

RESEARCH

Open Access



Natural deep eutectic solvents: cytotoxic profile

Maan Hayyan^{1,2*} , Yves Paul Mbous^{1,3}, Chung Yeng Looi⁵, Won Fen Wong⁴, Adeb Hayyan^{1,6}, Zulhaziman Salleh^{1,3} and Ozair Mohd-Ali⁷

Abstract

The purpose of this study was to investigate the cytotoxic profiles of different ternary natural deep eutectic solvents (NADESs) containing water. For this purpose, five different NADESs were prepared using choline chloride as a salt, alongside five hydrogen bond donors (HBD) namely glucose, fructose, sucrose, glycerol, and malonic acid. Water was added as a tertiary component during the eutectics preparation, except for the malonic acid-based mixture. Coincidentally, the latter was found to be more toxic than any of the water-based NADESs. A trend was observed between the cellular requirements of cancer cells, the viscosity of the NADESs, and their cytotoxicity. This study also highlights the first time application of the conductor-like screening model for real solvent (COSMO-RS) software for the analysis of the cytotoxic mechanism of NADESs. COSMO-RS simulation of the interactions between NADESs and cellular membranes' phospholipids suggested that NADESs strongly interacted with cell surfaces and that their accumulation and aggregation possibly defined their cytotoxicity. This reinforced the idea that careful selection of NADESs components is necessary, as it becomes evident that organic acids as HBD highly contribute to the increasing toxicity of these neoteric mixtures. Nevertheless, NADESs in general seem to possess relatively less acute toxicity profiles than their DESs parents. This opens the door for future large scale utilization of these mixtures.

Keywords: Natural deep eutectic solvents, Green solvents, Ionic liquids, Choline chloride, Cytotoxicity, Cancer cell line, COSMO-RS

Background

The use of volatile organic compounds (VOCs) has left in its wake countless considerations, most of which associated with safety and toxicity issues (Bushnell et al. 2007). Therefore, for various chemical and biological industries, one of the most pressing concern remains the development of 'greener', lower cost, and more efficient solvents. The discovery of ionic liquids (ILs) seemed to provide the solution to this predicament. ILs being low-melting organic salts composed of ionic species, which are often found in liquid state at temperatures below 100 °C (Ru and Konig 2012). ILs are characterized by a number of attractive properties such as high thermal stability, non-flammability, high solvability, chemical stability, low

volatility and tunability (Sowmiah et al. 2009). Notwithstanding their impressive contributions in various processes such as biotransformations (Domínguez de María and Maugeri 2011), biodiesel production (Ullah et al. 2015), extraction processes (Pereira et al. 2015), active pharmaceutical ingredients (Ferraz et al. 2011), and biomass treatment (da Costa Lopes et al. 2013); the use of ILs has often been marred with issues pertaining to high cost of synthesis, purification requirements, and toxicity (Gorke et al. 2010). The ambiguity surrounding their use has led to the emergence of an alternative class of solvents called deep eutectic solvents (DESs). As eutectics, DESs exhibit freezing points lower than those of their chief components (salt and HBD) (Smith et al. 2014). This depression in temperature is the result of the charge delocalization occurring via hydrogen bonding between the anion of the salt and the HBD (Ru and Konig 2012). DESs have been introduced as viable replacements of ILs because, on top of possessing similar physicochemical

*Correspondence: maan_hayyan@yahoo.com

¹ University of Malaya Centre for Ionic Liquids (UMCIL), University of Malaya, 50603 Kuala Lumpur, Malaysia
Full list of author information is available at the end of the article

aspects, they offer several other advantages, such as the low cost of their starting materials, ease of preparation, and no waste generation (Tang and Row 2013). As a result, they have been used in a wide number of applications such as extraction processes (Qi et al. 2015), biotransformations (Wu et al. 2014), nanoparticles assembly (Wagle et al. 2014), preservation of biomolecules (Dai et al. 2015), upstream and downstream biodiesel processing (Hayyan et al. 2010, 2013a, 2014), electrodeposition (Ru et al. 2015), and organic synthesis (Zhang et al. 2012).

The most encountered DESs are based on ChCl, which revolves around choline. Choline is a known component of Vitamin B, and plays important metabolic functions (Florindo et al. 2014). The qualifications of choline as a safe ingredient led to the assumption that DESs possess negligible toxicity/cytotoxicity profiles.

However, the leading cytotoxic assessments of DESs showed that they can be lethal to both terrestrial and marine organisms (bacteria and brine shrimp) (Hayyan et al. 2013b, c). Ammonium-based DESs, namely: [ChCl]-[Glycerol], [ChCl]-[Ethylene glycol], [ChCl]-[Triethylene glycol], [ChCl]-[Urea]; and methyltriphenyl phosphonium bromide (MTPB)-based DESs such [MTPB]-[Glycerol], [MTPB]-[Ethylene glycol], and [MTPB]-[Triethylene glycol], were used during these investigations. Both ammonium and phosphonium-based DESs were found toxic to brine shrimp, but only phosphonium-based DESs exhibited bacterial toxicity. The cytotoxicity of both ammonium and phosphonium-based DESs was higher than those of their individual components (Hayyan et al. 2013b, c). Therefore, the authors concluded that although ChCl and MTPB salts are not devoid of toxicity; their association with a HBD during DESs' preparation increases the eutectics' cytotoxicity considerably. This conclusion has been recurrent throughout several cytotoxic assessments of DESs (Hayyan et al. 2015; Radošević et al. 2015; Wen et al. 2015).

The HBD has proven to be of significant importance with regards to DESs' cytotoxic profiles. Recently, Radošević et al. (2015) examined the cytotoxic profile of [ChCl]-[Glucose], [ChCl]-[Glycerol], and [ChCl]-[Oxalic acid] on channel catfish ovary (CCO) fish cell line and the human breast adenocarcinoma cell line (MCF-7). Their results showed that the [ChCl]-[Oxalic acid] DES exhibited a significantly higher toxicity ($EC_{50} < 5$ mM) compared to the remaining ChCl-based DESs ($EC_{50} > 10$ mM). These results reinforced the importance of a careful selection of the HBD prior DESs synthesis. In yet another study, among four ChCl-based DESs namely, [ChCl]-[Urea], [ChCl]-[Acetamide], [ChCl]-[Ethylene glycol], and [ChCl]-[Glycerol]; the [ChCl]-[Ethylene

glycol] DES was shown to have the highest toxicity (Wen et al. 2015).

The cytotoxic mechanism of DESs outlines an increased membrane porosity due to the continuous DESs' induced damage of the plasma membrane (Hayyan et al. 2015). Accordingly, upon penetration, DESs' species effectively contribute to the increase of reactive oxygen species (ROS) concentrations, hereby subjecting the cell to increasing oxidative stress. The end-point of this cascade involves the complete destruction of the cell through apoptosis (Hayyan et al. 2015).

The key to producing less toxic DESs may reside in the use of materials of natural origin. Recently, Choi et al. (2011) revealed that a number of primary metabolic substances (e.g. sugars, amino acids, choline, and some organic acids) form intracellular eutectic mixtures to assist plants during specific developmental stages (germination, cryopreservation). These eutectics provide cells with a third type of solvent/media, completely different from lipids and water. The presence of these eutectic mixtures—termed natural deep eutectic mixtures (NADESs)—intracellularly, entails their cellular tolerance and presume safer cytotoxic profiles. Recent studies have provided a list of the composition of these mixtures as well as their molar ratios (Choi et al. 2011; Dai et al. 2013).

One of the fundamental precepts of this class dictates that if cellular media produce NADESs, the propensity for cytotoxicity must be minimal. Paiva et al. (2014) briefly investigated the cytotoxic profile of several NADESs namely, [ChCl]-[Glucose], [ChCl]-[Citric acid], [ChCl]-[sucrose], [ChCl]-[Tartaric acid], [ChCl]-[Xylose], [Citric acid]-[Glucose], [Citric acid]-[Sucrose], [Glucose]-[Tartaric acid] at various molar ratios. Using fibroblast-like cells, the authors assessed cellular viability following NADESs' treatment. The results pointed once again to the role of the HBD (organic acids) as major enhancer of cytotoxicity, because the most toxic NADESs were [Glucose]-[Tartaric acid], [ChCl]-[Tartaric acid], [ChCl]-[Citric acid], and [Citric acid]-[Glucose]. Understanding the various interactions or forces resulting from the association of NADESs chief elements can provide further elucidation of their resulting cytotoxic profiles.

A recent study has shown that NADESs physical properties can be tailored by adding water as a tertiary component. The authors showed that the strong hydrogen interactions within NADESs (accounting for their high viscosities) could be reduced upon addition of water (<50 % v/v). In fact, the resulting viscosities were found to be as low as those of water and other common VOCs (Dai et al. 2015). Consequently, water-based NADESs may represent yet another alternative to DESs of high viscosities, poor conductivities and higher toxicities.

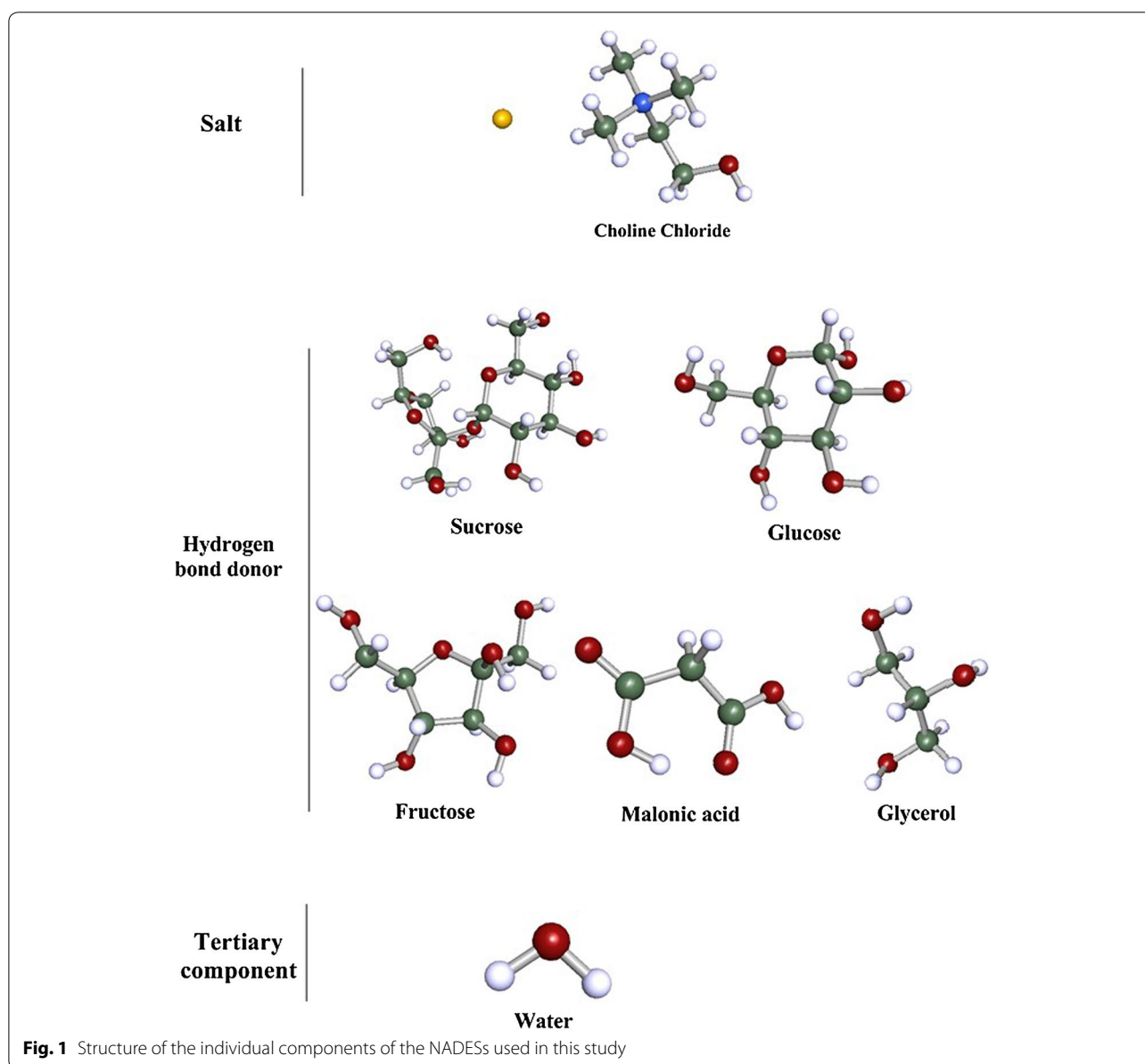
Hence, one of the objectives of this study was to investigate the effect of water on the cytotoxic profiles of NADESs by exploring the impact ternary NADES systems (containing water) have on several cancer cell lines. The understudied eutectics were ChCl-based NADESs prepared using five HBDs, namely, fructose, glucose, sucrose, glycerol, and malonic acid. Alternatively, we aimed to compare the effect of biomaterials against that of organic acids on mammalian cells. Lastly, we sought to model the interactions between NADESs and cellular membranes using COSMO-RS computational methodology in order to understand NADESs cytotoxic mechanism.

Methods

Chemicals and materials

The chemicals used for NADESs preparation were purchased from both Merck (sucrose, fructose and glucose) and Sigma-Aldrich (ChCl and malonic acid), Fig. 1. The purity of these chemicals was higher than 99 %.

The human cervical cancer cell line (HelaS3), the human ovarian cancer cell line (CaOV3), and the mouse skin cancer cell line (B16F10) were purchased from the American Type Culture Collection (ATCC, Manassas, VA). The human breast cancer cell line (MCF-7) was obtained from Cell Lines Service (300273; Eppelheim, Germany).



Both the Dulbecco's Modified Eagle Medium (DMEM) and the Roswell Park Memorial Institute medium (RPMI 1640) were obtained from Life Technologies, Inc., Rockville, MD. Fetal bovine serum (FBS) was supplied by Sigma-Aldrich.

Preparation of NADESs

Table 1 illustrates the composition, molar ratios and symbols of the NADES used throughout this study. The preparation method is similar to those previously described in the literature (Hayyan et al. 2013c).

Cell culture

HelaS3, CaOV3, and B16F10 were grown in DMEM supplemented with 10 % FBS, 1 % penicillin and streptomycin. MCF-7 were grown in RPMI supplemented with 10 % FBS, 1 % penicillin and streptomycin. The cells were kept in culture flasks inside an incubator providing a humidified atmosphere of 37 °C, with 5 % CO₂. The cells were grown to a necessary confluence of 70–80 %, necessary for the 3-(4, 5-dimethylthiazolyl-2)-2, 5-diphenyltetrazolium bromide (MTT) viability assay.

MTT viability assay

The MTT cell viability assay was performed as previously described (Hayyan et al. 2015). The IC₅₀ values were obtained from an average of at least 3 independent experiments. The standard error of the mean (SEM) derived from the repeated experiments were used to derive the variations from the average IC₅₀ values. The statistical analysis was performed using Graph Pad Prism 5 software. Statistical significance was defined when $P < 0.05$.

Computational methodology for COSMO-RS

Molecular geometry optimization

The geometry optimization of all species involved in this study was performed using the Turbomole program package. Using this program, the basic structure of the target molecule was drawn first. After which,

geometry optimization was performed at the Hartree–Fock level and 6-31G* basis set. The generation of cosmo file was then conducted through a single-point calculation by using DFT with Becke–Perdew and the Triple- ζ Zeta Valence Potential (TZVP) basis set. Finally, the *cosmo files* were exported to the COSMOthermX program with parameterization BP_TZVP_C30_1301.ctd.

DES representation in COSMOtherm-X

Since a single DES is composed of more than one molecule, employing its representation method in the COSMOtherm-X program is crucial. In this study, the electroneutral approach was adopted, whereby the DES was represented in COSMO-RS according to the mole composition of their constituents shown in Table 1 [the salt cation, salt anion, and hydrogen bond donor (HBD)]. Membrane phospholipids were designed according to the same principle; that is using the most basic composition of their constituents (Table 2).

Results and discussion

The cytotoxicity of the five understudied NADESs was assessed on various human and mice cancer cell lines, namely, HelaS3, CaOV3, MCF-7, and B16F10. Table 3 illustrates the IC₅₀ values obtained. The results indicate the following decreasing order of toxicity for HelaS3, MCF-7, and B16F10 cell lines: NADES₅ > NADES₃ > NADES₁ > NADES₂ > NADES₄. In CaOV3 case, NADES₂ was more toxic than NADES₁ resulting in a slightly different trend: NADES₅ > NADES₃ > NADES₂ > NADES₁ > NADES₄. However, if the SEM of the IC₅₀ values are included, the resulting IC₅₀ intervals of both NADES₁ (198.5–213.5 mM) and NADES₂ (185.5–200.5 mM) overlap, as the end values are close to one another.

Overall, we noticed a trend between NADESs' cytotoxicity and various factors namely the cellular requirements of cancer cells, the physical properties of NADESs (especially viscosity); the addition of water; and the nature of NADESs' raw materials as well as their interactions with the different functional groups present on the cell surface.

The merits of most DESs stem from the qualifications of ChCl; with specific referral to the metabolism and function of choline in mammalian cells. Choline is the preferred cellular raw material used for the synthesis of cellular membranes phospholipids, namely phosphatidylcholine, and sphingomyelin (Lodish et al. 2000; Plagemann 1971). Consequently, ChCl has been classified as a salt of relatively safe profile (although high intake is associated with adverse conditions). However, the DESs cytotoxic profiles obtained thus far do not share the negligible cytotoxic labeling of ChCl. This has

Table 1 Compositions, symbols, and molar ratios of the NADESs used in this study

Salt	HBD	Add-on	Molar ratio	Appearance	Symbol
ChCl	Fructose	Water	5:2:5	Moderately viscous liquid	NADES ₁
ChCl	Glucose	Water	5:2:5	Moderately viscous liquid	NADES ₂
ChCl	Sucrose	Water	4:1:4	Moderately viscous liquid	NADES ₃
ChCl	Glycerol	Water	1:2:1	Lightly viscous liquid	NADES ₄
ChCl	Malonic acid	–	1:1	Moderately viscous liquid	NADES ₅

Table 2 Composition of cellular membrane and ratio (Harwood and Weselake 2015)

Cell membrane elements	Composition and ratio			
	Fatty acid	Alcohol	Metabolites	Functional group
Phosphatidylcholine	Palmitic acid (2)	Glycerol (1)	Choline (1) Phosphate (1)	Phosphate (1)
Phosphatidylethanolamine	Linoleic acid (1); palmitic acid (1)	Glycerol (1)	Ethanolamine (1) Phosphate (1)	Phosphate (1)
Phosphatidylserine	Stearic acid (1); ceronic acid (1)	Glycerol (1)	Serine (1) Phosphate (1)	Phosphate (1)
Sphingomyelin	Oleic acid (1)	–	Choline (1) Phosphate (1)	Phosphate (1)
Glycolipids	Oleic acid (1)	–	Glucose (1) Sphingosine (1)	Phosphate (1)

Table 3 IC₅₀ of the studied NADESs on various cell lines

Solvent	IC ₅₀ (mM)			
	Hela S3	CaOV3	MCF-7	B16F10
NADES ₁	177 ± 7.3	206 ± 7.5	127 ± 9.22	195 ± 7.7
NADES ₂	182 ± 7.6	193 ± 7.5	186 ± 7.9	211 ± 8
NADES ₃	166 ± 5.8	154 ± 5.6	150 ± 5.5	136 ± 5.7
NADES ₄	427 ± 11	483 ± 11	457 ± 11	340 ± 10.3
NADES ₅	20 ± 8.4	15 ± 8.2	35 ± 8.3	35 ± 8.8

prompted the examination of the role of the HBD in these profiles.

From a cellular perspective, fructose, glucose, sucrose (50 % glucose and 50 % fructose), and glycerol are essential carbohydrates whose metabolism provide energy required for various cellular functions. Upon adsorption, fructose and glucose undergo glycolysis if energy is needed, or are stored as glycogen. The glycolytic pathway for these molecules leads to either the pentose phosphate pathway (for nucleic acid synthesis), the mitochondrial tricarboxylic acid pathway (for energy production), or de novo lipogenesis (for fatty acids synthesis). Cancer cells, especially, require more energy than normal cells, given their abnormal and exponential growth features. Therefore, they use a higher amount of energy or energy sources (glucose and fructose) for metastasis, growth, invasion and migration purposes (Port et al. 2012; Santos and Schulze 2012).

Likewise, glycerol is the precursor of triglycerides and phospholipids. It is activated by a phosphorylation reaction and forms glycerol-3-phosphate (G-3-P), which is then involved in the carbohydrate and lipid metabolism. Alternatively, glycerol also functions as a shuttle of electrons from the cytosol to the mitochondria by regenerating NAD⁺ from NADH (Laforenza et al. 2016). In both normal and cancer cell lines, glycerol can be used for

gluconeogenesis, although the main metabolite used for that purpose is different; probably glycogen. Nevertheless, there is evidence that in cancer cells, a higher than normal plasma concentration of glycerol (comparable in this case to NADES₄ treatment) contributes to increased glycerol turnover for gluconeogenesis and lipogenesis (Liu et al. 1995; Lundholm et al. 1982). Judging from these facts, a higher cellular tolerance of these carbohydrates-based eutectics is expected, and this is reflected by the IC₅₀ values recorded for NADES₁, NADES₂, NADES₃ and NADES₄.

In contrast, NADES₅ which boasts of organic acid as raw material, is the most lethal mixture. Dai et al. (2013) listed NADES₅ as eutectic used by plants for developmental or metabolic purposes. Although this is valid for certain plants tissues—where malonic acid accounts for as much as 4 % of the dry weight and up to 50 % of the total acid content and may be actively used during nitrogen assisted symbiosis or abiotic stress as a defense chemical; the scenario might be slightly different for mammalian cells (Kim 2002). Indeed, in mammalian systems, malonic acid is known to stall the Krebs cycle by inhibiting succinic dehydrogenase (mitochondria complex II); a crucial enzyme for the citric acid cycle and the electron transport chain (Hosoya and Kawada 1958). Consequently, this paralyzes ATP synthesis. Moreover, malonate is known to disrupt glycogenesis, lipid synthesis and carbon dioxide production during glycolysis (Hosoya et al. 1960). It comes as no surprise that calls have been made for malonate to be used as an anticancer agent. As a matter of fact, Fernandez-Gomez et al. (2005) showed that malonate causes SH-SY5Y neuroblastoma mitochondrial failure by inducing a rapid build-up of ROS, which overwhelms mitochondrial antioxidant capacity, ultimately leading to cellular apoptosis.

This shows that with regards to the HBD, the inclusion of organic acids seem to increase the overall toxicity

of NADESs. This is consistent with the other cytotoxic reports on DESs/NADESs (Paiva et al. 2014; Radošević et al. 2015; Zhao et al. 2015).

Zhao et al. (2015) observed that NADESs with organic acids as HBDs had a low pH (<6.5); when the optimal growth range for mammalian cells is 7.0–7.4. This change in environmental conditions is partially responsible for the high toxicity of NADES₅.

DESs investigations led to a similar observation. For instance, Radošević et al. (2015) observed the formation of intracellular calcium oxalate crystals following [ChCl]-[Oxalic acid] DES treatment of CCO and MCF-7 cell lines. Another perspective on organic acids as HBDs was shown by Paiva et al. (2014). The authors examined NADESs toxicity towards fibroblasts like-cells (L929) and reported that the most lethal solvents also had organic acids as HBDs (i.e. tartaric acid and citric acid). However, it must be noted that the solvents with the highest viability, also had organic acids as ingredients, although the remaining constituent was another HBD (sucrose) and not a salt (ChCl) (Paiva et al. 2014). It might be that the devastating effect of organic acids in NADESs is better countered with the use of biomaterials (e.g. sugars).

The arguments above do not presume to provide a complete understanding of the reasons behind the variation in IC₅₀ values; but serve to highlight that safer NADESs can be obtained by using biomaterials of cellular necessity. Of course, the interactions of these mixtures and their aggregation on cellular membranes as well as the neoteric properties of NADESs, remain aspects to be investigated. Meanwhile, physical properties of NADESs can also be used to better appreciate the obtained cytotoxic values.

Table 4 lists the known viscosities values at 30 °C of the understudied NADESs. The values for NADES₁, NADES₂, and NADES₅ were obtained from a recent study by Zhao et al. (2015). The viscosities of NADES₃ and NADES₄ were measured independently during this study. Sorting out viscosities in a decreasing order (NADES₃ > NADES₅ > NADES₂ > NADES₁ > NADES₄) reveals that they form a series almost similar to the cytotoxicity trend above.

According to Table 4, NADES₅ and NADES₃ possess the highest viscosities at 30 °C (respectively 616 and 853.3 mPa s). It is no surprise that they also possess the lowest IC₅₀ values on average across all examined cells, as high viscosity is often associated with increased lethality. Despite being less viscous than NADES₃, NADES₅ was identified as the most toxic material tested, with an IC₅₀ interval (15 ≤ IC₅₀ ≤ 35 mM) at least three times lower than NADES₃' interval (136 ≤ IC₅₀ ≤ 166 mM).

In a separate study, upon testing numerous DESs and NADESs (including similar NADES₁, NADES₂, and

Table 4 Viscosities of the understudied NADESs

NADES	Molar ratio	Viscosity (mPa s)	References
NADES ₁	5:2:5	584	Zhao et al. (2015)
NADES ₂	5:2:5	598	Zhao et al. (2015)
NADES ₃	4:1:4	853.3	–
NADES ₄	1:2:1	36	–
NADES ₅	1:1	616	Zhao et al. (2015)

NADES₅ used in this study) on several bacteria species (i.e. *Escherichia coli*, *Staphylococcus aureus*, *Salmonella enteritidis*, *Listeria monocytogenes*), Zhao et al. (2015) also identified NADES₅ as one of the most toxic mixture. In contrast, NADES₁ and NADES₂ were found to be non-inhibitory to all studied bacteria. Moreover, the most toxic DESs and NADESs ([ChCl]-[Toluenesulfonic acid], [ChCl]-[Oxalic acid], [ChCl]-[Levulinic acid], [ChCl]-[Malonic acid], [ChCl]-[Malic acid], [ChCl]-[Citric acid], [ChCl]-[Tartaric acid] all included organic acids as HBDs (Zhao et al. 2015). Just as in our study, where the malonic acid based NADES was the most toxic. This stipulates that despite resulting in high viscosities (compared to water and VOCs), NADESs composed of sugars are relatively less dangerous to biological machinery than those composed of organic acids.

On the other hand, NADES₄ exhibited the lowest viscosity at 30 °C (36 mPa s) as well as the highest cytotoxicity values. With reference to Table 4, NADES₂ is slightly more viscous than NADES₁. Their cytotoxicity values are roughly similar if the SEM is taken into account. Viscosity or micro-viscosity (in cellular terms) is an important property to consider in intracellular activities. Not only does it affect diffusion within biological systems, but it is also involved during processes such as protein–protein interactions, transportation of small solutes and macromolecules, and signal transduction in living cells. The local micro-viscosity in cells ranges from 1 to 400 mPa s (Liu et al. 2014). The highest viscosity values (≥200 mPa s) are usually associated with the microviscosity in the hydrophobic domains of living cells (lipid bilayers of cell surfaces); whereas, values between 1 and 3 mPa s are attributed to the aqueous phases of the cellular cytoplasm (Juneidi et al. 2015; Kuimova et al. 2009). A variation or a disturbance of these homeostatic values leads to the onset of various diseases (atherosclerosis, diabetes) as well as cell death (Deliconstantinos et al. 1995; Nadiv et al. 1994). The understudied NADESs possess viscosities higher than 500 mPa s; with the exception of NADES₄ (36 mPa s), which happens to be the least toxic mixture tested (with IC₅₀ values 1.6–32 times lower than the other NADESs). Hence, it is not difficult to perceive the substantial influence that these highly viscous

materials can have on cells. Just like DESs perforate cellular membranes (Hayyan et al. 2015), NADESs can probably enhance cellular membrane permeability. As such, the introduction of such viscous substances in cellular medium can result in a major variation of cytoplasmic microviscosity, and eventually lead to cell death.

Based on the available knowledge of DESs, the high viscosities of NADESs originate from the rigidity of their supramolecular complexes reposing on a strong hydrogen bond network. It entails that the disruption of this network will affect the viscosity of NADESs. In fact, Dai et al. (2015) recently provided evidence of the progressive rupture of this hydrogen bond network upon addition of water. The authors also showed that the supramolecular complexes of NADESs remain intact if the volume of added water is less than 50 %. Pass this threshold, the resulting mixture consists merely of dissociated NADES ingredients. This is a consequence of the complete rupture of hydrogen bonds stabilizing NADESs. The fact that the entire NADES structure repose on hydrogen bonds means that their progressive breakdown simultaneously induces a change in their physicochemical properties. Accordingly, Dai et al. (2015) reported a decrease in viscosity from 397 to 7.2 mPa s, following the addition of 25 % of water to [ChCl]-[Glucose]-[Water] (in this case NADES₂).

This argument is further justified by the fact that we have independently recorded a value of 36 mPa s for NADES₄ ([ChCl]-[Glycerol]-[Water]) viscosity; whereas Zhao et al. (2015) recorded a value of 177 mPa s for the [ChCl]-[Glycerol] DES (both at 30 °C). The effects of water can also be acknowledged through the observation that of all the understudied NADESs, the most toxic and most viscous was the one prepared without the use of water (NADES₅). Moreover, Dai et al. (2015) stated that the water activity of NADESs increases with increasing water content (or after water addition). Consequently, the polarity of the eutectics after addition of water may mimic that of water itself. This may influence the interactions of these solvents with cell surfaces.

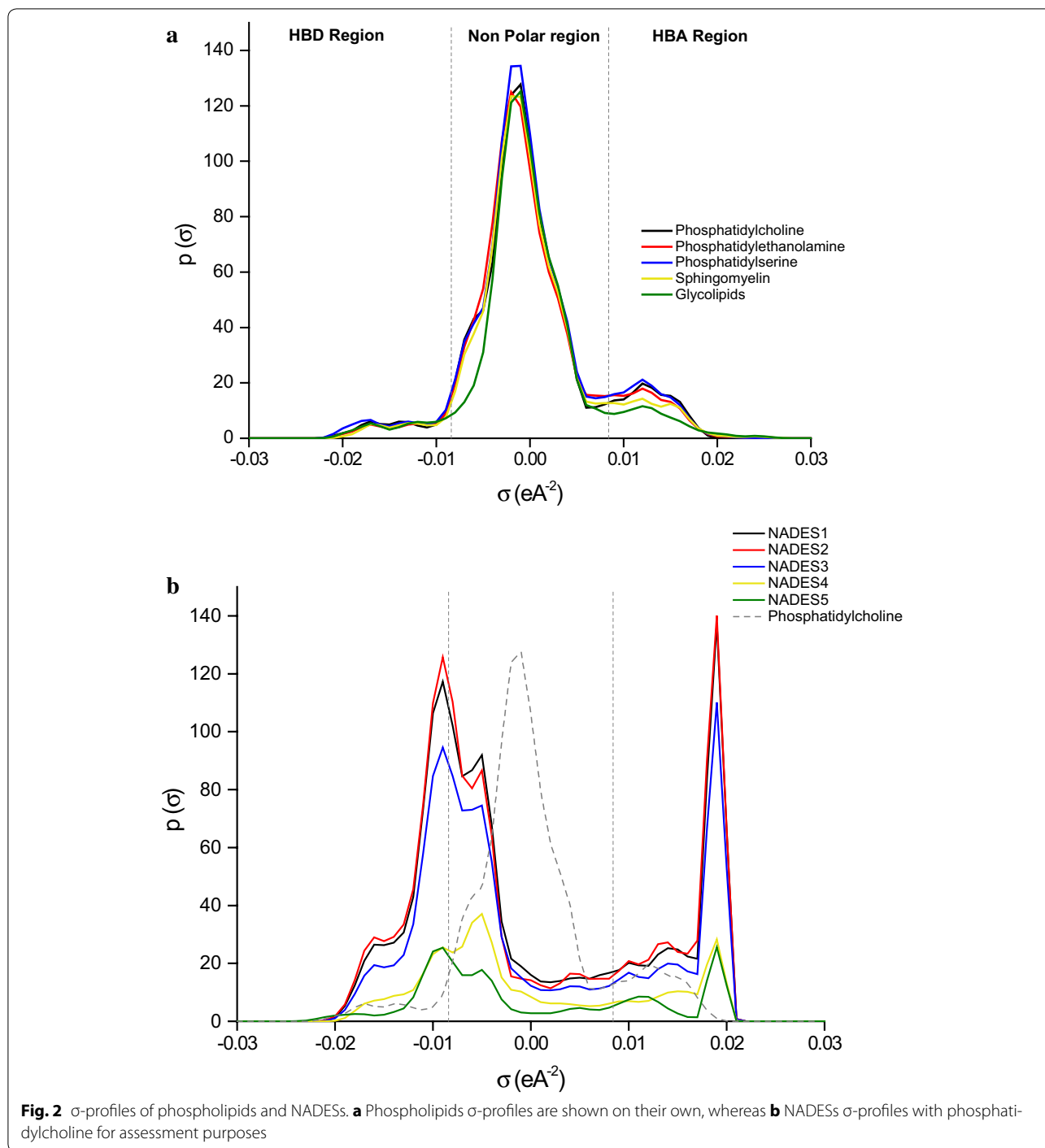
The importance of these interactions must be dully underscored, as DESs have shown that they promote cellular failure through an increase in membrane porosity (Hayyan et al. 2015). In order to have an idea of what unfolds upon NADESs treatment on cell surface, we used a computational methodology using the COSMO-RS software. COSMO-RS is a very useful and fast tool for the prediction of thermophysical and chemical properties of fluid mixtures (Klamt 2005). It is a model that combines an electrostatic theory of locally interacting molecular surface descriptors with statistical thermodynamics. Although mostly used to predict the thermodynamic properties of a mixture without prior experimental

data; it can also be applied to life sciences and molecular studies. Examples include the prediction of drug's partition coefficients and the computation of proteins pKa (Andersson et al. 2013; Buggert et al. 2009). Two major steps are involved in the COSMO-RS prediction process. The first step involves the creation of virtual conductor surroundings for the molecule by using the continuum solvation model. After performing the quantum chemical calculation through the density functional theory, a screening charge density known as sigma (σ) forms on the nearby conductor. The distribution of the screening charge density on the surface of the molecule is then converted into a function of surface composition, known as the σ -profile. The second step applies statistical thermodynamic principles to compute the molecular energy due to the electrostatic misfit, hydrogen bond, and Van der Waals interactions (Klamt and Eckert 2000). COSMO-RS can also be used to study the possible thermodynamic behaviour of an individual component in a mixture and its affinity or interactions with the other components through the σ -profile and σ -potential, respectively. The σ -profile describes the molecule polarity properties. Each peak in the σ -profile plot for a molecule corresponds to its constituent atoms depending on their screening charge densities. The negative partial charges of atoms cause positive screening charge densities, and vice versa.

Using the elements in Table 2, we modeled each of the listed phospholipids and examined their interactions with the NADESs. The sigma profiles of both phospholipids and NADESs are shown in Fig. 2. For comparison purposes, we only used phosphatidylcholine, as it is the most common membrane phospholipid, and also because, the phospholipids' σ -profiles and potentials are almost identical.

In the σ -profile, when the screening charge density is lower than $-0.0084 \text{ e}\text{\AA}^{-2}$ or exceeds $+0.0084 \text{ e}\text{\AA}^{-2}$; the molecule is considered sufficiently polar to induce hydrogen bonding.

Figure 2 is divided into three quadrants with corresponding σ values; the HBD region ($\sigma < -0.0084 \text{ e}\text{\AA}^{-2}$), the nonpolar region ($-0.0084 \leq \sigma \leq 0.0084 \text{ e}\text{\AA}^{-2}$) and the HBA region ($\sigma > 0.0084 \text{ e}\text{\AA}^{-2}$). Negative values represent positive polarities and vice versa. Hence, the elements in the HBD region of a molecule interact or attract elements in the HBA region (from another molecule) since they are of opposite polarities. Looking at Fig. 2a, the bulks of the peaks reside in the nonpolar region. This is most likely due to the long hydrophobic fatty acid chains making up these phospholipids and included during our modeling. Compared to NADESs, phosphatidylcholine σ -profile (similar to all other phospholipids) breadth is narrower; this is usually indicative of a less polar character (Fig. 2b) (Mulyono et al. 2014).



The peaks located between $+0.01$ and $+0.02 e\text{\AA}^{-2}$ in Fig. 2a, represent the negative charge of the O atoms present in the hydroxyl groups of the phospholipids ingredients (phosphate, glycerol, fatty acid). These O atoms may interact to form hydrogen bonds with the H atoms in NADESs that produce the peaks between -0.02 and $-0.01 e\text{\AA}^{-2}$. These peaks belong to the hydroxyl groups

of the sugars, polyols, or acids ingredients of NADESs and the H atom of ChCl. Likewise, the broad HBD region of the phospholipids between -0.01 and $-0.02 e\text{\AA}^{-2}$ represent H atoms in the glycol groups of glycerol or in the functional group of the acids. These may interact with the elements in the HBA region (which mainly comes from Cl atoms of ChCl at $0.02 e\text{\AA}^{-2}$) in Fig. 2b to form

hydrogen bonds;. The importance of these interactions is perhaps best reflected in the σ -potentials of Fig. 3. The σ -potential represents the interaction behavior and affinities between molecules in a system. On the σ -potential plot, a more negative value of μ (σ) indicates higher affinity, and vice versa. Figure 3a shows the σ -potentials of modelled phospholipids. The phospholipids show clear and strong affinities for HBD on the left side, and HBA on the right side, given their high outer negative ranges. Consequently, they will be more attractive for HBDs and HBAs of other molecules. Their nonpolar surfaces possess slightly negative values, which promotes average to

low interactions with other molecules nonpolar surfaces. NADESs, in contrast, vary slightly in their σ -potentials. NADES₁, NADES₂, and NADES₃, all possess very strong affinities for HBDs; relatively weaker affinities for HBAs, and low affinities for nonpolar surfaces. When compared to phosphatidylcholine (Fig. 3b), NADES₁, NADES₂, and NADES₃ affinities for HBD, and nonpolar regions is significantly higher; although relatively similar for HBA. It entails that these three NADESs can interact strongly with HBDs, HBAs and nonpolar surfaces of phospholipids than NADES₄ and NADES₅. These interactions may correlate with solvent accumulation and aggregation on

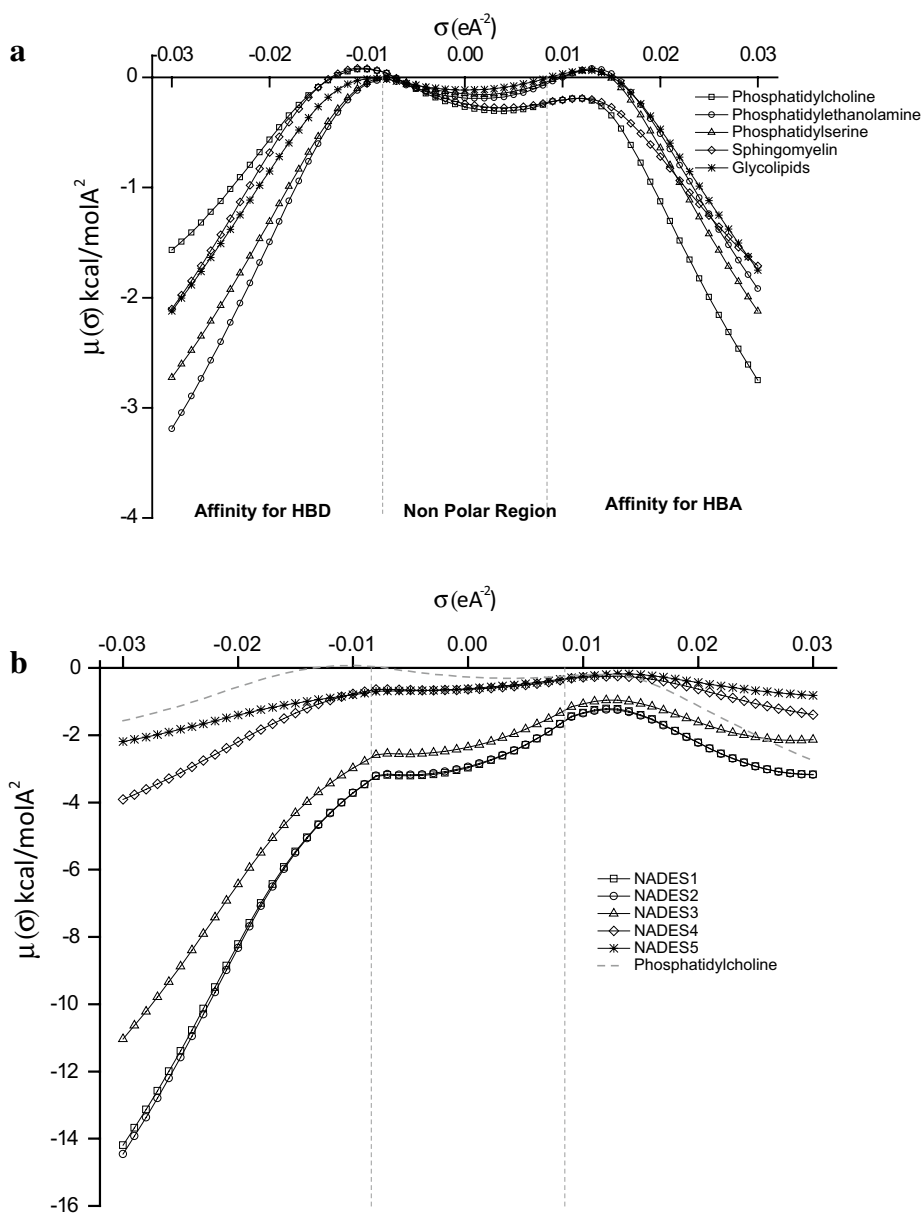


Fig. 3 σ -potentials of phospholipids and NADESs. **a** Phospholipids σ -potentials, **b** NADESs σ -profiles with phosphatidylcholine

the cell surface, which ultimately leads to cellular demise through reduced growth. An example of such critical interactions between groups of opposites polarities and affinities was shown by Cornmell et al. (2008). The authors emphasized that the interactions taking place between aqueous quaternary ammonium salts cations (such as cholinium cations) and the negatively charged groups present on cell surfaces may lead to the penetration of the latter in the cytoplasm. The consequences range from the loss of membrane integrity to a subsequent demise of the cell, through an increased permeability of the cell membrane to exogenous species (Cornmell et al. 2008).

NADES₄ and NADES₅ affinities are almost similar to phospholipids (phosphatidylcholine). These mild affinities of NADES₄ and NADES₅ for HBDs and HBAs of phospholipids suggest that perhaps their cytotoxic mechanism is not entirely focused on cellular aggregation, but rather depends on the resulting reactions engendered by the cellular adsorption. That is, Krebs cycle stalling and acidosis by NADES₅, and cellular poisoning by NADES₄ at a threshold concentration. It is interesting though to note that the solvents with the overall lesser affinities are both the most toxic and least toxic understudied eutectics.

Of course, this model is not an exact replica of what is found in membranes, especially in terms of ratio and functional groups occurrence. As such, the higher ratio of ChCl, or water and sugars in NADES₁, NADES₂, and NADES₃, and the hypothetical ratio of cell membrane elements, may explain the resulting fluctuating affinities. Phospholipids elements especially, consist of a set ratio of functional groups on the cell surface (carboxyl, phosphate and amino groups). The ratio of these functional groups dictates the entry and the rate of passage of extracellular materials (such as NADESs' species) in the intracellular medium, as their proportions differ according to cell type. These proportions regulate the diffusion of NADESs, and indirectly affect their effect on the cellular machinery.

The propensity of NADESs/DESs species to permeate through cellular membranes was suggested to obey a principle of colloidal biology, which is based on the Hofmeister phenomenon (Vlachy et al. 2009). An elucidation of the specifics of the principle of affinities between chaotropic and kosmotropic DESs/NADES species and cell surface groups would provide a strong tool for the prediction of the toxicity of these mixtures.

Conclusion

NADESs show similar physical characteristics to DESs. They exhibit high viscosities, poor conductivities and malleable densities at room temperature. These

characteristics are determined by the strong hydrogen networks holding together their supramolecular structures. Loosening this network brings about ideal conditions for the industrial use of these solvents. Changes in temperatures alter this network but so does the inclusion of water as a tertiary component. This study showed that NADESs are generally less toxic than DESs. Moreover, it emphasized the significant role of HBDs with regards to NADESs cytotoxic profiles. The use of biomaterials appears to be an important asset for lowering their cytotoxicity. Organic acids, as in previous reports, should be used with caution as they increase the deleterious attributes of NADESs. The COSMO-RS based computational approach proposed a hypothetical cytotoxic mechanism of NADESs mostly based on cellular aggregation. Although further assessment is needed to draw a comprehensive picture of the cytotoxicity mechanism of these neoteric mixtures; the results obtained in this work are encouraging with regards to their safety.

Authors' contributions

MH conceived of the study. MH, YPM, CYL, WFW, and ZS analyzed the data. MH, YPM, CYL, WFW, ZS, AH, OMA participated in both the interpretation of results and the preparation of the manuscript. All authors read and approved the final manuscript.

Author details

¹ University of Malaya Centre for Ionic Liquids (UMCiL), University of Malaya, 50603 Kuala Lumpur, Malaysia. ² Department of Civil Engineering, University of Malaya, 50603 Kuala Lumpur, Malaysia. ³ Department of Chemical Engineering, University of Malaya, 50603 Kuala Lumpur, Malaysia. ⁴ Department of Medical Microbiology, University of Malaya, 50603 Kuala Lumpur, Malaysia. ⁵ Department of Pharmacology, University of Malaya, 50603 Kuala Lumpur, Malaysia. ⁶ Institute of Halal Research University of Malaya (IHRUM), Academy of Islamic Studies, University of Malaya, 50603 Kuala Lumpur, Malaysia. ⁷ UiTM Medical Specialist Centre, University of Technology MARA, Jalan Hospital, 47000 Sungai Buloh, Selangor, Malaysia.

Acknowledgements

The authors would like to express their gratitude to the University of Malaya HIR-MOHE (D000003-16001) and to UMRG (RP037B-15AET) for their support throughout this research.

Competing interests

The authors declare that they have no competing interests.

Received: 27 January 2016 Accepted: 15 June 2016

Published online: 29 June 2016

References

- Andersson MP, Jensen JH, Stipp SLS (2013) Predicting pKa for proteins using COSMO-RS. *PeerJ* 1:e198
- Buggert M, Cadena C, Mokrushina L, Smirnova I, Maginn EJ, Arlt W (2009) COSMO-RS calculations of partition coefficients: different tools for conformation search. *Chem Eng Technol* 32:977–986. doi:10.1002/ceat.200800654
- Bushnell PJ, Boyes WK, Shafer TJ, Bale AS, Benignus VA (2007) Approaches to extrapolating animal toxicity data on organic solvents to public health. *Neurotoxicology* 28:221–226. doi:10.1016/j.neuro.2006.03.013
- Choi YH et al (2011) Are natural deep eutectic solvents the missing link in understanding cellular metabolism and physiology? *Plant Physiol* 156:1701–1705. doi:10.1104/pp.111.178426

- Cornnell RJ, Winder CL, Tiddy GJT, Goodacre R, Stephens G (2008) Accumulation of ionic liquids in *Escherichia coli* cells. *Green Chem* 10:836–841. doi:10.1039/B807214K
- da Costa Lopes AM, João KG, Rubik DF, Bogel-Lukasik E, Duarte LC, Andreus J, Bogel-Lukasik R (2013) Pre-treatment of lignocellulosic biomass using ionic liquids: wheat straw fractionation. *Bioresour Technol* 142:198–208. doi:10.1016/j.biortech.2013.05.032
- Dai Y, van Spronsen J, Witkamp G-J, Verpoorte R, Choi YH (2013) Natural deep eutectic solvents as new potential media for green technology. *Anal Chim Acta* 766:61–68. doi:10.1016/j.aca.2012.12.019
- Dai Y, Witkamp G-J, Verpoorte R, Choi YH (2015) Tailoring properties of natural deep eutectic solvents with water to facilitate their applications. *Food Chem* 187:14–19. doi:10.1016/j.foodchem.2015.03.123
- Deliconstantinos G, Villiotou V, Stavrides JC (1995) Modulation of particulate nitric oxide synthase activity and peroxynitrite synthesis in cholesterol enriched endothelial cell membranes. *Biochem Pharmacol* 49:1589–1600. doi:10.1016/0006-2952(95)00094-G
- Domínguez de María P, Maugeri Z (2011) Ionic liquids in biotransformations: from proof-of-concept to emerging deep-eutectic-solvents. *Curr Opin Chem Biol* 15:220–225. doi:10.1016/j.cbpa.2010.11.008
- Fernandez-Gomez FJ, Galindo MF, Gómez-Lázaro M, Yuste VJ, Comella JX, Aguirre N, Jordán J (2005) Malonate induces cell death via mitochondrial potential collapse and delayed swelling through an ROS-dependent pathway. *Br J Pharmacol* 144:528–537. doi:10.1038/sj.bjp.0706069
- Ferraz R, Branco LC, Prudêncio C, Noronha JP, Petrovski Z (2011) Ionic liquids as active pharmaceutical ingredients. *ChemMedChem* 6:975–985. doi:10.1002/cmdc.201100082
- Florindo C, Oliveira FS, Rebelo LPN, Fernandes AM, Marrucho IM (2014) Insights into the synthesis and properties of deep eutectic solvents based on cholinium chloride and carboxylic acids. *ACS Sustain Chem Eng* 2:2416–2425. doi:10.1021/sc500439w
- Gorke J, Srienic F, Kazlauskas R (2010) Toward advanced ionic liquids. Polar, enzyme-friendly solvents for biocatalysis. *Biotechnol Bioprocess Eng* 15:40–53. doi:10.1007/s12257-009-3079-z
- Harwood JL, Weselake RJ (2015) Lipid library. *American Oil Chemists' Society*. AOCs. <http://lipidlibrary.aocs.org/index.cfm>. Accessed 25 April 2016
- Hayyan M, Mjalli FS, Hashim MA, AlNashef IM (2010) A novel technique for separating glycerine from palm oil-based biodiesel using ionic liquids. *Fuel Process Technol* 91:116–120. doi:10.1016/j.fuproc.2009.09.002
- Hayyan A, Ali Hashim M, Mjalli FS, Hayyan M, AlNashef IM (2013a) A novel phosphonium-based deep eutectic catalyst for biodiesel production from industrial low grade crude palm oil. *Chem Eng Sci* 92:81–88. doi:10.1016/j.ces.2012.12.024
- Hayyan M, Hashim MA, Al-Saadi MA, Hayyan A, AlNashef IM, Mirghani MES (2013b) Assessment of cytotoxicity and toxicity for phosphonium-based deep eutectic solvents. *Chemosphere* 93:455–459. doi:10.1016/j.chemosphere.2013.05.013
- Hayyan M, Hashim MA, Hayyan A, Al-Saadi MA, AlNashef IM, Mirghani MES, Saheed OK (2013c) Are deep eutectic solvents benign or toxic? *Chemosphere* 90:2193–2195. doi:10.1016/j.chemosphere.2012.11.004
- Hayyan A, Hashim MA, Hayyan M, Mjalli FS, AlNashef IM (2014) A new processing route for cleaner production of biodiesel fuel using a choline chloride based deep eutectic solvent. *J Clean Prod* 65:246–251. doi:10.1016/j.jclepro.2013.08.031
- Hayyan M, Looi CY, Hayyan A, Wong WF, Hashim MA (2015) In vitro and in vivo toxicity profiling of ammonium-based deep eutectic solvents. *PLoS One* 10:e0117934. doi:10.1371/journal.pone.0117934
- Hosoya N, Kawada N (1958) Malonate metabolism in human placenta. *J Biochem* 45:363–365
- Hosoya N, Kawada N, Matsumura Y (1960) Regulatory effect of malonate on glucose metabolism in human earlier placenta. *J Biochem* 48:262–266
- Juneidi I, Hayyan M, Hashim MA (2015) Evaluation of toxicity and biodegradability for cholinium-based deep eutectic solvents. *RSC Adv* 5:83636–83647. doi:10.1039/C5RA12425E
- Kim Y-S (2002) Malonate metabolism: biochemistry, molecular biology, physiology, and industrial application. *BMB Rep* 35:443–451
- Klamt A (2005) COSMO-RS: from quantum chemistry to fluid phase thermodynamics and drug design. Elsevier, Amsterdam
- Klamt A, Eckert F (2000) COSMO-RS: a novel and efficient method for the a priori prediction of thermophysical data of liquids. *Fluid Phase Equilib* 172:43–72. doi:10.1016/S0378-3812(00)00357-5
- Kuimova MK et al (2009) Imaging intracellular viscosity of a single cell during photoinduced cell death. *Nat Chem* 1:69–73. http://www.nature.com/nchem/journal/v1/n1/supinfo/nchem.120_S1.html
- Laforenza U, Bottino C, Gastaldi G (2016) Mammalian aquaglyceroporin function in metabolism. *Biochim Biophys Acta* 1858:1–11. doi:10.1016/j.bbmem.2015.10.004
- Liu KJ-M, Drucker Y, Jarad J (1995) Hepatic glycerol metabolism in tumorous rats: a ¹³¹C nuclear magnetic resonance study. *Cancer Res* 55:761–766
- Liu T, Liu X, Spring DR, Qian X, Cui J, Xu Z (2014) Quantitatively mapping cellular viscosity with detailed organelle information via a designed PET fluorescent probe. *Sci Rep* 4:5418. doi:10.1038/srep05418. <http://www.nature.com/articles/srep05418#supplementary-information>
- Lodish H, Berk A, Zipursky SL, Matsudaira P, Baltimore D, Darnell J (2000) *Molecular cell biology*. W. H. Freeman, New York
- Lundholm K, Edström S, Karlberg I, Ekman L, Schersten T (1982) Glucose turnover, gluconeogenesis from glycerol, and estimation of net glucose cycling in cancer patients. *Cancer* 50:1142–1150
- Mulyono S, Hizaddin HF, Alnashef IM, Hashim MA, Fakeeha AH, Hadj-Kali MK (2014) Separation of BTEX aromatics from n-octane using a (tetrabutylammonium bromide + sulfolane) deep eutectic solvent—experiments and COSMO-RS prediction. *RSC Adv* 4:17597–17606. doi:10.1039/C4RA01081G
- Nadiv O, Shinitzky M, Manu H, Hecht D, Roberts CT, LeRoith D, Zick Y (1994) Elevated protein tyrosine phosphatase activity and increased membrane viscosity are associated with impaired activation of the insulin receptor kinase in old rats. *Biochem J* 298:443–450
- Paiva A, Craveiro R, Aroso I, Martins M, Reis RL, Duarte ARC (2014) Natural deep eutectic solvents—solvents for the 21st century. *ACS Sustain Chem Eng* 2:1063–1071. doi:10.1021/sc500096j
- Pereira MM, Pedro SN, Quental MV, Lima AS, Coutinho JAP, Freire MG (2015) Enhanced extraction of bovine serum albumin with aqueous biphasic systems of phosphonium- and ammonium-based ionic liquids. *J Biotechnol* 206:17–25. doi:10.1016/j.jbiotec.2015.03.028
- Plagemann PGW (1971) Choline metabolism and membrane formation in rat hepatoma cells grown in suspension culture. III. Choline transport and uptake by simple diffusion and lack of direct exchange with phosphatidylcholine. *J Lipid Res* 12:715–724
- Port AM, Ruth MR, Istfan NW (2012) Fructose consumption and cancer: is there a connection? *Curr Opin Endocrinol Diabetes Obes* 19:367–374. doi:10.1097/MED.0b013e328357f0cb
- Qi X-L, Peng X, Huang Y-Y, Li L, Wei Z-F, Zu Y-G, Fu Y-J (2015) Green and efficient extraction of bioactive flavonoids from *Equisetum palustre* L. by deep eutectic solvents-based negative pressure cavitation method combined with macroporous resin enrichment. *Ind Crops Prod* 70:142–148. doi:10.1016/j.indcrop.2015.03.026
- Radošević K, Cvjetko Bubalo M, Gaurina Srček V, Grgas D, Landeka Dragičević T, Radojčić Redovniković I (2015) Evaluation of toxicity and biodegradability of choline chloride based deep eutectic solvents. *Ecotoxicol Environ Saf* 112:46–53. doi:10.1016/j.ecoenv.2014.09.034
- Ru C, König B (2012) Low melting mixtures in organic synthesis—an alternative to ionic liquids? *Green Chem* 14:2969–2982. doi:10.1039/C2GC36005E
- Ru J et al (2015) Morphology-controlled preparation of lead powders by electrodeposition from different PbO-containing choline chloride-urea deep eutectic solvent. *Appl Surf Sci* 335:153–159. doi:10.1016/j.apsusc.2015.02.045
- Santos CR, Schulze A (2012) Lipid metabolism in cancer. *FEBS J* 279:2610–2623
- Smith EL, Abbott AP, Ryder KS (2014) Deep eutectic solvents (DESs) and their applications. *Chem Rev* 114:11060–11082. doi:10.1021/cr300162p
- Sowmiah S, Srinivasadesikan V, Tseng M-C, Chu Y-H (2009) On the chemical stabilities of ionic liquids. *Molecules* 14:3780
- Tang B, Row K (2013) Recent developments in deep eutectic solvents in chemical sciences. *Monatsh Chem* 144:1427–1454. doi:10.1007/s00706-013-1050-3
- Ullah Z, Bustam MA, Man Z (2015) Biodiesel production from waste cooking oil by acidic ionic liquid as a catalyst. *Renew Energy* 77:521–526. doi:10.1016/j.renene.2014.12.040

- Vlachy N, Jagoda-Cwiklik B, Vácha R, Touraud D, Jungwirth P, Kunz W (2009) Hofmeister series and specific interactions of charged headgroups with aqueous ions. *Adv Colloid Interface Sci* 146:42–47. doi:[10.1016/j.cis.2008.09.010](https://doi.org/10.1016/j.cis.2008.09.010)
- Wagle DV, Zhao H, Baker GA (2014) Deep eutectic solvents: sustainable media for nanoscale and functional materials. *Acc Chem Res* 47:2299–2308. doi:[10.1021/ar5000488](https://doi.org/10.1021/ar5000488)
- Wen Q, Chen J-X, Tang Y-L, Wang J, Yang Z (2015) Assessing the toxicity and biodegradability of deep eutectic solvents. *Chemosphere* 132:63–69. doi:[10.1016/j.chemosphere.2015.02.061](https://doi.org/10.1016/j.chemosphere.2015.02.061)
- Wu B-P, Wen Q, Xu H, Yang Z (2014) Insights into the impact of deep eutectic solvents on horseradish peroxidase: activity, stability and structure. *J Mol Catal B Enzym* 101:101–107. doi:[10.1016/j.molcatb.2014.01.001](https://doi.org/10.1016/j.molcatb.2014.01.001)
- Zhang Q, De Oliveira Vigier K, Royer S, Jerome F (2012) Deep eutectic solvents: syntheses, properties and applications. *Chem Soc Rev* 41:7108–7146. doi:[10.1039/C2CS35178A](https://doi.org/10.1039/C2CS35178A)
- Zhao B-Y, Xu P, Yang F-X, Wu H, Zong M-H, Lou W-Y (2015) Biocompatible deep eutectic solvents based on choline chloride: characterization and application to the extraction of rutin from *Sophora japonica*. *ACS Sustain Chem Eng* 3:2746–2755. doi:[10.1021/acssuschemeng.5b00619](https://doi.org/10.1021/acssuschemeng.5b00619)

Submit your manuscript to a SpringerOpen[®] journal and benefit from:

- ▶ Convenient online submission
- ▶ Rigorous peer review
- ▶ Immediate publication on acceptance
- ▶ Open access: articles freely available online
- ▶ High visibility within the field
- ▶ Retaining the copyright to your article

Submit your next manuscript at ▶ springeropen.com
