recently, incorporation of ferulate esters into hybrid poplar has been validated as a
strategy to improve enzymatic hydrolysis following an alkaline pre-treatment (Wilkerson et al., 2014).

Lignin content, pCA content, and the S/G ratio in maize stem internodes and rinds have been shown to increase with increasing maturity, while ferulate content does not change (Morrison et al., 1998). The simultaneous increase in these properties has been implicated in the decrease in maize digestibility with increasing maturity (Grabber et al., 2004; Jung and Casler, 2006). Both the initial pCA (Fig. 4H) and solubilized pCA (Fig. 4I) exhibited statistically significant correlations with the hydrolysis yields. These results were notable in that the correlation was the opposite for untreated biomass compared with NaOH-pre-treated biomass, whereby increasing release or content of pCA corresponded to decreasing hydrolysis yields for untreated cell walls and increasing hydrolysis yields for pre-treated cell walls. High pCA content in untreated maize has been correlated with low in vitro digestibility (Méchin et al., 2000; Zhang et al., 2011), although the finding that high initial p-coumarylation of lignin may be related to high hydrolysis yields following pre-treatment is novel. As discussed previously, the pCA content was positively correlated with the initial lignin content, so this could be an indirect measure of the impact of lignin content. As another alternative, pCA-containing lignins may be responsible for the uptake and inactivation of cellulolytic enzymes, as is known to occur for polyphenolic compounds such as tannins and lignins and phenolic acid monomers (Ximenes et al., 2011).

Conclusions

Understanding the relationship between cell-wall recalcitrance and pre-treatment is important in that this may lead to the identification of strategies for plant breeding or genetic engineering that may improve cell-wall deconstruction. For this study, 26 diverse maize lines were subjected to mild NaOH pre-treatment and a broad set of properties relating to cell-wall recalcitrance were characterized. By combining cell-wall structural characterization and enzymatic hydrolysis of cell-wall polysaccharides both prior to and following mild NaOH pre-treatment, a number of expected as well as non-intuitive results were identified. The hydrolysis yields of non-pre-treated cell walls were found to be positively correlated with WRV and S/G ratio and negatively correlated with xylan and acetate content. While the initial cell-wall xylan and acetate content were uncorrelated with each other, the acetate content was found to exhibit a number of strong correlations with other cell-wall properties. The pre-treated cell-wall hydrolysis yields were positively correlated with the ferulate released by pre-treatment, indicating that breaking of ferulate cross-links between cell-wall polymers is an important outcome of pre-treatment. As expected, statistically significant negative correlations were identified between the cell-wall lignin content and the hydrolysis yields for both untreated (R=–0.40, P=0.037) and NaOH-pre-treated maize (R=–0.50, P=0.009). It has long been known that cell-wall lignin content can be negatively correlated with enzymatic hydrolysis yields and in vitro digestibility, as well as in vivo digestibility in ruminants. However, the data demonstrated that the initial cell-wall lignin content and the pre-treated cell-wall content were not correlated with each other, that initial cell-wall lignin content was not correlated with hydrolysis yields following pre-treatment, and the cell-wall lignin content following pre-treatment was not correlated with untreated hydrolysis yields. This important finding indicates that, while enzymatic hydrolysis yields may be set by the cell-wall lignin content, the cell wall’s response to delignifying pre-treatment such as mild NaOH pre-treatment is not necessarily set by the initial lignin content. The pCA that was saponifiable by mild NaOH pre-treatment showed a negative correlation with the hydrolysis yields of untreated maize (R=–0.61, P=0.001) and the inverse response for pre-treated maize (R=0.40, P=0.044), indicating that cell walls with a high content of saponifiable pCA are more recalcitrant without treatment, yet respond better to pre-treatment than cell walls with a low content of saponifiable pCA. Importantly, this indicates that properties contributing to a ‘reduced recalcitrance’ phenotype following a specific pre-treatment are not necessarily the same properties that contribute to recalcitrance in untreated cell walls.

Supplementary data

Supplementary data are available at JXB online.

Supplementary Table S1. Genotype and reference source for the 27 maize lines used in this work.

Supplementary Table S2. Calculated proportionality constants (unscaled) between all cell-wall properties and hydrolysis yields.

Supplementary Table S3. Complete data set for cell-wall composition prior to and following pre-treatment, mass yields following pre-treatments, quantified properties, and hydrolysis yields.

Acknowledgements

The authors would like to acknowledge Robert Sykes (NREL) for generously performing py-MBMS analysis on the samples. This work was supported by the US Department of Energy Great Lakes Bioenergy Research Center (grant no. DOE BER Office of Science DE-FC02-07ER64494). ML and JDC were supported in part by a grant from the US National Science Foundation (grant no. NSF CBET 1336622). David Hodge is grateful for support by a travel award from the US Department of Energy, Office of Science, Office of Basic Energy Sciences (BES) and the Office of Biological and Environmental Research (BER), DE-FOA-0000995.

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