Lawrence Berkeley National Laboratory
Recent Work

Title
Survey of renewable chemicals produced from lignocellulosic biomass during ionic liquid pretreatment.

Permalink
https://escholarship.org/uc/item/71d9d3rw

Journal
Biotechnology for biofuels, 6(1)

ISSN
1754-6834

Authors
Varanasi, Patanjali
Singh, Priyanka
Auer, Manfred
et al.

Publication Date
2013-01-28

DOI
10.1186/1754-6834-6-14

Peer reviewed
Survey of renewable chemicals produced from lignocellulosic biomass during ionic liquid pretreatment

Patanjali Varanasi1,2, Priyanka Singh1, Manfred Auer1, Paul D Adams1, Blake A Simmons1,2 and Seema Singh1,2*

Abstract
Background: Lignin is often overlooked in the valorization of lignocellulosic biomass, but lignin-based materials and chemicals represent potential value-added products for biorefineries that could significantly improve the economics of a biorefinery. Fluctuating crude oil prices and changing fuel specifications are some of the driving factors to develop new technologies that could be used to convert polymeric lignin into low molecular weight lignin and or monomeric aromatic feedstocks to assist in the displacement of the current products associated with the conversion of a whole barrel of oil. We present an approach to produce these chemicals based on the selective breakdown of lignin during ionic liquid pretreatment.

Results: The lignin breakdown products generated are found to be dependent on the starting biomass, and significant levels were generated on dissolution at 160°C for 6 hrs. Guaiacol was produced on dissolution of biomass and technical lignins. Vanillin was produced on dissolution of kraft lignin and eucalyptus. Syringol and allyl guaiacol were the major products observed on dissolution of switchgrass and pine, respectively, whereas syringol and allyl syringol were obtained by dissolution of eucalyptus. Furthermore, it was observed that different lignin-derived products could be generated by tuning the process conditions.

Conclusions: We have developed an ionic liquid based process that depolymerizes lignin and converts the low molecular weight lignin fractions into a variety of renewable chemicals from biomass. The generated chemicals (phenols, guaiacols, syringols, eugenol, catechols), their oxidized products (vanillin, vanillic acid, syringaldehyde) and their easily derivatized hydrocarbons (benzene, toluene, xylene, styrene, biphenyls and cyclohexane) already have relatively high market value as commodity and specialty chemicals, green building materials, nylons, and resins.

Keywords: Lignin valorization, Ionic liquid pretreatment, Renewable chemicals, Biofuels

Background
Lignocellulosic biomass is primarily composed of three biopolymers: cellulose, hemicelluloses and lignin [1]. Holocellulosic biopolymers are considered the most valuable components of lignocellulose and are utilized for the production of various products including paper and biofuels [2-4]. Lignin constitutes roughly a third of the biomass and is typically burned to produce waste heat and/or electricity within paper mills and biorefineries [1,5]. Lignin is a naturally occurring heterogeneous phenylpropanoid-based biopolymer, and provides mechanical support and water transport to the plant and inhibits the action of various biological agents (e.g., insects) on the plants [6]. It is estimated that 50 million tons of lignin is produced annually from pulp and paper industries worldwide [7]. The high energy content of lignin, the presence of highly reactive groups, and the fact that it will be generated in large quantities as second generation biorefineries are deployed represents a significant opportunity for the production of a wide range of renewable chemicals and materials that can be sold as co-products (Figure 1).

For example, lignin sulfonates produced from kraft pulping are currently utilized as phenol-formaldehyde plastics, binders, adhesives, mud-sand cements in drilling oil-wells, dispersants, or flotation agents, emulsifiers

* Correspondence: seesing@sandia.gov
1Joint Bioenergy Institute, Physical Biosciences Division, Lawrence Berkeley National Laboratory, Emeryville, CA, USA
2Sandia National Laboratories, Biological and Materials Science Center, Livermore, CA, USA

© 2013 Varanasi et al; licensee BioMed Central Ltd. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/2.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.
and stabilizers, grinding agents, electrolytic refining agents, protein precipitants, tanning agents, sequestering agents, storage battery plates, lime plaster, crystal growth inhibitor, ingot mold wash and as flame retardants [8,9]. Starch-based films incorporated with lignin have higher water resistance and increased elongation that make them effective packaging materials [10]. Lignin is also used in the production of conducting polymer lignosulfonic acid-doped polyamine [11]. Carbon fibers produced from lignin based materials require a lower amount of thermo-stabilization and possess high tensile strength [12]. Lignin has been used to produce various polymers like ARBOFORM, polyesters and polyurethanes and various polymer blends with PVC, polyolefins, and rubbers are being currently developed [13-15]. Lignin has been used to produce various polymers like ARBOFORM, polyesters and polyurethanes and various polymer blends with PVC, polyolefins, and rubbers are being currently developed [13-15]. Lignin has been used to produce various polymers like ARBOFORM, polyesters and polyurethanes and various polymer blends with PVC, polyolefins, and rubbers are being currently developed [13-15]. Lignin has been used to produce various polymers like ARBOFORM, polyesters and polyurethanes and various polymer blends with PVC, polyolefins, and rubbers are being currently developed [13-15]. Lignin has been used to produce various polymers like ARBOFORM, polyesters and polyurethanes and various polymer blends with PVC, polyolefins, and rubbers are being currently developed [13-15]. Lignin has been used to produce various polymers like ARBOFORM, polyesters and polyurethanes and various polymer blends with PVC, polyolefins, and rubbers are being currently developed [13-15].

In addition to these applications, lignin is a potential renewable source for many low molecular weight chemicals like benzene, phenol, guaiacol, vanillic acid, methanol, acetic acid, and dimethyl sulfoxide (DMSO) [17,18]. These lignin products are considered “value-added” chemicals (Figure 2) that could substantially impact the profit margins of a lignocellulosic biorefinery, but significant hurdles remain before they can be fully realized. One of the most significant of these is the realization of an efficient, cost-effective, and scalable means of fractionating lignocellulose into polysaccharide- and lignin-rich streams that enable downstream conversion into fuels and chemicals. Recently, some ionic liquids (ILs) have been shown to be very effective in the delignification of lignocellulosic biomass and depolymerization of lignin [19-23]. While much of the focus on ILs in the biomass field has been on their impact on the crystalline nature of cellulose, little attention has been paid in their potential utility as a means to valorize lignin.

Recent efforts in converting lignin to its monomeric products using ILs have focused on the technical lignins extracted from lignocellulosic biomass [25-27]. Cox and Ekerdt have shown that during IL dissolution, lignin
depolymerization occurs through breakdown of alkyl-aryl ether linkages [26]. But no lignin breakdown products were reported to be observed during this process. Through electro-catalytic oxidative cleavage of lignin, Reichert et al. produced various aromatic compounds like guaiacol, vanillic acid, vanillin, acetovanillone, syringol, syringaldehyde, and syringic acid from alkali lignin [25]. Though a total yield of 6% was observed, no information about the relative quantities of each compound was reported. Stark et al. produce syringaldehyde and 2,6-dimethoxy-1,4-benzoquinone from oxidative depolymerization of Beech lignin. Although an impressive yield (66.3%) of the products was obtained the process utilized high pressure (84 × 10^5 Pa of air) and very long reaction times (24 h) [27]. Lignin cleavage to monomers has also been accomplished using Bacillus sp. LD003 by Bandounas et al., but the incubation times for microbial degradation were very long (1–2 days) [28]. In this work, we investigate the ability of the IL 1-ethyl-3-methylimidazolium acetate ([C2mim][OAc]) to depolymerize lignin and produce valuable products from a series of technical lignins (kraft lignin, low sulfonate alkali lignin) as well as samples of switchgrass, pine, and eucalyptus during pretreatment.

Results and discussion
Extraction of lignin byproducts from IL pretreatment using [C2mim][OAc]
Measured amounts of three different biomass types representing grasses, softwood and hardwood (switchgrass, pine, and eucalyptus respectively) and technical lignins (kraft and low sulfonate alkali) were treated with [C2mim][OAc] at 160 and 120°C for 6 hrs. Since IL and water are both polar, recovery of polar lignin breakdown products posed significant challenges. Our choice of extraction solvents for non-polar products included pentane, hexane, heptane and benzene. Out of the solvents tried, benzene enabled the most recovery and hence we have included the data on benzene extracted products in this report. Although yields were not the same using different solvents, patterns of lignin degradation and recovery were similar for all the solvents tested. Table 1 shows the percent biomass recovered for various types of biomass pretreated at 160°C. For all the conditions studied there is a loss of mass observed, indicating that lignin and other biomass constituents remain solubilized in the supernatant. At low biomass loading levels, low sulfonate alkali lignin showed the maximum solubilization, followed with switchgrass. Percent recoveries were found to be similar for kraft lignin and eucalyptus. Interestingly, at higher loadings, the extent of solubilization was found to be very different and the observed extent of solubilization at 20% loadings were switchgrass = eucalyptus > pine dust = low sulfonate alkali > kraft lignin. In addition, similar extent of mass solubilization from technical lignin and lignocellulosic biomass indicates high levels of impurities (other material than lignin) present in the technical lignins we used.
Lignin breakdown products from technical lignins

The non-polar lignin breakdown products extracted from the supernatant, along with their elution times during GC-MS, are shown in Table 2. The lignin breakdown products are observed to depend on the sample type and the dissolution temperature. For the technical lignins studied, guaiacol and allyl guaiacol were the major products at all biomass loadings (Figure 3). Higher quantities of guaiacol (5 g/kg of biomass) were produced from kraft lignin when compared to low sulfonate alkali lignin. The quantity of guaiacol produced (per kg of starting material) decreases as a function of increasing biomass loading. Higher quantities of ally guaiacol are produced from low sulfonate alkali lignin (2 g/kg) than from kraft lignin (1 g/kg). Other products like methyl guaiacol, ethyl guaiacol, vinyl guaiacol, vanillin, guaiacyl acetone are also present at smaller concentrations. Production of these minor products is observed to increase with increases in biomass loading from 3 wt% to 10 wt %, but is observed to decrease on further increasing the biomass loading to 20 wt%. Similar quantities of guaiacyl acetone are produced on dissolution of both kraft lignin and low sulfonate alkali lignin. Higher quantities of ethyl guaiacol, vinyl guaiacol and vanillin were produced from kraft lignin, whereas a higher quantity of methyl guaiacol was obtained from low sulfonate alkali lignin. As these technical lignins were derived from softwood [29,30] and contain very small quantities of S-lignin in the original feedstocks, syringyl compounds were not significant. Similar compounds were observed by Stark et al. from the oxidative depolymerization of beech lignin [27] and by Reichert on electrolysis oxidative cleavage of alkali lignin [25].

The acidic or basic dissolution conditions that may be involved during the dissolution of lignin in IL can be predicted based on the products formed after dissolution. Vanillin has been previously produced from alkaline treatment and nitrobenzene oxidation of kraft lignin [31]. Adler et al. show the formation of guaiacol as a result of acidolysis of guaiacylglycerol-β-guaiacyl ether [32]. They show that the guaiacol is the main product due to the cleavage of β-aryl ether linkages. Adler et al. (1966) also predict guaiacylacetone to be one of the products of lignin acidolysis, but this monomer was present in small quantities in our supernatant solutions. Formation of aldehyde derivatives (coniferaldehyde) like allyl guaiacol during acid pretreatment of lignin is well cited in the literature [31-33]. Presence of higher quantities of guaiacol and allyl guaiacol indicated acidic dissolution conditions in IL under these conditions. We have recently published on the dual acidic and basic behavior of [C2mim][OAc] as a function of temperature, and the products identified here strongly confirm these previous findings [34].

Lignin breakdown products from switchgrass

The total amount of byproducts produced from lignocellulosic biomass was observed to be lower than that produced from technical lignins. This is reasonable given that lignin constitutes only ~1/3 of the dry weight of biomass. Guaiacol was obtained on dissolution of all lignocellulosic biomass in [C2mim][OAc] at 160°C for 6 hrs (Figure 4). Allyl guaiacol and syringol are also produced by dissolution of switchgrass (Figure 4a). As in the case of kraft lignin, the amount of guaiacol and allyl

---

Table 1 Percent biomass recovered as a function of biomass loading during dissolution

| Biomass       | Biomass loading during dissolution | % Recovered after dissolution |
|---------------|-----------------------------------|------------------------------|
| Kraft Lignin  | 3%                                | 60 ± 2                       |
|               | 10%                               | 69 ± 7                       |
|               | 20%                               | 84 ± 4                       |
| Low Sulfonate Alkali Lignin | 3%                         | 33 ± 1                       |
|               | 10%                               | 43 ± 1                       |
|               | 20%                               | 45 ± 3                       |
| Switchgrass   | Untreated                         | N/A                          |
|               | 3%                                | 40 ± 1                       |
|               | 10%                               | 41 ± 4                       |
|               | 20%                               | 35 ± 1                       |
| Pine Dust     | Untreated                         | N/A                          |
|               | 3%                                | 78 ± 5                       |
|               | 10%                               | 56 ± 1                       |
|               | 20%                               | 44 ± 5                       |
| Eucalyptus    | Untreated                         | N/A                          |
|               | 3%                                | 64 ± 1                       |
|               | 10%                               | 58 ± 1                       |
|               | 20%                               | 36 ± 1                       |

| Name          | Compound                     | RT*  |
|---------------|------------------------------|------|
| 1             | Guaiacol                     | 8.8  |
| 2             | 4-Ethyl Guaiacol             | 10.4 |
| 3             | 4-Vinyl Guaiacol             | 11.1 |
| 4             | Eugenol                      | 11.61|
| 5             | Syringol                     | 12.03|
| 6             | 4-(1-propenyl) Guaiacol      | 12.97|
| 7             | Vanillin                     | 13.54|
| 8             | Allyl Syringol               | 14.49|
| 9             | Guaiacylacetone              | 14.14|
| 10            | Anthracene-d10               | 16.97|

*Retention time (signifies retention time of the compounds in the GC/MS).
* Internal standard.
guaiacol produced decrease with increase in biomass loading. The production of syringol-type lignin compounds like syringol, allyl syringol and aceto syringone increases with increase in biomass loading. This increase implies breakdown of syringyl lignin increases with increase in biomass loading. All the minor products (methyl guaiacol, ethyl guaiacol, vinyl guaiacol, guaiacyl acetone and acetosyringone) except vanillin are produced on dissolution of switchgrass (Figure 4b), and the amount of these products increases with biomass loading. The increase in production of these minor compounds and decrease in production of major compounds indicates incomplete breakdown of lignin with increase in biomass loading.

**Lignin breakdown products from pine**

Guaiacol and allyl guaiacol were the only major products from dissolution of pine (Figure 4c). This is expected as pine consists of mostly guaiacyl lignin and has low (or no) syringyl lignin. In the case of pine, no correlation between the biomass loading and the total amount of guaiacol released was observed. The quantity of guaiacol and allyl guaiacol produced decreases on increasing the biomass loading from 3% to 10% and further increases on increasing the biomass loading to 20%. Methyl guaiacol, ethyl guaiacol, vinyl guaiacol and guaiacyl acetone are produced as minor products from pine (Figure 4d). The amount of guaiacyl acetone decreases with increase in biomass loading. At higher biomass loadings there is a small increase in the minor products generated.

**Lignin breakdown products from eucalyptus**

Unlike pine, eucalyptus contains a higher amount of syringyl lignin than guaiacyl lignin and all major products (guaiacol, allyl guaiacol, syringol and allyl syringol) are produced (Figure 4e). The amount of guaiacol and allyl guaiacol decreases with increases in biomass loading. The decrease in the production of these compounds indicated decrease in guaiacyl-lignin breakdown with increase in biomass loading. But the quantity of syringol produced increases on increasing the biomass loading from 3% to 10% and decreases on further increasing the biomass loading. Similar to dissolution of switchgrass, breakdown of guaiacyl-lignin decreases and syringyl-lignin increases with increase in biomass loading. At higher biomass loading levels of eucalyptus, increasing quantities of ethyl guaiacol, vanillin, guaiacyl acetone and acetosyringone were produced (Figure 4f).
As shown in the case of kraft lignin, changing the dissolution temperature changes the products that can be recovered from lignocellulosic biomass. A higher quantity of unsaturated guaiacols and aldehydes can be produced by decreasing dissolution temperature, and although not tested here, reaction time. It has been previously reported that vinyl guaiacol is produced from switchgrass when it is pretreated under alkaline conditions [32]. It has also been reported in the literature that guaiacylacetone is produced under acidic treatment conditions [32]. Guaiacol and syringol have been reported as the lignin products under acidic pretreatment conditions [32, 33]. The reaction mechanisms involved in the formation of these compounds need to be further investigated.

**Temperature dependence of lignin breakdown products**

It is clear from the above discussions that the patterns of degradation products are very different from different biomass and also at different processing temperatures. Based on the final desired products, the dissolution conditions can be tuned to optimize the recovery of certain products. For example, the cleavage of the methyl ketone group from vanillin at higher temperature leads to guaiacol formation, and therefore if it is desired to generate

![Figure 4 Major and minor lignin breakdown products from switchgrass (a & b), pine (c & d) and eucalyptus (e & f) after dissolution at 160°C for 6 hrs with different biomass loading.](image-url)
more vanillin from biomass it can be obtained in higher quantities by lowering the process temperatures to 120°C (Figure 5). Similarly eugenol and vinyl guaiacol show the most dramatic impact of dissolution temperatures at the two temperatures studied. These examples provide evidence of a very flexible IL technology for lignin breakdown and product optimization. It is important to note that since the putative lignin glass transition temperature (polymeric lignin softens at broad temperature range instead of having a sharp melting point) is around 140-165°C based on the source of lignin [35], it is expected that polymeric lignin will be increasingly depolymerized at higher temperatures. The unique characteristics of [C2mim][OAc] induced behavior makes it a promising technology for the selective production of these chemicals while also serving as an efficient means of pretreating biomass [34].

Conclusion
In this study [C2mim][OAc] was used to produce monomeric aromatic compounds from two types of technical lignins and three types of lignocellulosic biomass (pine, switchgrass, and eucalyptus) during pretreatment. Several guaiacyl monomers were found to be present in the supernatant of technical lignins and biomass samples after dissolution in [C2mim][OAc] at 160°C for 6 hrs. Guaiacol was the common product from both technical lignins and biomass, and was produced at higher levels at 3% biomass loading. Higher biomass loadings did not generate more products per kg of starting material. Syringyl monomers were produced on dissolution of switchgrass and eucalyptus. Furthermore, the dissolution conditions can be changed to produce higher amount of the desired byproduct. The total amount of non-polar lignin products ranged from 0.5–5.7 g/kg of biomass. The amount of vanillin produced ranged from 0.04–2.0 g/kg of biomass under two processing temperatures tested. The products reported in this work only represents the non-polar monomeric components that were soluble in benzene after dissolution. Polar lignin products are expected to be present in the supernatant and efforts are underway to extract and quantify them. These results indicate that certain ILs used for pretreatment may also hold significant promise in the conversion of polymeric lignin to smaller aromatics and desired renewable chemical outputs.

Materials and methods
Kraft lignin was supplied from MeadWestvaco Corp., Richmond, VA. Low sulfonate alkali lignin was purchased from Sigma Aldrich. Switchgrass (Panicum virgatum, cultivar MPV2) was provided by the laboratory of Dr. Ken Vogel. Samples of Pinus radiata and Eucalyptus globulus were provided by Arborgen. 1-ethyl-3-methylimidazolium acetate ([C2mim][OAc], 98% purity) was used as the solvent to depolymerize and dissolve lignin from the biomass. Benzene was used as the extraction solvent. Guaiacol, ethyl guaiacol, vinyl guaiacol, vanillin, eugenol, syringol, 4-allyl syringol, guaiacyl acetone and 2-methoxy, 4-propenyl phenol were used as standards to confirm the position and the mass to charge ratio of the Gas Chromatography/Mass Spectrometry (GC/MS) peaks. All other chemicals used in this study were purchased from Sigma Aldrich and used as received.

Figure 5 Lignin breakdown products (vanillin and eugenol) from low sulfonate alkali lignin and kraft lignin after dissolution at 120 and 160°C for 6 hrs at 3% biomass loading.
Lignin dissolution in IL
Samples were ground to 40 mesh before the dissolution process (Thomas-Wiley Mini Mill fitted with a 40-mesh screen; Model 3383-L10 Arthur H. Thomas Co., Philadelphia, PA, USA). Technical lignins and biomass were dissolved in [C2mim][OAc] at 120 and 160°C in a conventional oven (Thelco Laboratory Oven, Jouan Inc, Virginia) for 6 hrs. Solid loading was varied from 3 wt% (300 mg in 9.7 mL of [C2mim][OAc]) to 10 wt% and 20 wt%, and components were mixed at room temperature before being placed in the oven. To this mixture 10 μl of anthracene-D_{10} was added as internal standard (IS), for quantification in GC/MS analysis. 35 mL of hot water (95°C) was added to the sample to precipitate the dissolved biomass (mostly glucans, unsolubilized lignin). The mixture of [C2mim][OAc], water, and biomass was then centrifuged to separate the solid (recovered biomass) and liquid ([C2mim][OAc] and water). This mixture of [C2mim][OAc] and water will be referred to as the supernatant for the rest of this report. After the collection of supernatant, the biomass was further washed ten times with 100 mL of water (10 mL/wash).

Lignin extraction from the supernatant
A total of 10 mL of benzene was added to the supernatant in two steps. The resultant mixture (benzene+ sample) was mixed thoroughly and was then phase separated using a centrifuge. This allowed for all the non-polar compounds to be extracted from the supernatant to the benzene phase. This benzene solution was further concentrated under nitrogen to a final volume of 1 mL.

Gas chromatography–mass spectrometry (GC-MS)
The analysis of the lignin breakdown compounds present after benzene recovery was performed using a GC-MS (Thermo Electron Corporation with Trace GC Ultra, Polaris-Q MS and TriPlus auto sampler). The compounds were separated using a TR-SMS (30 m, 0.25 mm ID, 0.25 μm) chromatographic column. 10 μl of the sample was injected into the GC at an inlet temperature of 220°C and was operated in a split mode (split flow of 12 mL/min, split ratio = 10). Helium was used as a carrier gas with a constant flow rate of 1.2 mL/min. The temperature of the GC was held at 45°C for 1 min, was then increased at a rate 10°C/min up to 300°C and was held at this temperature for 1 min. The MS was used until the end of GC run with a solvent delay of 3.5 min. The ion source was maintained at a temperature of 250°C and the MS was operated in scan mode. Anthracene-D_{10} was used as the internal standard as it is not present in the biomass samples. The standards of each compound were used to calculate the individual response factor.

Quantification: calculation of product yield
Single-point calibration was used to calculate the response factor of the eluted compounds with respect to the internal standard (IS). Standards containing 2 mM of the compound and 20 μg of IS in 1 mL of Benzene were used for calibration. The area under the spectral peak of the compound and the IS were used to calculate the response factor (RF) (equation 1).

\[
RF = \frac{\text{Area}_{\text{compound}}/\text{weight of the compound injected}}{\text{Area}_{\text{IS}}/\text{weight of IS injected}}
\]

RFs were determined independently for each standard using GC/Ms.

The amount of each product was calculated as

\[
\text{Amount of product A in the supernatant (g)} = \frac{\text{Amount of the starting biomass (Kg)}}{\text{Amount of product A in the supernatant (g)}}
\]

Competing interests
The strategy described in this paper has been included in a patent application.

Authors’ contributions
SS designed and coordinated the study; PV, PS, MA, PDA, BAS and SS conducted the experiments and data analysis. PV, BAS and SS wrote the manuscript and all authors read and approved the final manuscript.

Acknowledgements
This work was part of the DOE Joint BioEnergy Institute (http://www.jbei.org) supported by the US Department of Energy, Office of Science, Office of Biological and Environmental Research, through contract DE-AC02-05CH11231 between Lawrence Berkeley National Laboratory and the US Department of Energy.

Received: 9 September 2012 Accepted: 2 January 2013 Published: 28 January 2013

References
1. Binder JB, Raines RT: Fermentable sugars by chemical hydrolysis of biomass. Proc Natl Acad Sci USA 2010, 107:4516–4521.
2. Sun Y, Cheng J: Hydrolysis of lignocellulosic materials for ethanol production: a review. Bioresour Technol 2002, 83:1–11.
3. Louime C, Uckelmann H: Cellulosic ethanol: securing the planet future energy needs. Int J Mol Sci 2008, 9:838–841.
4. Jørgensen H, Kristensen JB, Felby C: Enzymatic conversion of lignocellulose into fermentable sugars: challenges and opportunities. Biofuels, Bioprod Biorefin 2007, 1:119–134.
5. Petrus L, Noordermeer MA: Biomass to biofuels, a chemical perspective. Green Chemisrty 2006, 8:861–867.
6. Jurgen E: Secondary metabolites and plant defense. In Plant Physiology. 4th edition. Edited by Lincoln T, Eduardo Z. Sunderland, Massachusetts: Sinauer Associates, Inc; 2006:316–341.

7. NNFC: Renewable Chemicals Factsheet: Lignin. http://www.mnfcc.co.uk/publications/nnfcc-renewable-chemicals-factsheet-lignin.

8. Lauten RA, Myrvold BO, Gundersen SA: New Developments in the Commercial Utilization of Lignosulfonates. In Surfactants from Renewable Resources. John Wiley & Sons, Ltd; 2010: 269–283.

9. Hoyt CH, Goheen DW: Utilization of Lignin- Polymeric Products. In Lignins. Edited by Sarkanen KV, Ludwig CH. USA: John Wiley & Sons, Inc; 1971:833–859.

10. Northey RA: The Use of Lignosulphonates as Water Reducing Agents in the Manufacture of Gypsum Wallboard. In Chemical Modification, Properties, and Usage of Lignin. Edited by Hu TQ. New York, USA: Kluwer Academic/Plenum Publishers; 2002:139–150.

11. Berry BC, Viswanathan MN: Lignosulonic Acid-Doped Polyaniline (Ligno-Pani)- A Versatile Conducting Polymer. In Chemical Modification, Properties, and Usage of Lignin. Edited by Hu TQ. New York, USA: Kluwer Academic/Plenum Publishers; 2002:211–40.

12. Kodia JF, Kubo S, Gilbert RD, Venditti RA: Lignin- based Carbon Fibers. In Chemical Modification, Properties, and Usage of Lignin. Edited by Hu TQ. New York, USA: Kluwer Academic/Plenum Publishers; 2002:121–138.

13. Nagel H, Pfister J, Nagele E, Ionee ER, Eisenreich N, Eckl W, Eyerer P; Arboform- A thermostatic, processable material from lignin and natural fibers. In Chemical Modification, Properties, and Usage of Lignin. Edited by Hu TQ. New York, USA: Kluwer Academic/Plenum Publishers; 2002:101–120.

14. Feldman D: Lignins and its Polyblends-A Review. In Chemical Modification, Properties, and Usage of Lignin. Edited by Hu TQ. New York, USA: Kluwer Academic/Plenum Publishers; 2002:281–100.

15. Montanari, Gandini A, Belgacem MN, Guo Z-X: Lignins as Macromonomers for Polymers and Polyurethanes. In Chemical Modification, Properties, and Usage of Lignin. Edited by Hu TQ. New York, USA: Kluwer Academic/Plenum Publishers; 2002:57–80.

16. Dimmel DR, Bozelli JJ, Oepen DG, Savidakis MC: Pulping Catalysts from Lignin-the Diels-Alder Step. In Chemical Modification, Properties, and Usage of Lignin. Edited by Hu TQ. New York, USA: Kluwer Academic/Plenum Publishers; 2002:199–220.

17. Goheen DW: Utilization of Lignin: Low Molecular Weight Chemicals. In Lignins. Edited by Sarkanen KV, Ludwig CH. USA: John Wiley & Sons, Inc; 1971:797–824.

18. Gandini A: The irruption of polymers from renewable resources on the scene of macromolecular science and technology. Green Chemistry 2011, 13:1061–1083.

19. Singh S, Simmons BA, Vogel KP: Visualization of biomass solubilization and cellulose regeneration during ionic liquid pretreatment of switchgrass. Biotechnol Bioeng 2009, 104:586–75.

20. Stark K, Taccardi N, Bosmann A, Wasserscheid P: Oxidative depolymerization of lignin in ionic liquids. ChemSusChem 2010, 3:719–723.

21. George A, Tran K, Morgan TJ, Benke PI, Berrueco C, Lorente E, Wu BC, Keasling JD, Simmons BA, Holmes BM: The effect of ionic liquid cation and anion combinations on the macromolecular structure of lignins. Green Chemistry 2011, 13:3375–3385.

22. Arora R, Manisseri C, Li CL, Ong MD, Scheller HV, Vogel K, Simmons BA, Singh S: Monitoring and Analyzing Process Streams Towards Understanding Ionic Liquid Pretreatment of Switchgrass (Panicum virgatum L). BioEnergy Research 2010, 3:134–145.

23. Cheng G, Kents MS, He L, Varanasi P, Dibble D, Arora R, Deng K, Hong K, Melnichenko YB, Simmons BA, Singh S: Effect of Ionic Liquid Treatment on the Structures of Lignins in Solutions: Molecular Subunits Released from Lignin. Langmuir 2012, 28:11850–11857.

24. Holladay JE, Bozelli JJ, White JF, Johnson D: Top Value-Added Chemicals from Biomass Volume II—Results of Screening for Potential Candidates from Bioenergy Lignin; 2007: Report prepared by Pacific Northwest National Laboratory for US DOE, Report number PNNL-16983, available for download at http://www1.eere.energy.gov/biomass/pdfs/pnnl-16983.pdf.

25. Reichert E, Wintringer R, Volmer DA, Hempelmann R: Electro-catalytic oxidative cleavage of lignin in a protic ionic liquid. Phys Chem Chem Phys 2012, 14:5214–5221.

26. Cox BJ, Ekerdt JG: Depolymerization of oak wood lignin under mild conditions using the acidic ionic liquid 1-H-3-methylimidazolium chloride as both solvent and catalyst. Bioresource Technol 2012, 118:584–588.