Gale, E., Wirawan, R. H., Silveira, R. L., Pereira, C. S., Johns, M. A., Skaf, M. S., & Scott, J. L. (2016). Directed discovery of greener cosolvents: new cosolvents for use in ionic liquid based organic electrolyte solutions for cellulose dissolution. *ACS Sustainable Chemistry and Engineering*, 4(11), 6200-6207.
<https://doi.org/10.1021/acssuschemeng.6b02020>

Peer reviewed version

Link to published version (if available):
[10.1021/acssuschemeng.6b02020](https://doi.org/10.1021/acssuschemeng.6b02020)

[Link to publication record in Explore Bristol Research](#)
PDF-document

This is the author accepted manuscript (AAM). The final published version (version of record) is available online via ACS at <https://pubs.acs.org/doi/10.1021/acssuschemeng.6b02020>. Please refer to any applicable terms of use of the publisher.

University of Bristol - Explore Bristol Research

General rights

This document is made available in accordance with publisher policies. Please cite only the published version using the reference above. Full terms of use are available:
<http://www.bristol.ac.uk/red/research-policy/pure/user-guides/ebr-terms/>

Directed Discovery of Greener Cosolvents: New Cosolvents for Use in Ionic Liquid Based Organic Electrolyte Solutions for Cellulose Dissolution

Ella Gale,^{†,‡} Remigius H. Wirawan,^{†,‡} Rodrigo L. Silveira,[§] Caroline S. Pereira,[§] Marcus A. Johns,^{||,⊥} Munir S. Skaf,[§] and Janet L. Scott^{*,†,‡}

[†]Department of Chemistry, [‡]Centre for Sustainable Chemical Technologies, ^{||}Department of Chemical Engineering, and [⊥]EPSRC Centre for Doctoral Training in Sustainable Chemical Technologies, University of Bath, Bath, BA2 7AY, U.K.

[§]Institute of Chemistry, University of Campinas, Campinas, Sao Paulo, 13084-862, Brazil

Supporting Information

ABSTRACT: Cellulose is an abundant, cheap, renewable, yet recalcitrant, material, which, if dissolved, may be formed into a wide range of materials, composites, and mixtures. Much attention has recently been focused on the use of mixtures of ionic liquids and some solvents (so-called organic electrolyte solutions, OESs) as efficient cellulose dissolution solvents, but many of the cosolvents used lack green credentials—a perennial problem where dipolar aprotic solvents are the solvents of choice. We present a rational approach, based on definition of ranges of solvent parameters gathered together in recently published databases, to find “greener” cosolvents for OES formation. Thus, γ -butyrolactone is identified as a suitable OES former for dissolution of microcrystalline cellulose and biobased γ -valerolactone as a marginally less efficient, but significantly safer, alternative. Comparison of cosolvent efficiency reveals that previous use of measures of mass, or concentration, of cellulose dissolved may have masked the similarities between 1-methylimidazole, dimethyl sulfoxide (DMSO), *N,N*-dimethylformamide, *N,N'*-dimethylimidazolidinone, *N,N*-dimethylacetamide, *N*-methylpyrrolidinone, and sulfolane (seldom considered), while comparison on a molar basis reveals that the molar volume of the solvent is an important factor. Reference-interaction site model (RISM) calculations for the DMSO/1-ethyl-3-methylimidazolium acetate OES suggest competition between DMSO and the acetate anion and preferential solvation of cellulose by the ionic liquid.

KEYWORDS: 1-Ethyl-3-methylimidazolium acetate, Ionic liquid, Cellulose, Dissolution, Organic electrolyte solutions (OESs), Reference-interaction site model (RISM), γ -Butyrolactone, γ -Valerolactone

INTRODUCTION

Cellulose is an abundant and renewable biopolymer produced in large quantities in the plant kingdom. Estimates vary, but of the 56×10^9 tonnes of biomass produced annually, approximately 40–60% is cellulose.¹ It is a critical structural biopolymer, providing rigidity to plants cell walls, and, while highly hydrophilic, it is also very difficult to solubilize and, thus, to reform into new structures, forms, or composites. As it is not a thermoplastic polymer, processing of cellulose often relies on functionalization to yield a soluble polymer (which means sacrificing its exquisite structural characteristics resulting from hydrogen bonding) or dissolution, either using solvent mixtures, or via reversible reaction, e.g. the xanthate process for producing viscose.² Alternative direct dissolution methods do exist, such as the Lenzing process, using *N*-methylmorpholine-*N*-oxide hydrate (NMMO) to dissolve cellulose, but for various reasons, including mitigation of use of hazardous materials (e.g., CS₂ in the xanthate process), excessive cost, difficulties in recovery and

recycling of solvent systems, and ease of application, alternatives are still sought.

A seminal paper by Swatloski and Rogers, describing the dissolution of cellulose in ionic liquids (ILs) followed by antisolvent phase inversion to yield regenerated cellulose in a different form,³ precipitated an avalanche of reports of both similar solvent systems and production of cellulose based materials using this method of processing.^{4–10} A recent review provides an excellent overview of regenerated cellulose materials, including those from ILs or containing IL mixtures.¹¹ Positive aspects of ionic liquid based dissolution/regeneration of cellulose

Special Issue: Building on 25 Years of Green Chemistry and Engineering for a Sustainable Future

Received: August 21, 2016

Revised: September 27, 2016

Published: September 28, 2016

include the lack of vapor pressure of the ionic liquid (mitigating fugative emissions, such as those always attendant on, for example, the xanthate process, or dissolution in NMMO) and the apparent ease of recycling of the solvent salt by recovery of the antisolvent using distillation. The latter is facilitated by the vast difference in vapor pressure between antisolvents, such as lower alcohols or water, and the ionic liquid. It should be noted that such separation is seldom as facile as reported, can be costly with respect to energy use, and final removal of traces of water can be challenging,¹² although rigorously dried ILs are not required for cellulose dissolution;¹³ recently, ILs containing 40 wt % water (effectively aqueous solutions of ILs) have even been reported for cellulose dissolution.¹⁴

A range of ILs has been demonstrated to dissolve cellulose, and combinations of 1,3-dialkylimidazolium cations with chloride, acetate, or alkylphosphate ester anions have proven particularly effective—data pertaining to solubility of cellulose in these, and a plethora of other ILs, are comprehensively summarized in a review by Wang, Gurau, and Rogers.¹⁵ In 2011, Rinaldi reported the use of combinations of ILs and dipolar aprotic solvents, mixtures which he termed “organic electrolyte solutions (OESs)” (these are analogues of well-known cellulose solvent systems, such as LiCl/dimethylacetamide), to dissolve cellulose,¹⁶ and this methodology has also been enthusiastically adopted by many, as the use of a cosolvent may mitigate the cost of the IL and facilitate processing by reduction of viscosity of cellulose solutions. The utility of OESs was presaged by an earlier report of the use of an amino acid derived IL, *N,N*-diethyl-*N*-(2-methoxyethyl)-*N*-methylammonium alanine, in combination with DMSO, to dissolve up to 23 wt % of cellulose.¹⁷

While many computational^{18–22} and experimental²³ studies dedicated to understanding the source of the enhanced dissolution^{24–26} (at least with respect to *rate* of dissolution, e.g. Rinaldi’s “instantaneous dissolution”¹⁶), or tailoring OESs²⁷ have appeared, there has not, to our knowledge, been a systematic study geared toward maximizing the efficiency of dissolution while minimizing the quantity of IL required. As many of the objections to the use of ILs at scale are based on cost, minimizing the quantity of IL required could be key to the production at scale of any of the interesting materials and composites reported.

Although there is no doubt that the discovery of OES solvent systems for cellulose dissolution has vastly expanded the range of solvents useful in cellulose processing, the dipolar aprotic solvent components of OESs present some health and safety challenges. For example, dimethylformamide carries the Globally Harmonized System of Classification and Labeling of Chemicals (GHS) phrase H360 “*may damage fertility or the unborn child*” and is included in the candidate list of Substances of Very High Concern under REACH regulation (Article 57c), and dimethyl sulfoxide (the most effective cosolvent in Rinaldi’s list), which may be an attractive replacement on toxicological grounds, has a propensity to undergo highly energetic decomposition when subjected to strong bases, electrophiles, and heat.²⁸

Hydrogen bond basicity has long been held to be an important factor in cellulose dissolution by ILs,²⁹ and Rinaldi considered the effect of addition of IL to dipolar aprotics on Kamlet–Taft solvent parameters and suggested that the molecular solvent (and the IL) should have high hydrogen-bond basicity (β) and no hydrogen-bond acidity (α).¹⁶ While this is a useful guide, many potential solvents that fulfill these requirements are not good cosolvents for cellulose and it is often the absolute amount of cellulose that can be dissolved rather than “instantaneous dissolution” that is the required quantity. Herein, we report the

relative efficacy of a range of cosolvents for the dissolution of microcrystalline cellulose using the ionic liquid 1-ethyl-3-methylimidazolium acetate ([EMIm][OAc]) (one of the most widely used ILs for cellulose dissolution) to form OESs and use comparisons of solvent parameters to reveal new candidates for OES formation. The focus is on using a rational approach to selecting the greenest possible cosolvent for preparation of cellulose solutions.

EXPERIMENTAL SECTION

Materials. 1-Ethyl-3-methylimidazolium acetate (>95% purity, Sigma-Aldrich) and microcrystalline cellulose (Sigma-Aldrich, 20 μ m particle size) were dried at 80 °C under reduced pressure to constant mass. Dipolar aprotic solvents (all >99% purity, Sigma-Aldrich) were used as received, except for sulfolane, which was dried over activated 3 Å molecular sieves. The water content of the ionic liquid and solvents was measured by Karl Fisher titration and found to be: [EMIm][OAc] 1.1%; 1-methylimidazole 0.2%; dimethyl sulfoxide 1.1%; *N,N*-dimethylformamide 0.3%; *N,N'*-dimethylimidazolidinone 0.7%; *N,N*-dimethylacetamide 0.4%; sulfolane 0.1%; propylene carbonate 0.1%; γ -valerolactone 0.8%; *N,N,N',N'*-tetramethylurea 0.3%; *N*-methylpyrrolidinone 0.8%; and γ -butyrolactone 0.3%.

Solubility Tests. Solubility tests were conducted at 70 °C. As the cellulose solutions formed are viscous and not amenable to filtration, solubility was determined by addition of small aliquots of dried microcrystalline cellulose (MC) to mixtures of [EMIm][OAc]/cosolvent of appropriate composition. The maximum amount of MC that dissolved, a , and minimum amount that did not dissolve, b , were recorded, giving an under- and overestimate for the maximum amount of MC dissolvable, v . Repeated determinations, using smaller incremental added weights of MC then led to a narrow range between “soluble” and “insoluble”. (As MC tended to form aggregates when added directly to OESs, a slightly modified procedure was followed once approximate solubility values were known at any given OES composition: to a weighed quantity of dried MC was added a measured quantity of the selected cosolvent; after stirring for 1 min to wet and disperse the MC powder, calculated quantities of IL were added, and then, small, weighed aliquots were added until no further MC would dissolve.) Apparently “insoluble” samples were maintained for a minimum of 16 h at 70 °C to confirm the lack of further dissolution.

Each pair of points (a_i , b_i) was given a weight, w_i :

$$w_i = \sqrt{\frac{1}{v_i^2}} \quad (1)$$

based on, v_i , the errors on a_i and b_i , as given by $v_i = (b_i - a_i)/2$, such that the uncertainties on points a_i and b_i are $a_i + v_i$ and $b_i - v_i$ respectively. Other sources of error are much smaller in comparison.

Data Analysis and Fitting. Weighted fits were computed using NonLinearModelFit in Mathematica, with weights w_i as given in eq 1 and the following

$$c = 1 + \text{Max}[c] - e^{(m\chi_{\text{IL}} + d)^2} \quad (2)$$

which arises from modeling dissolution as a one-dimensional process, where $\sqrt{\ln[\Delta c]}$ varies linearly with χ_{IL} ($\Delta c = 1 + \text{Max}[c] - c$), c is the weight percent of cellulose, and $\text{Max}[c]$ is the maximum measured weight percent of dissolved MC. For comparison, the data were normalized using the maximum value of c from the fit. The maximum value of a fit over a range was computed using the Mathematica “Maximize” function within the range of measured data (e.g., $\text{Min}[\chi_{\text{IL}}] < \chi_{\text{IL}} \leq 1$). Weighted averages, $\bar{x}$, of a set of n points $\{x_i\}$ were computed from

$$\bar{x} = \frac{\sum_{i=1}^n w_i x_i}{\sum_{i=1}^n w_i} \quad (3)$$

Chart 1. Cosolvents Tested in Organic Electrolyte Solutions with the Ionic Liquid [EMIm][OAc]

Figure 1. (a) Normalized weight percent of MC dissolved in OESs versus mole fraction of ionic liquid, χ_{IL} (dissolved MC reaches a maximum of 22–23 wt % and curves of weight percent MC dissolved versus χ_{IL} appear in the Supporting Information). Points are pairs of over- and underestimates on the maximum amount of MC soluble in each OES at that composition, the uncertainties on these measurements are presented as error bars and used to determine their weighting in the fit (eq 1). Lines are the fits calculated using eq 2, yielding $R^2 > 0.99$ in all cases. (b) Dissolution efficiency expressed as grams of MC dissolved per gram of IL versus χ_{IL} , relative to the efficiency at $\text{Max}[c]$. Lines are transformed fitted curves. The best OESs can dissolve >70% of the maximum amount of MC soluble in pure IL at χ_{IL} as low as 0.3, with up to twice the efficiency per unit mass of IL.

using the WeightedData function in Mathematica. The “efficiency measure” was calculated as the mass of MC dissolved over the mass of ionic liquid, divided by the maximum mass of MC dissolvable.

Cosolvent Search. To search for cosolvents expected to yield OESs capable of dissolving MC on the basis of similarities in key solvent parameters, Excel spreadsheets of solvent parameters from Catalán³⁰ and Laurence et al.³¹ were loaded into Mathematica. Minimum and maximum values for each solvent parameter were input to define a search range, which could then be padded by a percentage of the selected range. A simple search over the data sets provided a count of the number of times each solvent (in the superset) appeared within the search range and the solvent list was sorted according to the number of “hits”. As not all the solvent entries were complete, the percentage data matched was also determined. Annotated sample Mathematica code is given in the Supporting Information. The solvent parameters were not normalized.

Reference-Interaction Site Model (RISM): 3D-RISM Theory. The 3D-RISM theory is a fully atomistic molecular theory of solvation based on first principle statistical mechanics that provides a computational framework to obtain solvation properties for a variety of molecular systems in solution. Starting from the molecular coordinates and interaction potentials, the ensemble-averaged solvation structure and thermodynamics can be effectively computed within the 3D-RISM approach. Details of the theory and the computational methods involved are provided elsewhere.^{32–34}

RISM: Computational Modeling and System Setup. The radial pair distribution functions for a mixture of molar fraction composition of 0.95 DMSO and 0.05 IL were obtained by the dielectrically consistent RISM theory (DRISM) complemented by the one-dimensional Kovalenko–Hirata (KH) closure.^{32,33,35} The pair distribution functions were later used as input for the 3D-RISM calculations. The DRISM equations were iteratively solved in a radial grid of 0.1 Å resolution containing $2^{13} = 8192$ points, considering the temperature of 298 K, the DMSO density of 1.091 g cm^{-3} , and dielectric constant of 46.4.³⁶ To solve the DRISM equations for the mixture, calculations were performed for increasing values of IL concentration, from 0 to 0.05. Solvent distribution around a glucan chain of DP = 12 previously equilibrated in DMSO was obtained using the 3D-RISM theory with the 3D Kovalenko–Hirata closure.^{32–34} The 3D-RISM equations were iteratively solved in grid of size $128 \text{ Å} \times 128 \text{ Å} \times 128 \text{ Å}$ containing

$256 \times 256 \times 256$ points. For both DRISM and 3D-RISM calculations, the modified direct inversion in the iterative subspace (MDIIS) solver³² was employed to reach a relative mean square accuracy of 10^{-10} for DRISM and 10^{-4} for 3D-RISM. Partial atomic charges and Lennard-Jones parameters for acetate and cellulose were taken from the CHARMM force field.^{37,38} The parameters for DMSO and EMIM were taken from Strader and Feller³⁹ and Kelkar et al.,⁴⁰ respectively. The solvent distribution was represented in terms of the specific solvation free energy density (SFED) function. SFED indicates spatial regions around the solute that concentrate solute–solvent stabilizing interactions.^{34,41,42} The specific SFED, defined as SFED per unit of solvent molecules, provides the intrinsic solvation capacity of each component of the solvent.

The glucan chain used in the 3D-RISM calculations was equilibrated in DMSO using molecular dynamics (MD) simulations. Using Packmol,⁴³ the glucan chain was immersed in a cubic box of side 90 Å containing 6000 DMSO molecules. The MD simulation was performed employing periodic boundary conditions. Electrostatics were evaluated with particle mesh Ewald⁴⁴ and the short-range interactions were truncated at 12 Å. A time step of 2 fs was used to solve the equations of motion. The bonds involving hydrogen atoms were constrained at their equilibrium lengths. The simulation was propagated for 2 ns at 300 K and 1 atm using NAMD.⁴⁵ Three different glucan conformations were randomly taken from the simulation for postprocessing with 3D-RISM.

RESULTS AND DISCUSSION

The goals of this study were twofold: (i) to compare the relative efficiency of IL use in cellulose dissolution in OESs comprised of different solvents with [EMIm][OAc] with a view to mitigating use of costly ILs and (ii) to use a rational approach to discover new greener solvents for the formation of OESs to be used in cellulose dissolution.

Microcrystalline cellulose (MC) was selected as the test substrate as its relatively low average degree of polymerization of 150–300⁴⁶ provides solutions of reasonable viscosity. The elevated temperature used was selected as readily accessible using a wide range of cosolvents of variable boiling points, while remaining below the temperatures at which cellulose degradation

has been shown to be significant.⁴⁷ (While it is acknowledged that an increase in temperature does not thermodynamically favor the dissolution of cellulose,⁴⁸ it does enhance the rate of dissolution.) The structures of all solvents tested as OES components with [EMIm][OAc] are presented in Chart 1 and included the following: 1-methylimidazole (1-MI); dimethyl sulfoxide (DMSO); *N,N*-dimethylformamide (DMF); *N,N'*-dimethylimidazolidin-2-one (DMI); *N,N*-dimethylacetamide (DMAc); sulfolane; γ -valerolactone (γ -val); γ -butyrolactone (γ -but); propylene carbonate (PC); *N,N,N',N'*-tetramethylurea (TMU); and *N*-methylpyrrolidinone (NMP).

As expected from Rinaldi's earlier results,¹⁶ addition of modest quantities of an IL to a range of dipolar aprotic solvents led to enhanced dissolution of MC, and as the mole fraction of IL, χ_{IL} , increased, the quantity of MC dissolved tended to the value soluble in pure [EMIm][OAc], Figure 1a.

Clearly, comparison of the OESs on the basis of mole fraction IL required to effect dissolution of a given quantity of MC suggests that some solvents, notably 1-MI, DMSO, DMF, etc., are "good" cosolvents, while others, such as TMU, are significantly less effective. This reflects the conclusion reached by Olsson et al. with respect to 1-MI⁴⁹ and expands on Rinaldi's findings (of instant dissolution),¹⁶ completing the data across a range of IL concentrations for a number of solvents. In general, the IL will be the most costly (and sometimes the least green) component of the OES, so we consider a measure of the efficiency of IL use in MC dissolution, Figure 1b. Differences between the relative efficacies of the solvents are starkly highlighted, with 1-MI yielding the greatest mass of MC dissolved per unit mass of IL. DMSO follows a close second, while DMF, DMAc, NMP, sulfolane, and DMI are similar, Table 1. TMU, in spite of its significant structural similarity to DMI, is

Table 1. Limiting Quantities of [EMIm][OAc] Required to Produce an OES for Dissolution of Cellulose for All Solvents Tested and Measures of the Efficiency of the OESs Formed

	minimum χ_{IL} required to initiate dissolution	χ_{IL} required for maximum efficiency	maximum efficiency measure
1-MI	0.052	0.168	2.16
DMSO	0.019	0.160	1.58
DMF	0.058	0.284	1.35
γ -but	0.065	0.318	1.30
DMI	0.096	0.333	1.45
DMAc	0.056	0.349	1.19
NMP	0.100	0.344	1.39
sulfolane	0.115	0.399	1.29
γ -val	0.184	0.609	1.20
TMU	0.451	0.752	1.08

the poorest solvent tested using the efficiency measure. Interestingly, the most efficient IL cosolvent, 1-MI, is not the solvent with the lowest limiting χ_{IL} required to effect dissolution of MC. Indeed, there are clues that 1-MI might be distinct from the other solvents as the shape of the curve reflecting quantity of cellulose dissolved versus IL concentration does not match that of all other solvents tested, possibly pointing to some mechanistic peculiarity. It is notable that this is the only cosolvent in the group with a nitrogen atom hydrogen bond acceptor (all others are oxygen atom acceptors); nonetheless, as our focus was on efficient and green cosolvents, this was not explored further.

The inclusion of some solvents, such as γ -but and γ -val, is not obvious and does not follow from the previous reports of instantaneous dissolution, or any of the plethora of publications that have drawn on that work for inspiration. In addition, the alert reader will note that PC has not yet been considered—we return to discussion of these greener solvents later.

While the weight percent MC dissolved in an OES of given composition is a useful comparative value, this treatment does not provide for direct comparison of the efficiency of the dissolution of MC in a given OES on a molar basis. Considering the mole fraction of the cellobiose unit versus the mole fraction of IL in the OES provides solubility curves that more accurately reflect the molecular composition of the solutions, Figure 2.

Figure 2. Mole fraction of the MC cellobiose repeat unit, χ_{Cell} (calculated from the previously normalized values), versus mole fraction of IL in the OES. As before, lines are transformed fits of eq 2. Presenting the data in this way removes the effect of cosolvent molar volume. All effective cosolvents now exhibit similar curves with only TMU and, to a lesser extent, γ -val appearing to be significantly different from the others.

Represented thus, on a comparative molar basis, the differences between the solvents are significantly reduced, and it becomes clear that the apparently significant differences noted by Rinaldi and highlighted in Figure 1 are due to the variable molar volume of the individual solvents. Thus, sulfolane, with $V_m = 95.27 \text{ cm}^3 \cdot \text{mol}^{-1}$ is a less effective cosolvent than DMSO, with $V_m = 71.30 \text{ cm}^3 \cdot \text{mol}^{-1}$, but by a far smaller margin than previously implied.

With efficiency considered, we turn to the discussion of solvent selection and greenness. Rinaldi and others used Kamlet–Taft parameters as the basis for comparison of solvents, but here we use the recently compiled databases of Catalán³⁰ and Laurence and co-workers³¹ providing solvent parameters for 160 and >300 solvents, respectively. Catalán defines four empirically derived parameters: solvent polarizability (SP); solvent dipolarity (SdP); solvent acidity (SA); and solvent basicity (SB). Notably, in this scale, the so-called nonspecific solvent/solute interactions are described by two parameters (polarizability and dipolarity), thus avoiding confounding these two different effects.³⁰ Laurence et al. define solvent parameters describing solute/solvent interactions: dispersion induction (DI); electrostatic (ES); solute Lewis base/solvent Lewis acid (α_1); and solute hydrogen bond donor/solvent hydrogen bond acceptor (β_1) interactions.³¹ Clearly these scales and parameters are not dissimilar, and it may be possible to select a single set of parameters to describe solvent/solute interactions. However, the goal here is to use such solvent parameter scales as tools for solvent selection. Specifically, we seek "greener" solvents for use in OESs for dissolution of cellulose to facilitate large scale production of novel materials, such as cellulose composites.

Table 2. Solvent Parameters from the works of Catalán³⁰ and Laurence³¹ for the Six Solvents Selected as Effective Cosolvents in OESs with [EMIm][OAc], γ -but, and Two Solvents with Some Structural Similarities with TMU and PC

	Catalán				Laurence			
	SP	SdP	SA	SB	DI	ES	α_1	β_1
DMSO	0.830	1.000	0.072	0.647	0.84	1.00	0.00	0.71
DMF	0.759	0.977	0.031	0.613	0.78	0.87	0.00	0.69
1-MI	0.834	0.959	0.069	0.658	0.86	0.97	0.00	0.70
sulfolane	0.830	0.896	0.052	0.365	0.85	0.93	0.00	0.34
NMP	0.812	0.959	0.024	0.613	0.83	0.80	0.00	0.76
DMAc	0.763	0.987	0.028	0.650	0.79	0.85	0.00	0.75
max ^a	0.834	1.000	0.072	0.658	0.86	1.00	0.00	0.76
min ^a	0.759	0.896	0.024	0.365	0.78	0.80	0.00	0.34
range ^a	0.075	0.104	0.048	0.293	0.08	0.20	0.00	0.42
percent of total range ^a	7.5	9.5	4.5	29.3	7.0	17.5	0.0	33.5
γ -but ^b	0.775	0.945	0.057	0.399	0.79	0.94	0.00	0.45
TMU ^c	0.778	0.878	0.000	0.624	0.81	0.71	0.00	0.75
PC ^d	0.746	0.942	0.106	0.341	0.77	1.07	0.00	0.40

^aThe minimum and maximum values of the six selected solvents were used to search the databases, and the percentage range over the parameter is included as a measure of specificity. ^bNew greener cellulose dissolving OES identified in this analysis. ^cA poor cosolvent not included in the range analysis—tested due to superficial structural similarity with DMI. ^dBears a superficial structural similarity to γ -but and γ -val.

While the terms solute and solvent may be a little confusing, here both [EMIm][OAc] and cellulose are common to all systems, thus the only variable to be considered is the cosolvent used to form the OES. Using the six solvents confirmed to form effective OESs for dissolution of cellulose when combined with [EMIm][OAc] that appear in both databases (DMSO, DMF, 1-MI, sulfolane, NMP, DMAc), we define ranges for each of the four parameters described by Catalán and Laurence, Table 2. These are represented graphically in Figure 3, and it is clear that the parameters describing these solvents fall into narrow bands.

Figure 3. Catalán and Laurence parameters presented as ranges for each of the four parameters defined in each database. The values for the six representative solvents selected are represented in red and it is clear that these fall into narrow bands (as expected for solvents with similar characteristics). Parameters describing the newly identified greener solvent, γ -but, are indicated as ---* and those for TMU (a rather poor solvent) as ---. (Note: solvent parameters have not been normalized, and some ranges have a small number of solvents with values >1, off scale in the figure.)

We hypothesized that interrogation of the Catalán³⁰ and Laurence³¹ databases (conveniently provided as Supporting Information for their publications) for solvents falling in, or close to, these ranges might offer up some candidates for new OES production. Remarkably, interrogation of both databases for solvents falling within the ranges defined by the parameters for the six selected solvents $\pm 5\%$ of the range yields only one further solvent expected to form an OES capable of efficiently dissolving MC: γ -but (full sorted lists are contained in the Supporting Information). Testing of this solvent showed that it does, indeed,

form an efficient cellulose dissolving OES, with a solubility curve falling well within the group comprised of the six solvents selected plus DMI (Figure 1) and with efficiency closely matching that of DMF and other dipolar aprotic solvents (Table 1 and Figure 2).

This encouraging result led to consideration of other solvents that might provide greener options for OES based processing of cellulose. While γ -but is not associated with any of the known, or potential, reproductive hazards identified for DMF, NMP, and others in the group, it nonetheless remains *harmful if swallowed* (H302), *causes serious eye damage* (H318), and *may cause drowsiness or dizziness* (H336). It is produced in large quantities as an industrial solvent and intermediate in the production of, for example, pyrrolidinones, but there are significant concerns about the ease with which it may be converted into γ -hydroxybutyrate (GHB), a compound listed in Schedule II of the 1971 UN Convention on Psychotropic Substances,⁵⁰ or simply ingested directly as an intoxicant (γ -but is metabolized to GHB).⁵¹ As the dose–effect curve is steep, the difference between doses yielding only “*euphoria, relaxation, reduced inhibition and sedation*” and those progressing to “*vomiting, urinary and fecal incontinence, agitation, convulsions, bradycardia, respiratory depression, coma and death*” can be slight.⁵¹ Thus, γ -butyrolactone is a controlled substance in some countries. In the hope of retaining solvent efficacy, but mitigating the potential for unwitting (or intentional) narcotic doses, the structurally similar compound, γ -val, was tested.

γ -Val proved to form a reasonably effective OES for cellulose dissolution when combined with [EMIm][OAc] (Figures 1 and 2) and would be an effective alternative to γ -but for generation of MC solutions significantly below the saturation concentration of MC in most of the OESs (~ 22 – 23 wt %). γ -Val can be produced from biomass⁵² and has appropriate physical properties for use as a solvent (including modest vapor pressure, limiting fugative emissions); its credentials as a “green” solvent have been described.⁵³ Finally, as it can be readily separated from water, γ -val may be more amenable to recovery post phase inversion in water—the process frequently used to generate cellulose composites, or formed, materials from OES solution.

Following the success of OES formation with two cyclic lactones (γ -but and γ -val), PC, a solvent which bears some

structural similarities to these, was (optimistically) tested. PC combined with [EMIm][OAc] was shown to form an OES capable of dissolving some cellulose, but required $>0.5\chi_{IL}$ and the solubility plateau noted for all other OESs was only approached in solutions of almost pure IL. Indeed, PC could perhaps be better described as a “diluent or contaminant that is not quite an anti-solvent for MC”. The solvent parameters for PC do not, at first glance, appear to be very far removed from the ranges indicated (Table 2), but inspection of the sorted lists (Supporting Information) suggests that PC is, indeed, not predicted to be a good OES former for cellulose dissolution. The values for SA and ES fall well outside the ranges defined.

The significant effect of small changes in molecular structure of OES cosolvent on cellulose solubility is reflected in another pair of good/modest solvents with very small structural differences: the cyclic urea DMI ($C_5H_{10}N_2O$) combined with [EMIm][OAc] yields an OES with good dissolving power for MC, while the open chain urea TMU ($C_5H_{12}N_2O$) combined with the same IL requires a high IL concentration to effect dissolution.

Designation of the most discriminatory parameters may be based on the tightness of the range within which “good” cosolvents fall. A critical parameter would be expected to yield a tight range, while, for a parameter less important in defining MC dissolution in the OES, a wider spread of values might be noted. Thus, the value “percent of total range” (Table 2) may be used: the smaller the value, the more similar the parameter for the selected solvents and, on a qualitative basis, the more dissimilar the selected solvents from other solvents in the database (this would only be quantitative if all solvents were uniformly distributed over the parameter space). In both databases, the parameters describing the basic nature of the cosolvents (SB and β_1) spread over ranges of 30% or more, suggesting that this is not a key characteristic of a good cosolvent. Conversely, α_1 is limited to zero for all solvents found to form MC dissolving OESs and the similar solvent acidity measure, SA , in the Catalán database is restricted to a narrow range of low values. Given the plethora of studies that have suggested strong interactions between cellulose and acetate ions,^{54–56} this is not surprising, but does perhaps provide an indication of why PC, while appearing to fall well within most ranges, is a poor cosolvent in this context: SA for PC = 0.106 placing this solvent 70% above the upper bound of the range defined for SA using the six selected cosolvents. Finally, polarity (SdP) and polarizability (SP) show intermediate spread for acceptable OES formers, and this is reflected in the DI values in the Laurence database, while ES values appear to be less useful in selecting cosolvents.

To gain a molecular level view of the distribution of IL components relative to dissolved cellulose chains in MC solutions in OESs, RISM calculations were conducted. DMSO was selected as the solvent of choice for this study since it required the least quantity of added IL ($\chi_{IL} = 0.019$) to generate an OES capable of initiating MC dissolution.

Figure 4 shows a map of specific solvation free energy density (SFED) to represent interactions of DMSO, [EMIm]⁺, and acetate with a glucan chain in three different conformations. Solvent–solvent and solute–solvent interactions are fully taken into account in these computations. The results show that DMSO is able to solvate the glucan chain by acting as an H-bond acceptor, concentrating around the hydrogen atoms of the exposed hydroxyl groups. The acetate anions also concentrate around the hydroxyl hydrogen atoms and exhibit a stronger H-bond acceptor character than DMSO. Therefore, acetate and DMSO seem to compete for the same sites on cellulose, with a

Figure 4. Specific solvation free energy density for DMSO (blue), [EMIm]⁺ (green), and acetate (orange) around a glucan chain in three different conformations. The surfaces indicate spatial regions around the solute that concentrate stabilizing solute–solvent interactions (negative solvation free energy values). The isosurfaces were built considering the isovalue of $-18 \text{ kcal}\cdot\text{mol}^{-1}\cdot\text{\AA}^{-3}$ (chosen for best visualization).

preferential solvation by acetate. Alternatively, this could be viewed as DMSO fulfilling a similar H-bond accepting role to acetate, albeit more weakly. We also observe that acetate can concentrate over the glucose pyranose rings and interact with their (weakly positive) hydrogen atoms. In contrast, [EMIm]⁺ is distributed around the oxygen atoms of the exposed hydroxyl groups and glycosidic bonds, with a smaller intensity compared to acetate. Therefore, whereas DMSO, as a dipolar aprotic solvent, is capable of acting only as H-bond acceptor, the ionic liquid can act as both H-bond acceptor and donor, establishing stronger interactions with the glucan chain than does the DMSO cosolvent. The fact that IL–glucan interactions are stronger suggests that small amounts of IL would be able to create the first solvation shell around the cellulose glucan chains, while DMSO would be predominant in the outer solvation shells, thereby interacting less directly with cellulose. In the dynamical equilibrium underlying the solvation phenomenon, with rapid exchange between solvation shell and free species, the ability of DMSO to fulfill some of the H-bond accepting role of acetate, might also explain why the DMSO/[EMIm][OAc] OES is effective even at very low χ_{IL} . This propensity of the IL to solvate cellulose fits well with the shape of the dissolution curves, that is as long as the cellulose glucan chains are largely solvated by IL, further increase of the IL concentration would not lead to substantial increase in cellulose–IL interactions.

CONCLUSIONS

A range of OESs comprising mixtures of [EMIm][OAc] and various solvents (selected from common low to medium molecular weight dipolar aprotic solvents) have been shown to have very similar efficacy for dissolution of cellulose when compared using appropriate measures of mole fractions of solutes and solvents. Thus, 1-MI, DMSO, DMF, DMAc, NMP, DMI, and sulfolane all provide OESs that are effective cellulose solvents at low mole fraction of added ionic liquid. Definition of ranges of solvent parameters derived from the databases of Catalán and Laurence and co-workers and comparison of the relative degree of restriction of the ranges for each confirms others' assertions that cosolvents must not compete with

cellulose with respect to H-bond donation to the IL anion, but the wide range of “allowed” basicity (H-bond acceptor) values, and RISM studies on one system, suggest that hydrogen bond accepting capability may not be deleterious to efficient dissolution of cellulose at low χ_{IL} . Most importantly, given the hazards associated with many of the widely used OES cosolvent components (particularly the reproductive hazards associated with many dipolar aprotic solvents), we have identified an alternative, biobased solvent, γ -valerolactone, as a suitable cosolvent for formation of cellulose dissolving OESs with [EMIm][OAc]. (While γ -butyrolactone also proved to be an excellent cosolvent, significant concerns associated with its use in the preparation of a dangerous drug of abuse, militate against its use.) This offers significant improvements, with regards to safety profiles and sourcing of solvents from renewable sources, for large scale processing of cellulose to form more sustainable materials.

■ ASSOCIATED CONTENT

■ Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: [10.1021/acssuschemeng.6b02020](https://doi.org/10.1021/acssuschemeng.6b02020).

Original weight percent solubility data, annotated Mathematica code for database sorting, fitting coefficients and sorted tables of solvents from the Catalán and Laurence databases (PDF)

■ AUTHOR INFORMATION

Corresponding Author

*E-mail: j.l.scott@bath.ac.uk.

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

The authors thank the UK Engineering and Physical Sciences Research Council (EPSRC) for Ph.D. studentship funding for R.H.W. and M.A.J. via the EPSRC Doctoral Training Centre in Sustainable Chemical Technologies, University of Bath (Grant No. EP/G03768X/1), and the British Council for funding via the Global Innovation Initiative program, which, in particular, facilitated UK/Brazilian collaboration. R.L.S., C.S.P., and M.S.S. thank the Sao Paulo Research Foundation (FAPESP) for financial support (Grants CEPID 2013/08293-7, 2014/10448-1, and 2015/25031-1). RISM calculations were performed at the Center for Computational Engineering and Sciences (CCES) at University of Campinas, Brazil.

■ ABBREVIATIONS

IL: ionic liquid; [EMIm][OAc]: 1-ethyl-3-methylimidazolium acetate; OES: organic electrolyte solution; RISM: reference-interaction site model; H-bond: hydrogen bond

■ REFERENCES

- (1) Klemm, D.; Heublein, B.; Fink, H.-P.; Bohn, A. Cellulose: Fascinating Biopolymer and Sustainable Raw Material. *Angew. Chem., Int. Ed.* **2005**, *44*, 3358–3393.
- (2) Cross, C.; Bevan, E. *Research on Cellulose, 1895–1900*; Longmans, Green and Co.: London, 1901.
- (3) Swatoski, R. P.; Spear, S. K.; Holbrey, J. D.; Rogers, R. D. Dissolution of Cellulose with Ionic Liquids. *J. Am. Chem. Soc.* **2002**, *124*, 4974–4975.
- (4) Zhang, H.; Wu, J.; Zhang, J.; He, J. 1-Allyl-3-methylimidazolium Chloride Room Temperature Ionic Liquid: A New and Powerful Nonderivatizing Solvent for Cellulose. *Macromolecules* **2005**, *38*, 8272–8277.
- (5) Vitz, J.; Erdmenger, T.; Haensch, C.; Schubert, U. S. Extended Dissolution Studies of Cellulose in Imidazolium Based Ionic Liquids. *Green Chem.* **2009**, *11*, 417–424.
- (6) Zavrel, M.; Bross, D.; Funke, M.; Büchs, J.; Spiess, A. C. High-throughput Screening for Ionic Liquids Dissolving (Ligno-)cellulose. *Bioresour. Technol.* **2009**, *100*, 2580–2587.
- (7) Aaltonen, O.; Jauhiainen, O. The Preparation of Lignocellulosic Aerogels from Ionic Liquid Solutions. *Carbohydr. Polym.* **2009**, *75*, 125–129.
- (8) Kosan, B.; Michels, C.; Meister, F. Dissolution and Forming of Cellulose with Ionic Liquids. *Cellulose* **2008**, *15*, 59–66.
- (9) Turner, M. B.; Spear, S. K.; Holbrey, J. D.; Rogers, R. D. Production of Bioactive Cellulose Films Reconstituted from Ionic Liquids. *Biomacromolecules* **2004**, *5*, 1379–1384.
- (10) Zhang, H.; Wang, Z. G.; Zhang, Z. N.; Wu, J.; Zhang, J.; He, J. S. Regenerated-Cellulose/Multiwalled- Carbon-Nanotube Composite Fibers with Enhanced Mechanical Properties Prepared with the Ionic Liquid 1-Allyl-3-methylimidazolium Chloride. *Adv. Mater.* **2007**, *19*, 698–704.
- (11) Wang, S.; Lu, A.; Zhang, L. Recent advances in regenerated cellulose materials. *Prog. Polym. Sci.* **2016**, *53*, 169–206.
- (12) Huddleston, J. G.; Visser, A. E.; Reichert, W. M.; Willauer, H. D.; Broker, G. A.; Rogers, R. D. Characterization and Comparison of Hydrophilic and Hydrophobic Room Temperature Ionic Liquids Incorporating the Imidazolium Cation. *Green Chem.* **2001**, *3*, 156–164.
- (13) Hauru, L. K. J.; Hummel, M.; King, A. W. T.; et al. Role of Solvent Parameters in the Regeneration of Cellulose from Ionic Liquid Solutions. *Biomacromolecules* **2012**, *13*, 2896–2905.
- (14) Abe, M.; Fukaya, Y.; Ohno, H. Fast and facile dissolution of cellulose with tetrabutylphosphonium hydroxide containing 40 wt% water. *Chem. Commun.* **2012**, *48*, 1808–1810.
- (15) Wang, H.; Gurau, G.; Rogers, R. D. Ionic Liquid Processing of Cellulose. *Chem. Soc. Rev.* **2012**, *41*, 1519–1537.
- (16) Rinaldi, R. Instantaneous dissolution of cellulose in organic electrolyte solutions. *Chem. Commun.* **2011**, *47*, 511–513.
- (17) Ohira, K.; Yoshida, K.; Hayase, S.; Itoh, T. Amino Acid Ionic Liquid as an Efficient Cosolvent of Dimethyl Sulfoxide to Realize Cellulose Dissolution at Room Temperature. *Chem. Lett.* **2012**, *41*, 987–989.
- (18) Huo, F.; Liu, Z.; Wang, W. Cosolvent or Antisolvent? A Molecular View of the Interface between Ionic Liquids and Cellulose upon Addition of another Molecular Solvent. *J. Phys. Chem. B* **2013**, *117*, 11780–11792.
- (19) Zhao, Y.; Liu, X.; Wang, J.; Zhang, S. Insight into the Cosolvent Effect of Cellulose Dissolution in Imidazolium-Based Ionic Liquid Systems. *J. Phys. Chem. B* **2013**, *117*, 9042–9049.
- (20) Velioglu, S.; Yao, X.; Devémy, J.; Ahunbay, M. G.; Tantekin-Ersolmaz, S. B.; Dequidt, A.; Costa Gomes, M. F.; Pádua, A. A. H. Thermodynamics of Cellulose Solvation in Water and the Ionic Liquid 1-Butyl-3-methylimidazolium Chloride. *J. Phys. Chem. B* **2014**, *118*, 14860–14869.
- (21) Mostofian, B.; Smith, J. C.; Cheng, X. Simulation of a Cellulose Fiber in Ionic Liquid Suggests a Synergistic Approach to Dissolution. *Cellulose* **2014**, *21*, 983–997.
- (22) Rabideau, B. D.; Agarwal, A.; Ismail, A. E. Observed Mechanism for the Breakup of Small Bundles of Cellulose Ia and Ib in Ionic Liquids from Molecular Dynamics Simulations. *J. Phys. Chem. B* **2013**, *117*, 3469–3479.
- (23) Andanson, J.-M.; Bordes, E.; Devemy, J.; Leroux, F.; Padua, A. A. H.; Costa Gomes, M. F. Understanding the role of cosolvents in the dissolution of cellulose in ionic liquids. *Green Chem.* **2014**, *16*, 2528–2538.
- (24) Xu, A.; Wang, J.; Wang, H. Effects of anionic structure and lithium salts addition on the dissolution of cellulose in 1-butyl-3-methylimidazolium

zolinium-based ionic liquid solvent systems. *Green Chem.* **2010**, *12*, 268–275.

(25) Xu, A. R.; Zhang, Y. J.; Zhao, Y.; Wang, J. J. Cellulose Dissolution at Ambient Temperature: Role of Preferential Solvation of Cations of Ionic Liquids by a Cosolvent. *Carbohydr. Polym.* **2013**, *92*, 540–544.

(26) Rodríguez, A.; Graciani, M. D. M.; Moyá, M. L. Effects of Addition of Polar Organic Solvents on Micellization. *Langmuir* **2008**, *24*, 12785–12792.

(27) Gericke, M.; Liebert, T.; El Seoud, O. A.; Heinze, T. Tailored Media for Homogeneous Cellulose Chemistry: Ionic Liquid/Co-Solvent Mixtures. *Macromol. Mater. Eng.* **2011**, *296*, 483–493.

(28) Wang, Z.; Richter, S. M.; Gates, B. D.; Grieme, T. A. Safety Concerns in a Pharmaceutical Manufacturing Process Using Dimethyl Sulfoxide (DMSO) as a Solvent. *Org. Process Res. Dev.* **2012**, *16*, 1994–2000.

(29) Remsing, R. C.; Swatloski, R. P.; Rogers, R. D.; Moyna, G. Mechanism of cellulose dissolution in the ionic liquid 1-*n*-butyl-3-methylimidazolium chloride: a ^{13}C and $^{35/37}\text{Cl}$ NMR relaxation study on model systems. *Chem. Commun.* **2006**, 1271–1273.

(30) Catalán, J. Towards a Generalized Treatment of the Solvent Effect Based on Four Empirical Scales: Dipolarity (SdP, a New Scale), Polarizability (SP), Acidity (SA), and Basicity (SB) of the Medium. *J. Phys. Chem. B* **2009**, *113*, 5951–5960.

(31) Laurence, C.; Legros, J.; Chantzis, A.; Planchat, A.; Jacquemin, D. A Database of Dispersion-Induction DI, Electrostatic ES, and Hydrogen Bonding α_1 and β_1 Solvent Parameters and Some Applications to the Multiparameter Correlation Analysis of Solvent Effects. *J. Phys. Chem. B* **2015**, *119*, 3174–3184.

(32) Kovalenko, A. Three-dimensional RISM theory for molecular liquids and solid-liquid interfaces. In *Molecular Theory of Solvation*; Hirata, F., Ed.; Kluwer Academic: Dordrecht, The Netherlands, 2003; pp 169–275.

(33) Kovalenko, A. Multiscale Modeling of Solvation in Chemical and Biological Nanosystems and in Nanoporous Materials. *Pure Appl. Chem.* **2013**, *85*, 159–199.

(34) Ratkova, E. L.; Palmer, D. S.; Fedorov, M. V. Solvation Thermodynamics of Organic Molecules by the Molecular Integral Equation Theory: Approaching Chemical Accuracy. *Chem. Rev.* **2015**, *115*, 6312–6356.

(35) Perkyns, J. S.; Pettitt, B. M. A Dielectrically Consistent Interaction Site Theory for Solvent-Electrolyte Mixtures. *Chem. Phys. Lett.* **1992**, *190*, 626–630.

(36) Skaf, M. S. Static dielectric properties of a model for liquid DMSO. *Mol. Phys.* **1997**, *90*, 25–34.

(37) MacKerell, A. D., Jr.; Bashford, D.; Bellott, M.; Dunbrack, R. L., Jr.; Evanseck, J. D.; Field, M. J.; Fischer, S.; Gao, J.; Guo, H.; Ha, S.; Joseph-McCarthy, D.; Kuchnir, L.; Kucsera, K.; Lau, F. T. K.; Mattos, C.; Michnick, S.; Ngo, T.; Nguyen, D. T.; Prodhom, B.; Reiher, W. E., III; Roux, B.; Schlenkrich, M.; Smith, J. C.; Stote, R.; Straub, J.; Watanabe, M.; Wiorkiewicz-Kuczera, J.; Yin, D.; Karplus, M. All-atom Empirical Potential for Molecular Modeling and Dynamics Studies of Proteins. *J. Phys. Chem. B* **1998**, *102*, 3586–3616.

(38) Guvench, O.; Hatcher, E.; Venable, R. M.; Pastor, R. W.; MacKerell, A. D., Jr. CHARMM Additive All-atom Force Field for Glycosidic Linkages Between Hexopyranoses. *J. Chem. Theory Comput.* **2009**, *5*, 2353–2370.

(39) Strader, M. L.; Feller, S. E. A Flexible All-Atom Model of Dimethyl Sulfoxide for Molecular Dynamics Simulations. *J. Phys. Chem. A* **2002**, *106*, 1074–1080.

(40) Kelkar, M. S.; Shi, W.; Maginn, E. J. Determining the Accuracy of Classical Force Fields for Ionic Liquids: Atomistic Simulation of the Thermodynamic and Transport Properties of 1-Ethyl-3-methylimidazolium Ethylsulfate ([emim][EtSO₄]) and its Mixtures with Water. *Ind. Eng. Chem. Res.* **2008**, *47*, 9115–9126.

(41) Silveira, R. L.; Stoyanov, S. R.; Gusarov, S.; Skaf, M. S.; Kovalenko, A. Plant Biomass Recalcitrance: Effect of Hemicellulose Composition on Nanoscale Forces that Control Cell Wall Strength. *J. Am. Chem. Soc.* **2013**, *135*, 19048–19051.

(42) Silveira, R. L.; Stoyanov, S. R.; Gusarov, S.; Skaf, M. S.; Kovalenko, A. Supramolecular Interactions in Secondary Plant Cell Walls: Effect of Lignin Chemical Composition Revealed with the Molecular Theory of Solvation. *J. Phys. Chem. Lett.* **2015**, *6*, 206–211.

(43) Martínez, L.; Andrade, R.; Birgin, E. G.; Martínez, J. M. Packmol: A Package for Building Initial Configurations for Molecular Dynamics Simulations. *J. Comput. Chem.* **2009**, *30*, 2157–2164.

(44) Darden, P.; York, D.; Pedersen, L. Particle mesh Ewald: an $N\log(N)$ method for Ewald sums in large systems. *J. Chem. Phys.* **1993**, *98*, 10089–10092.

(45) Phillips, J. C.; Braun, R.; Wang, W.; Gumbart, J.; Tajkhorshid, E.; Villa, E.; Chipot, C.; Skeel, R. D.; Kalé, L.; Schulten, K. Scalable Molecular Dynamics with NAMD. *J. Comput. Chem.* **2005**, *26*, 1781–1802.

(46) Shlieout, G.; Arnold, K.; Müller, G. Powder and Mechanical Properties of Microcrystalline Cellulose with Different Degrees of Polymerization. *AAPS PharmSciTech* **2002**, *3*, 45–54.

(47) Clough, M. T.; Geyer, K.; Hunt, P. A.; Son, S.; Vagt, U.; Welton, T. Ionic Liquids: Not Always Innocent Solvents for Cellulose. *Green Chem.* **2015**, *17*, 231–243.

(48) Andanson, J.-M.; Pádua, A. A. H.; Costa Gomes, M. F. Thermodynamics of cellulose dissolution in an imidazolium ionic liquid. *Chem. Commun.* **2015**, *51*, 4485–4487.

(49) Olsson, C.; Hedlund, A.; Idström, A.; Westman, G. Effect of Methylimidazole on Cellulose/Ionic Liquid Solutions and Regenerated Material Therefrom. *J. Mater. Sci.* **2014**, *49*, 3423–3433.

(50) UN Convention on Psychotropic Substances, 1971. https://www.unodc.org/pdf/convention_1971_en.pdf (accessed August 04, 2016).

(51) World Health Organisation (WHO) report, Gamma-butyrolactone (GBL) Critical Review Report, Agenda item 4.3, Expert Committee on Drug Dependence, Thirty-sixth Meeting, Geneva, June 16–20, 2014. http://www.who.int/medicines/areas/quality_safety/4_3_Review.pdf (accessed August 04, 2016).

(52) Luterbacher, J. S.; Rand, J. M.; Alonso, D. M.; Han, J.; Youngquist, J. T.; Maravelias, C. T.; Pfleger, B. F.; Dumesic, J. A. Nonenzymatic Sugar Production from Biomass Using Biomass-Derived gamma-Valerolactone. *Science* **2014**, *343*, 277–280.

(53) Horváth, I. T.; Mehdi, H.; Fábos, V.; Boda, L.; Mika, L. T. γ -Valerolactone - a sustainable liquid for energy and carbon-based chemicals. *Green Chem.* **2008**, *10*, 238–242.

(54) Huang, Y. B.; Xin, P. P.; Li, J. X.; Shao, Y. Y.; Huang, C. B.; Pan, H. Room-temperature dissolution and mechanistic investigation of cellulose in a tetra-butylammonium acetate/dimethyl sulfoxide system. *ACS Sustainable Chem. Eng.* **2016**, *4*, 2286–2294.

(55) Andanson, J.-M.; Bordes, E.; Devémy, J.; Leroux, F.; Pádua, A. A. H.; Costa Gomes, M. F. Understanding the role of cosolvents in the dissolution of cellulose in ionic liquids. *Green Chem.* **2014**, *16*, 2528–2538.

(56) Velioglu, S.; Yao, X.; Devémy, J.; Ahunbay, M. G.; Tantekin-Ersolmaz, S. B.; Dequidt, A.; Costa Gomes, M. F.; Pádua, A. A. H. Solvation of a cellulose microfibril in imidazolium acetate ionic liquids: effect of cosolvent. *J. Phys. Chem. B* **2014**, *118*, 14860–14869.