



A REVIEW OF GREEN SOLVENT IONIC LIQUIDS: AS A FUTURE SOLVENT

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ABSTRACT

Currently, chemists seem more fascinated with exploiting less hazardous green solvent ionic liquids for countering the detrimental consequence of organic solvents on the environment. Ordinarily, ionic liquids having salt-like structures comprise cations and anions. Ionic liquids alleviate various enzyme-catalyzed reactions having excellent yields, increasing capacities and functions, enhancing recyclability, increasing stability, recovery, and cost-effectiveness, and these are cardinal elements for applications in industry. Furthermore, growing nanotechnology, polymer, electrochemistry, environmental chemistry, pharmaceutical medicinal chemistry or pharmaceutical chemistry, and biochemistry in the area of ionic liquids have been flourished among investigators. Ionic liquids-supported materials and techniques are excellent for efficient and sustainable procedures. Because of their assignment as designer solvents, through which the physico-chemical dimensions of an Ionic liquid can be adjusted for better functioning for peculiar applications. The review delivers the gradual improvement, growth or development, or progress and prospects of ionic liquids and their physicochemical properties as well as examples of applications in a unified fashion.

Keywords: Ionic liquids, Physio-chemical properties, Separation Science, Nanomaterials, Ionanofluids, Fossil fuel cleaner.

1. INTRODUCTION

Magical chemical ionic liquids, having unique tunable properties with a variety of applications in every field have attracted tremendous attention from scientists ecumenically. The history of liquids effectively started in 1914 when the inception of the first ionic liquid (IL) ethyl ammonium nitrate ($[\text{EtNH}_3][\text{NO}_3]$) with a melting point of 12°C was reported. Years later, a procession of remarkable related ionic liquids has been revealed unusually, have established enormous pursuit in the chemical industry as well as in academia because of their tremendous greener properties and exist as a liquid at below 100°C temperature. Ionic liquids (ILs) have no boundary between organic and inorganic materials because of their tunable properties. Furthermore, target-specific ionic liquids with tunable properties can be designed by changing cation-anion combinations. Examples of several normally employed cationic constituents are pyrrolidinium [1], imidazolium cations [2], ammonium cations [3], pyridinium cations [4], phosphonium cations [5], and piperidinium cations [6] with variant substituents and anionic constituents tetrafluoroborate anions, alkyl sulphonate anions, alkyl

tosylate anions, and hexafluorophosphate anions for production of ILs. As a result, ILs have many advantages over the traditional organic solvents (Table1) such as very low volatility, trifling vapor pressure [7], less hazardous, inflammability [8], solubility, and more environmental compatibility, viscosity [9], high chemical and thermal stability [10], prominent ionic conductivity [11-14]. Usually, the viscosity of ILs relies on and enhances with increasing hydrogen bonding, van der Waals forces, length of the alkyl chain, and degree of fluorination or anion. Furthermore, normally, at room temperature, ILs's viscosity has a scope between 10 and 500 mPa. [15-18].

2. PHYSIO-CHEMICAL PROPERTIES OF IONIC LIQUIDS

Remarkably, some of their canonical dimensions are [19-29]:

- Potentiality to solvate numerous organic, organo-metallic, and inorganic substances.
- Prominent polarity.
- Loose-fitting coordinating large-size ions.

- In general, very low vapor pressures as well as low volatility.
- Thermal constancy up to 300°C approximately.
- Eminent thermal conductance and an enormous electrochemical window.
- Liquid windows of ILs are up to 200°C to empower broad kinetic control.
- Nonaqueous polar options to phase transfer procedures.
- They have been acknowledged as universal solvents.
- They are capable to solvate still gases such as H₂, CO₂, O₂, and CO. They have been applied under supercritical CO₂.
- In ILs, the dissolubility deciding components are anions and cations from which these are framed.
- They do not coordinate with macrocycle metal complexes such as enzymes, etc. chiefly, the ionic properties of ILs enhance the rate of reactions.
- They are stable and can be stored without decomposition for a long time.
- ILs have found extensive use in the control of stereo selectivity.
- The viscosity of imidazolium-modified ILs can be controlled by modifications in branching.
- They have melting point preferably below 100°C.
- The dielectric constant of ILs has been implied to <30.

3. APPLICATIONS OF IONIC LIQUIDS

On account of their fascinating properties, ILs have been exploited in a wide area of applications which are listed below [30-54]:

Medium extraction [30], physico-chemical procedures [31], ILs as a medium for many reactions [31], in the modification of the mobile phase of HPLC [32], metals electrodeposition, and semiconductors in ILs [33], in the analysis of chemicals [34], dye-sensitized solar cells [35, 36], in the nuclear fuel chain: extraction and electrodeposition [37], nuclear-supported separations [38], oil shale treating [39], separation of petrochemical relevance [40], development of functional nanoparticles as well as different inorganic nanostructures [41], ILs as mediums for electrochemistry [42], ILs as mediums in polymerization procedures [43], chemical as well as biochemical conversions [44], materials chemistry [45], ILs as a biocatalyst, or biocatalysts in ILs [46-55], utilization as green catalysts, chromatographic actions; ILs have been depicted as immense per-formances in

separation techniques [56], as separators for extraction of metal ions, Ionic liquids modified sensing-Chemosensing has been examined by employing ILs and detected the fast reduction in viscosity, in spectroscopy; The applications of ILs in mass spectroscopy are speedily raising after being accomplished by Armstrong's group. These are some of the universal applications of ILs. Several applications that are associated with oil as well as with the gas industry comprise:

Removal of sulfur from fuels, naphthenic acids extraction, removal of nitrogen from gasoline, utilization as a demulsifiers agent, eliminate contamination, selective gas extraction and mercury elimination in natural gas, production of Biofuels, in improving oil recovery, CO₂ is captivating and sequestrant, application as asphaltene and wax retardant, recovery of heavy oils or bitumen with ILs and in oil fields, deep eutectic media is being extended. Ionic liquids (ILs) have come out into view as a mesmerizing green reusable substitute for toxic, dangerous, combustible, and extremely volatile organic compounds [58-63]. ILs are widely exploited in chemistry, nanotechnology [64, 65], engineering [66-68], biotechnology [69-74], pharmaceutical industries [75-80]. ILs have many exertion like as natural fibers processing, carbon dioxide capture [81-83], separation, battery [84, 85], biofuel formation, extraction [86], biological activities, solvents [87, 88], catalysis [89-92], fuel cells [93] and biomedical uses [94]. Several reports have included a total or partial degradation of the epoxy thermoset polymers as well as utilized the end productions for making functional materials for various applications [95-98]. Other approaches applied by many investigators include chemical decomposition by the base or acid reactions [99-101] or by the implementation of a catalyst for degradation of the Epoxy-resin [102-105].

4. IONIC LIQUIDS AND IONIC SALTS

Ionic liquids are magnificently distinct from ionic salts such as NaCl [106]. Ionic salts exist as solid at room temperature and have electrostatics interaction with normally metallic cations and non-metallic anions. On the whole, ionic materials such as NaCl can be in a liquid or molten state at high temperatures so can not be used in organic synthesis. Whereas, ionic liquids, a category of ionic salts which have melting points below 100°C, are soluble in organo-metallic as well as inorganic-organic compounds and are used in organic synthesis [107].

Table 1: Comparison between Ionic liquids and organic solvents [57]

Serial Number	Property	Ionic liquids	Organic solvents
1	Number of solvents	More than 1,000,000	More than 1000
2	Applicableness	Multifunctional	One work
3	Catalytic capability	Frequent and tunable	Uncommon
4	Chiral Property	Frequent and tunable	Uncommon
5	Refractive Index	From 1.5 to 2.2	From 1.3 to 1.6
6	Viscosity (cP)	From 22 to 40,000	From 0.2 to 100
7	Polar property	Employ doubtful conceptions of polarity	Employ traditional conceptions of polarity
8	Density (g/cm^3)	From 0.8 to 3.3	From 0.6 to 1.7
9	Recyclability	Economical clamant	Green clamant
10	The capability of being tuned	The almost limitless scope entails designed solvents	The narrow scope of solvents accessible
11	Solvency	Potent Solvency	Feeble Solvency
12	Inflammability	Generally inflammable	Generally flammable
13	Monetary Value	Mostly low-priced	Mostly from two to hundreds of times than that of organic solvents
14	Vapor pressure	Worthless vapor pressure	Follow the Clausius-Claperton equation

5. APPLICATIONS OF IONIC LIQUIDS

5.1. Applications in separation science

Outstanding holdings of IL, ability to synthesize with different solubility in water as well as in organic solvents such as methanol or ethanenitrile makes their use in separation technique newsworthy. Due to the viscous nature, ILs have been utilized in chromatography as an additive and after adding High-Performance Liquid Chromatography solvents viscosities of ILs reduced [108, 109]. In a special separation technique capillary electrophoresis (CE), ILs are used as background electrolytes as well as additives. Additionally, ILs modify the capillary wall and electrophoretic separations [110-112]. ILs are also used in the separation of flavonoids from phenolic compounds and imidazolium ionic liquid, and carprofen, naproxen, ketoprofen, and suprofen from (C_4mim) (NTf_2). Natural product researchers are using ILs as a green method to extract and isolate from plants-like alkaloids, flavonoids, terpenoids, and phenolics [113-116].

5.2. Environmental acceptable green extraction techniques

Conventionally, scientists ordinarily utilize hazardous volatile organic compounds (VOCs) in extraction techniques to have organic extracts like soxhlet, distillation, percolation, maceration, distillation, and infusion. On the other hand, all green chemistry scientists have long attempted for replacing conventional methods which have an environmental

impact with green methods such as supercritical fluid extraction, ultrasound-assisted extraction, microwave-assisted extraction, high-speed homogenization, and pulsed electric field, and pressurized solvents. ILs scale down solvent uptake and extraction time. Supercritical fluid extraction has fantabulous extraction properties but high operational costs than ILs [117-121].

5.3. Role of ionic liquids as a catalyst for the preparation of biodiesel

Negligible vapor pressure, high thermal stability, phase transition behavior, solubility and miscibility with reactants, acidity and basicity, switchable ionic liquids as green solvents, ionic liquids in biodiesel synthesis, catalyst, solvent for enzyme-catalyzed transesterification, catalyst support, recycling of ionic liquids, ionic liquids recovery, deep eutectic solvents (DES) such as choline chloride, a peculiar category of ILs regarded as a new generation of ionic liquids that form hydrogen bonding. Additionally, hydrogen-bonding interactions in deep eutectic solvents (DES) are energetically favorable as a result of more solubility. Researchers have used a combination of deep eutectic solvents and ILs in biodiesel preparation [122,123].

5.4. Protein solubility and stability in ionic liquids

Many publications on protein solubility and consistency have been published in aqueous solution but in some cases solvation and constancy of proteins in water are

pocket-size and these difficulties can be subdued by using ILs as co-solvents with an aqueous medium in two-phase systems or as dexteros ILs [124-126] and also ameliorate the separation effectiveness of essential biomolecules like carbohydrates, amino acids, alkaloids, proteins, antibiotics, and various solvents [127, 128]. The stability of a protein is important for the purification and extraction process for their diligence because alterations in the protein surroundings can modify its pure form. Classical methods of protein refinement like liquid-liquid extraction, electrophoresis, and chromatography, have convinced disadvantages admitting huge prices, time uptake, deprivation in biological functions, and insufficiency of lustiness. Ionic liquid-based aqueous biphasic systems can get over these disadvantages [129-132]. IL-supported aqueous state extraction for protein separation is a desirable alternative.

5.5. Biocatalysis in the presence of ionic liquids

The foremost booming study of biocatalysis in presence of ILs is done in 2000 [133]. Functions and consistency of a vast figure of enzymes for example tremendously employed lipases [134-137], oxidoreductases, proteases [138, 139], and alcohol dehydrogenases, and compatibility with other enzymes have been measured. The functioning of enzymes in ILs is stimulated by numerous components such as anions which can form hydrogen bonding with enzymes and make configurational modifications, cations that have the power to form van der waals interactions, the alkyl chain extension enhance enzyme constancy, as well as viscosity, hydrophobicity that can be elevated by an alkyl chain, lengthened, affects the folding or refolding of enzymes [140-143]. Both hydrolases enzymes as lipases have mostly habituated in reactions transesterification, perhydrolysis, ammonolysis, polyester synthesis [144-151], proteases such as thermolysin have applied commonly in peptide synthesis [152], chymotrypsin has been used in transesterification [153-156], esterase has exploited in transesterification [157], glycosidase like as galactosidase has employed in N-acetylactosamine synthesis and oxidoreductases for examples formate dehydrogenase in the regeneration of NADH [158], baker's yeast in reduction of ketones, peroxidases in the oxidation of guaiacol, laccase C in the oxidation of syringaldazine [159-161], have been expended, hold back natural functions with ionic liquids. In many reactions, proteases and lipases show enantioselectivity in ILs [162].

5.6. Ionic liquids as a fossil fuel cleaner

For the expeditious formation of cleaner fossil fuels, ILs possess various anion and cation additions that have been programmed to efficaciously draw out sulfur chemical compounds likewise dibenzothiophene out of fossil fuels under atmospheric considerations. On account of its extremely noxious and destructive nature, the quantity of H₂S is hard to govern in natural gas, and the quantity of H₂S should be less than 6mg/m³. Consequently, the establishment of ILs for withdrawing H₂S has appealed to modernize attention freshly. Many kinds of ILs have been examined for their dissolubility of H₂S, and their functioning. The dissolubility and discrimination of H₂S are due to attractions in anions and H₂S, favorable prominent CO₂ concentrations. Chemists have reported that imidazolium-based ionic liquids can be employed as an extraordinary excerpt to remove the sulfur from liquid fuels, and dibenzothiophene [163,164].

5.7. Ionic liquids and polysaccharide cellulose

Polysaccharide cellulose-supported kinds of stuff have been exercised in the global industries [165] and have fascinating attributes like biological harmony, biodegradability, best solubility, recyclable, chemical as well as thermal constancy [166, 167]. Because cellulose is indissoluble in traditional solutions, current research universally proposes to discover novel, effective, and green solvents for biomass such as cellulose, so attempts have concentrated on the exploitation of ionic liquids for receiving solubility of cellulose. Currently, a research report has found the association between the dissolution and the degree of polymerization (DP) of cellulose [168], and the dissolution of cellulose can be decreased with the enhancing DP.

5.8. Role of ionic liquids in nanomaterials

The stability of nanomaterials in the various solvents has been essential for beautifying versatile usage in divergent areas. Moreover, the solution works a cardinal function in altering and calibrating the physiochemical attributes of the nanomaterials [169, 170]. Imidazolium-supported ILs have been depicted for ameliorating dimensions of the nanomaterials such as carbon nanotubes [171], and offer surface monotony of nanoparticles without throwing away properties like no alteration in nanoparticle size as well as in size dispersion, allowing for reusability as catalysis, and comfortably separation from the reaction solutions [171]. According to reference, imidazolium-supported ILs have behaved as capping agents [172,173]. IL-nanomaterial composites

1-butyl-3-methylimidazolium tetrafluoroborate has been applied to make IL-ZnO composites to increase the potent capacity of dasatinib and doxorubicin for breast antitumor curatives [174]. 1-ethyl-3-methylimidazolium methanesulfonate stabilized PbS, CdS nanoparticles [175,176].

Graphene oxide-supported ionic liquids methods have been developed to find out traces of Hg in water [177]. Moreover, carbon quantum dots capped with imidazolium-supported ILs by pyrolysis have been groomed and presently, citrate-based ILs have been applied to develop the Ag nanoparticles capped with IL [178], and nanosheets. According to Okoli, [179] the sole function of ILs is the generation of best attribute metal alloy nanoparticles to raise catalytic functions. Imidazolium modified ILs such as 1-methacryloyloxypropyl-3-methylimidazolium bromide [180] for Al_2O_3 , TiO_2 , Fe_3O_4 , 1-butyl-3-methylimidazolium methyl sulfate [181] for ZnSe with 70-100 nm size, PbS from 1-n-butyl-3-methylimidazolium hexafluorophosphate with 100 nm size [182], 1-alkyl-3-methylimidazolium N-bis(trifluoromethane sulfonyl) [183] for Ni with 4.9–5.9 nm size, n-butyltrimethylammonium N-bis(trifluoromethylsulfonyl)imide [184] for Au in rage 140nm, 1-n-butyl-3-methyl trifluoromethane sulfonate for Ir with 2.4-2.6 nm size, were applied to synthesis nanoparticle [185].

5.9. Electrochemical applications of ionic liquids

Application of early acknowledged IL-supported activity has been done in 1996. The intrinsic conductance of ILs builds ILs desirable for varied uses in electrochemistry like as chromium electroplating of scionix procedure [186-188], providing a securer option for toxicity of chromium (IV) salts, in reinforced current power, exempt from cracked, rusting resistive coatings, in a diverse metal procedures employment such as metal plating and metal polishing [189-192].

Various commercial ILs have launched demand in batteries for ameliorating protection, enhanced eminent-voltage steadiness, in ion-Li batteries work as an electrolyte [193, 194]; and also, in Zn modified air batteries, Zn based Br batteries [195-198], gas sensors, supercapacitors, cheap option for crystalline silicon-based cells, in photovoltaic cells which are dye sensitive [199-201]. Various companies are acknowledged to employ the ILs's electrochemical attributes like as IoLiTec as well as C-Tech Innovation for aluminum plating, BASF has applied imidazolium modified ILs to

aluminum plating, and Xtalic has employed nanoscale aluminum metal alloy plating [200, 202-206].

5.10. The function of ionic liquids in alkylation

According to recent reports, ILs are an appropriate and securer substitute to invent gasoline from lower hydrocarbons by alkylation [207], however, in the conventional methods alkylation procedure is made up of catalytic action of corrosive acids. In cooperation with QUILL, Chevron started the study of ILs for catalyzing alkylation as a catalyst in 1999 and performed research establishment with their ISOALKYTM discipline for five years long till 2010 for perfecting chemical reaction circumstances [208-210]. The unit of ISOALKYTM which is authenticated by Honeywell UOP takes a lower quantity of catalyst because of their more prominent capability than based-acid operations, enhanced production yield, and attributes with analogous manufacturing as well as operating prices, and also in magnanimous technical applications [211-213].

5.11. The activity of ionic liquids as a capture

In 2005, wind products accounted that they had generated a novel technique to collect and transfer hazardous gases which are broadly applied in the electrochemical industry for doping silicon [214, 215]. Cooperation with investigators in PETRONAS and QUILL resulted in the exploitation of a new technique of withdrawing mercury (Hg) from natural gas currents, with an extraordinarily less execution period [216-220]. Springing up the concern in supported ionic liquids phase (SILP) modified catalyst process headed to the research of proportion synthesis process [221].

5.12. Utility of ionic liquids in coating

The traditional synthesis process involves the dissolution of catalyst in proper solution adopted by the add-on of IL and a poriferous catalyst-based substance and sequential separation of the solvent. Extraction should be done slowly to assuring dispersal of the catalyst as well as IL throughout the poriferous base. According to documented reports, SILP modified substances can be manifoldly produced by employing a fluidized bed spray-based coating, through which supported substances are liquified by applying managed temperature inert gases, catalyst dissolution, and IL in a dissolution is consecutively sprinkled on the support, likewise, this procedure can be applied to prepare ionic liquid supported solid catalyst, incorporating a conventional catalyst covered with a fine film of IL. In

addition, this process has been commercially exploited for a hydrogenation procedure, also impacting activity [222-225].

5.13. Use of ionic liquids in polymer chemistry

ILs having eminent optical limpidity with less hazing work as additives such as antistatic additives for ameliorating neatness, and enhancing security. ILs have harmony with many usual working methods or techniques as well as polymers [226-229]. According to Evonik, ILs as an auxiliary dissipate additive for uniformly reinforcing water-modified pigments in water as well as solvent-modified paints and stopping sedimentation [230,231]. Moreover, IoLiTec is acknowledged for making and conveying its investigation and exploitation of IL modified functioning additives to a broad kind of utilizations: diffusing agents, optic brighteners, neatness additives, formation of inorganic substances having definite particle sizes on a commercial scale, and preparation of alcohols on the pilot-scale [222, 232, 233]. In natural fiber welding procedures, ILs have been exploited to procedure natural fiber materials for making congealable networks to sustain the original polymer structure [234-236]. The qualities of the novel material, as well as the magnitude of the welding procedure, have been governed by time, temperature/heat, pressure or ionic liquids nature, quantity, and placement [237]. Trouble in recyclate of plastic is being accepted as a challenge by ILs modified technique and the mainsail problem overcome by plastics recycling because it is frequently much economy to landfill, and incinerate waste matters [238].

5.14. Ionic liquids as active ingredients of pharmaceutical preparations

Crystal polymorphism and less solubility are the most considerable difficulty in the pharmaceutical industry [239-243]. ILs have been depicted as a substitute for conventional drug delivery systems for increasing dissolubility and consistency. Currently produced reports have proved that many ILs hold protein structures [244] so ILs are assured as a stabilizing media to protein remedies [245-248] and also are capable to substitute water and more expensive lyophilic substances in protein supported pharmaceutical formulations. According to currently published reports, ILs have a destructive effect on bacterial and fungal growth, due to this ILs can be used for removing pathogenic micro-organisms tolerant to antibiotics and different medicines [249, 250]. There are numerous

illustrations of exploiting ILs in medicinal chemistry, disinfectant detergents [251], as constituents of drugs, tumor treatment [252-254], and novel bioactive substances such as antiseptics [255-258], to oppose microbial biofilms which have various bacteria [259]. Benzalkonium and ammonium supported ILs auspicious action against fungi and bacteria peculiarly for *Streptococcus mutans*. Various ILs have broad-spectrum actions to oppose pathogenic micro-organisms [259]. Another major trouble regarding the bioavailability of drugs is difficulty in traveling across biological membranes because of the prominent hydrophilic nature and this drawback can be overcome by the proper choice of different active cations and anions to combine these with hydrophilic ones, also by the coating of ILs [259, 260]. Employing ILs as constituents of drugs demands organized determination of toxicological attributes and cytotoxicity [261,262]. Moreover, by exploiting ILs as constituents of pharmaceutical formulations, the osmotic coefficient potency should be precisely determined. Phosphates, as well as choline, have been picked out as initiating.

5.15. Selection of ionic liquids and preparation of ionanofluids

For heat transfer-supported uses, ILs can be chiefly picked out grounded on their physical properties (Table 2) especially of eminent thermal conductance, heat capacity, negligible viscosity, and dissolubility in water. Applications and properties of prominent functioning ionanofluids (INF) depend on the processes by which they are formed. The formation process of high functioning ionanofluids is a one-step process and a two-step process. In a one-step process, nanomaterials to ionanofluids are established directly without supported ILs, and seldomly employed. However, in a generally employed two-step process, ionanofluids are usually developed by using nanomaterials supported ILs, after that homogeneousness is chiefly developed by ultrasonication.

Several kinds of nanomaterials like graphene, carbon nanotubes, and imidazolium-modified ionic liquids have been exploited for the grooming of ionanofluids. The formation process of ionanofluids is quite free from ambiguity, however, it has a stimulating interest in verifying thoroughly uniform dispersal of nanoparticles and long-time constancy of developed ionanofluids. Stability of developed ionanofluids can be ameliorated by adding sonication and surface-active agents and in other manners by surfacing treatment or alteration of

nanoparticles. It is essential to notice that specific alertness should be devoted to ultrasonic ionanofluids because beyond normal limits sonication can devolve the substance in both physical and chemical conditions, evaporate substance ionanofluids, and in the alteration of concentration of nanomaterials. Surfactants have been

also suggested not to exercise because they can degenerate or can get dormant at average and higher temperatures. Determination of stability of ionanofluids admits measuring Zeta potential, SEM, TEM investigation thermophysical properties of ILs and INFs used on convective heat transfer applications [263-265].

Table 2: Thermophysical properties of ionic liquids expanded as basis fluids for ionanofluids

Ionic Liquid	Thermophysical Property	Thermophysical Property's value	Condition	Reference
1-butyl-3-methylimidazolium bis(trifluoromethylsulfonyl)imide [C ₄ mim] [NTf ₂]	Viscosity (mPa·s)	99.6	298 K	[266]
	Density (kg/m ³)	1.436	298 K	[267]
	Thermal Conductivity (W/m K)	0.126	300 K	[268]
	Heat Capacity (kJ/kg·K)	1.352	298 K	[269]
1-hexyl-3-methylimidazolium bis(trifluoromethylsulfonyl)imide [C ₆ mim] [NTf ₂]	Viscosity (mPa·s)	70.5	298 K	[270]
	Density (kg/m ³)	1.372	298 K	[271]
	Thermal Conductivity (W/m K)	0.122	293 K	[272]
	Heat Capacity (kJ/kg·K)	1.426	298 K	[271]
1-n-butyl-3-methylimidazoliumtrifluoromethanesulfonate [C ₄ mim] [CF ₃ SO ₃]	Viscosity (mPa·s)	76.0	298 K	[273]
	Density (kg/m ³)	1.306	293 K	[274]
	Thermal Conductivity (W/m K)	0.142	293 K	[272]
	Heat Capacity (kJ/kg·K)	1.484	298 K	[275]
1-butyl-methylimidazolium hexafluorophosphate [C ₄ mim] [PF ₆]	Viscosity (mPa·s)	257.0	298 K	[273]
	Density (kg/m ³)	1.372	293 K	[274]
	Thermal Conductivity (W/m K)	0.145	293 K	[276]
	Heat Capacity (kJ/kg·K)	1.432	308 K	[276]
1-hexyl-3-methylimidazolium hexafluorophosphate [C ₆ mim] [PF ₆]	Viscosity (mPa·s)	485.8	298 K	[277]
	Density (kg/m ³)	1.293	298 K	[277]
	Thermal Conductivity (W/m K)	0.142	293 K	[274]
	Heat Capacity (kJ/kg·K)	1.358	293 K	[277]
1-n-butyl-3-methylimidazolium dicyanamide [C ₄ mim] [DCA]	Thermal Conductivity (W/m K)	0.176	298 K	[278]
	Heat Capacity (kJ/kg·K)	1.827	296 K	[279]
1-ethyl-3-methylimidazoliummethylsulfate [C ₂ mim] [EtSO ₄]	Viscosity (mPa·s)	125.4	293 K	[280]
	Density (kg/m ³)	1.236	298 K	[280]
	Thermal Conductivity (W/m K)	0.1706	293.4 K	[281]
	Heat Capacity (kJ/kg·K)	1.57	293 K	[282]
1-n-butyl-3-methylimidazolium bis(trifluoromethanesulfonylimid) [C ₄ mim] [(CF ₃ SO ₂) ₂ N]	Viscosity (mPa·s)	51.1	313 K	[283]
	Density (kg/m ³)	1.426	313 K	[283]
	Thermal Conductivity (W/m K)	0.1114	293.4 K	[281]
	Heat Capacity (kJ/kg·K)	1.373	313 K	[283]
1-hexyl-3-methylimidazolium tetrafluoroborate [C ₆ mim] [BF ₄]	Viscosity (mPa·s)	250.0	298 K	[284]
	Density (kg/m ³)	1.149	298 K	[284]
	Thermal Conductivity (W/m K)	2.21	298 K	[284]
	Heat Capacity (kJ/kg·K)	0.166	298 K	[284]
1-Butyl-3-methylimidazolium tetrafluoroborate [C ₄ mim] [BF ₄]	Viscosity (mPa·s)	85.37	303.15 K	[285]
	Density (kg/m ³)	1.198	303.15 K	[285]
	Thermal Conductivity (W/m K)	0.163	298 K	[272]
	Heat Capacity (kJ/kg·K)	1.614	298 K	[272]
1-Ethyl-3-methylimidazolium methanesulfonate [C ₂ mim] [CH ₃ SO ₃]	Viscosity (mPa·s)	149.1	298 K	[286]
	Density (kg/m ³)	1.239	298 K	[286]
	Thermal Conductivity (W/m K)	0.190	298 K	[287]
	Heat Capacity (kJ/kg·K)	1.629	298 K	[287]
Butyltrimethylammoniumbis	Viscosity (mPa·s)	105.4	298 K	[288]

(trifluoromethylsulfonyl)imide [N4111] [NTf ₂]	Density (kg/m ³)	1.392	298 K	[289]
	Thermal Conductivity (W/m K)	0.122	303 K	[289]
	Heat Capacity (kJ/kg·K)	1.70	303 K	[289]
N-butyl-N- methylpyrrolidiniumbis(trifluoro methanesulfonyl)imide [C ₄ mppyrr [NTf ₂]	Viscosity(mPa·s)	68	303 K	[290]
	Density (kg/m ³)	1.382	298 K	[291]
	Thermal Conductivity (W/m K)	0.124	303 K	[289]
1-Methylimidazolium tetrafluoroborate [HMIM] [BF ₄]	Heat Capacity (kJ/kg·K)	1.58	303 K	[289]
	Viscosity (mPa·s)	250	298 K	[292]
	Density (kg/m ³)	1.123	298 K	[293]
Trihexyltetradecylphosphoniumbis (trifluoromethylsulfonyl)imide [(C ₆) ₃ PC14] [NTf ₂]	Thermal Conductivity (W/m K)	0.166	298 K	[292]
	Heat Capacity (kJ/kg·K)	2.265	298 K	[292]
	Viscosity (mPa·s)	318	298 K	[294]
1-Ethyl-3-Methylimidazolium Diethyl Phosphate [EMIM] [DEP]	Density (kg/m ³)	1.065	298 K	[295]
	Thermal Conductivity (W/m K)	0.137	298 K	[295]
	Heat Capacity (kJ/kg·K)	1.788	333 K	[295]
1-n-butyl-3-methylimidazolium chloride [C ₄ mim] [Cl]	Viscosity (mPa·s)	274	298 K	[296]
	Density (kg/m ³)	1.148	298 K	[296]
	Thermal Conductivity (W/m K)	0.1749	303 K	[297]
1-Ethyl-3-Methylimidazolium dicyanamide [EMIM] [DCA]	Heat Capacity (kJ/kg·K)	1.998	293 K	[297]
	Viscosity (mPa·s)	545	333 K	[298]
	Density (kg/m ³)	1.087	293 K	[298]
Trihexyltetradecylphosphoniumph osphate[(C ₆) ₃ PC14] [Phosph]	Thermal Conductivity (W/m K)	0.176	293 K	[298]
	Heat Capacity (kJ/kg·K)	1.982	298 K	[299]
	Viscosity (mPa·s)	13.2	300 K	[300]
1-butyl-3-methylimidazolium bromide[C ₄ mim] [Br]	Density (kg/m ³)	1.1	298 K	[301]
	Thermal Conductivity (W/m K)	0.135	298 K	[295]
	Heat Capacity (kJ/kg·K)	2.12	298 K	[295]
1-butyl-3-methylimidazolium iodide[C ₄ mim] [I]	Viscosity (mPa·s)	215	303 K	[298]
	Density (kg/m ³)	1.298	293 K	[298]
	Thermal Conductivity (W/m K)	0.16	293 K	[298]
	Heat Capacity (kJ/kg·K)	1.421	298 K	[269]
	Viscosity (mPa·s)	379	303 K	[298]
	Density (kg/m ³)	1.489	293 K	[298]
	Thermal Conductivity (W/m K)	0.131	293 K	[298]
	Heat Capacity (kJ/kg·K)	1.165	298 K	[302]

6. CONCLUSION

To respond to the harmful consequence of chemical substances on the natural world, institutional as well as industrial research concerns have transferred towards environment-friendly ILs with negligible vapor pressure, high thermal stability, phase transition behavior, solubility, and miscibility with reactants, acidity, and basicity, switchable ionic liquids as green solvents, ionic liquids in biodiesel synthesis, catalyst, solvent for enzyme-catalyzed transesterification, catalyst support, recycling of ionic liquids, ionic liquids recovery. Ionic liquids are not future solvents exclusively due to their green dimensions, however, that is not to express in words that belongings, like negligible flammability or worthless volatility, do not

encourage the overall process. The concept is that ionic liquids are made up solely of ions has led to their future solvent applications with a large range of electrochemical windows, prominent conductance, and high thermal stability.

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Conflicts of interest

The authors declare no conflict of interest

8. REFERENCES

- Pont AL, Marcilla R, De Meazza I, Grande H, Mecerreyes D. *Journal of Power Sources*, 2009; **188(2)**:558-563.
- Freire MG, Neves CM, Marrucho IM, Coutinho JA, Fernandes AM. *The Journal of Physical Chemistry A*, 2009; **114(11)**:3744-3749.
- Kogelnig D, Stojanovic A, Galanski M, Groessel M, Jirsa F, Krachler R, Keppler BK. *Tetrahedron Letters*, 2008; **49(17)**:2782-2785.
- Gao H, Luo M, Xing J, Wu Y, Li Y, Li W, Liu Q, Liu H. *Industrial & Engineering Chemistry Research*, 2008; **47(21)**:8384-8388.
- Breitbach ZS, Armstrong DW. *Analytical and Bioanalytical Chemistry*, 2008; **390(6)**:1605-1617.
- Paduszyński K, Domańska U. *The Journal of Chemical Thermodynamics*, 2013; **60**:169-178.
- Earle MJ, Esperança JM, Gilea MA, et al. *Nature*, 2006; **439(7078)**:831-834.
- Ye C, Liu W, Chen Y, Yu L. *Chemical Communications*, 2001; **21**:2244-2245.
- Seddon KR. *Journal of Chemical Technology and Biotechnology*, 1997; **68(4)**:351-356.
- He Z, Alexandridis P. *Physical Chemistry Chemical Physics*, 2015; **17(28)**:18238-18261.
- Noda A, Hayamizu K, Watanabe M. *The Journal of Physical Chemistry B*, 2001; **105(20)**:4603-4610.
- Welton T. *Chemical Reviews*, 1999; **99**: 2071.
- Shariati K, Gutkowski CJ, Peters. *AIChE Journal*, 2005; **51**:1532.
- Brennecke JF, Maginn EJ. *AIChE Journal*, 2001; **47**:2384.
- Varma R. *Green Chemistry*, 1999; **1(1)**:43-55.
- Davoodnia A, Heravi MM, Safavi-Rad Z, Tavakoli-Hoseini N. *Synthetic Communications*, 2010; **40(17)**:2588-2597.
- Endres F, ElAbedin SZ. *Physical Chemistry Chemical Physics*, 2006; **8(18)**:2101-2116.
- Bonhote P, Dias AP, Papageorgiou N, Kalyanasundaram K, Grätzel M. *Inorganic Chemistry*, 1996; **35(5)**: 1168-1178.
- Keskin S, Kayrak-Talay D, Akman U, Hortaçsu Ö. *J. Supercrit Fluids*, 2007; **43**:150-180.
- Welton T. *Chem Rev*, 1999; **99**:2071-2084.
- Gorman J. *Sci News*, 2001; **160**:156-158.
- Brennecke JF, Maginn EJ. *AIChE J*, 2001; **47**:2384-2388.
- Yang Q, Dionysiou DD. *J. Photochem Photobiol A Chem*, 2004; **165**:229-240.
- Seddon KR. *Kinet Catal*, 1996; **37**:693-697.
- Martinez-Palou RJ. *Mex Chem Soc*, 2007; **51**:252-264.
- Hoffmann J, Nuchter M, Ondruschka B, Wasserscheid P. *Green Chem*, 2003; **5**: 296-299.
- Shariati A, Peters CJ. *Supercrit Fluids*, 2005; **34**:171-176.
- Shariati A, Gutkowski K, Peters CJ. *AIChE J*, 2005; **51**:1532-1540.
- Zhao H, Xia S, Ma PJ. *Chem Technol Biotechnol*, 2005; **80**:1089-1096.
- Kubota F, Goto M. *Res Develop Jpn*, 2006; **13**:23-36.
- Dupont J, Suarez PAZ. *Phys Chem Chem Phys*, 2006; **8**:2441-2452.
- Jorapur YR, Chi DY. *Bull. Korean Chem Soc*, 2006; **27**:345-354.
- Polyakova Y, Koo YM, Row KH. *Biotech Bioproc Eng*, 2006; **11**:1-6.
- Abedin EISZ, Endres F. *Chem. Phys. Chem*, 2006; **7**:58-61.
- Cao Y, Zhang J, Bai Y, Li R, Zakeeruddin SM, Grätzel M, Wang PJ. *Phys Chem*, 2008; **112**:13775-13781.
- Kuang D, Klein C, Zhang Z, Ito S, Moser JE, Zakeeruddin SM, Grätzel M. *Small*, 2007; **3**:2094-2102.
- Pan X, Dai SY, Wang KJ. *Acta Phys. Chim. Sinica*, 2005; **21**:697-702.
- Gaillard C, Moutiers G, Mariet C. *ACS Symposium Series*, 2005; **902**:19-32.
- Koel M. *ACS Symposium Series*, 2005; **902**:72-82.
- Gutowski KE, Bridges NJ, Cocalia VA. *ACS Symposium Series*, 2005; **902**:33-48.
- Antonietti M, Kuang DB, Smarsly B. *Angewandte Chem In Ed*, 2004; **43**:4988-4992.
- Endres F. *Phys Chem Chem Phys*, 2004; **218**:255-283.
- Kubisa P. *Prog Polym Sci*, 2004; **29**:3-44.
- Jain N, Kumar A, Chauhan SMS. *Tetrahedron*, 2005; **61**:1015-1060.
- Yang Z, Pan WB. *Technol*, 2005; **37**:19-28.

46. Magnusson DK, Bodley JW, Adams DF. *J Solution Chem*, 1984; **13**:583-587.
47. Erbedinger M, Mesiano AJ, Russell AJ. *Biotechnol Prog*, 2000; **16**:1129-1135.
48. Schoefer SH, Kaftzik N, Wasserscheid P, Kragl U. *Chem Commun*, 2001; **5**:425-426.
49. Kim KW, Song B, Choi MY, Kim MJ. *Org Lett*, 2001; **3**:1507-1509.
50. Itoh T, Akasaki E, Kudo K, Shirakami S. *Chem Lett*, 2001; **30**:262-263.
51. Park S, Kazlauskas RJ. *J Org Chem*, 2001; **66**:8395-8401.
52. Langer R, Klausener A, Rodefeld L. *Chem. Abstr.*, 2001; **134**:85168u.
53. Kragl U, Kaftzik N, Schoefer SH, Eckstein M, Wasserscheid P, Hilgers C. *Chim Oggi*, 2001; **19**:22-24.
54. Husum TL, Jorgensen CT, Christensen MW, Kirk O. *Biocatal Biotransform*, 2001; **19**:331-338.
55. Zhao H, Malhotra SV. *Biotechnol Lett*, 2002; **24**:1257-1260.
56. Pandey S. *Analytica Chimica Acta*, 2006; **556(1)**:38-45.
57. Plechkova NV, Seddon KR. *Chemical Society Reviews*, 2008; **37(1)**:123-50.
58. Rogers RD, Seddon KR. ACS Symposium Series. Vol **818**. American Chemical Society, Washington, DC; 2002.
59. Welton T. *Chem Rev*, 1999; **99**:2071-2084.
60. Earle MJ, Esperanca JMSS, Gilea MA, Lopes JNC. *Nature*, 2006; **439**:831-834.
61. Zolfigol MA, Khazaei A, Moosavi-Zare AR, Zare A. *Appl Catal A*, 2011; **400**:70-81.
62. Zolfigol MA, Khazaei A, Moosavi-Zare AR, Zare A. *J Org Chem*, 2012; **77**:3640-3645.
63. Khazaei A, Zolfigol MA, Moosavi-Zare AR, Afsar J. *Chin J Catal*, 2013; **34**:1936-1944.
64. Parvulescu VI, Hardacre C. *Chem Rev*, 2007; **107**:2615-2665.
65. Antonietti M, Kuang D, Smarsly B, Zhou Y. *Angew Chem Int Ed*, 2004; **43**:4988-4992.
66. Visser AE, Swatloski RP, Griffin ST, Hartman DH. *Sep Sci Technol*, 2001; **36**:785-804.
67. Muzart J. *Catal*, 2006; **348**:275-295.
68. Ibrahim F, Moniruzzaman M, Yusup S, Uemura Y. *J Mol Liq*, 2015; **211**:370-372.
69. Moniruzzaman M, Ono T. *Biochem Eng J*, 2012; **60**:156-160.
70. Quental MV, Caban M, Pereira MM, Stepnowski P. *Biotechnol J*, 2014; **9**:1565-1572.
71. Moniruzzaman M, Ino K, Kamiya N, Goto M. *Org Biomol Chem*, 2012; **10**:7707-7713
72. Moniruzzaman M, Ono T. *Bioresour Technol*, 2013; **127**:132-137.
73. Financie R, Moniruzzaman M, Uemura Y. *Biochem Eng J*, 2016; **110**:1-7.
74. Elgharbawy AA, Alam MZ, Moniruzzaman M, Goto M. *Biochem Eng J*, 2016; **109**:252-267.
75. Hough WL, Smiglak M, Griffin S, Reichert WM. *New J Chem*, 2009; **33**:1429-1436.
76. Moniruzzaman M, Tahara Y, Tamura M, Kamiya N, Goto M. *Chem Commun*, 2010; **46**:1452-1454.
77. Moniruzzaman M, Goto M. *J. Chem. Eng. Jpn.*, 2011; **44**:295-314.
78. Moniruzzaman M, Tamura M, Tahara Y, Kamiya N. *Int J Pharm*, 2010; **400**:243-250.
79. Araki S, Wakabayashi R, Moniruzzaman M, Kamiya N, Goto M. *Med Chem Comm*, 2015; **23**:2124-2128.
80. Williams HD, Sahbaz Y, Ford L, Nguyen T. *Chem Commun*, 2014; **50**:1688-1690.
81. Kelemen Z, Peter-Szabo B, Sze kely E, Hollo czki O, Firaha DS, Kirchner B, Nagy J, Nyulaszi L. *Chem Eur J*, 2014; **20**:13002-13008.
82. Choi J, Benedetti TM, Jalili R, Walker A, Wallace GG, Officer DL. *Chem Eur J*, 2016; **22**:14158-14161.
83. Kelley SP, Flores LA, Shannon MS, Bara JE, Rogers RD. *Chem Eur J*, 2017; **23**:14332-14337.
84. Elia GA, Ulissi U, Mueller F, Reiter J, Tsiouvaras N, Sun YK, Scrosati B, Passerini S, Hassoun J. *Chem Eur J*, 2016; **22**:6808-6814.
85. Hu J, Diao H, Luo W, Song YF. *Chem Eur J*, 2017; **23**:8729-8735.
86. Vander Hoogerstraete T, Blockx J, de Coster H, Binnemans K. *Chem Eur J*, 2015; **21**:11757-11766.
87. Rogers RD, Seddon KR. *Science*, 2003; **302**:792-793.
88. Seddon KR. *Nat Mater*, 2003; **2**:363.
89. Bruce DW, Gao Y, Canongia Lopes JN, Shimizu K, Slattery JM. *Chem Eur J*, 2016; **22**:16113-16123.
90. Ta L, Axelsson A, Bijl J, Haukka M, Sunden H. *Chem Eur J*, 2014; **20**:13889-13893.
91. Kunov-Kruse AJ, Weber CC, Rogers RD, Myerson AS. *Chem Eur J*, 2017; **23**:5498-5508.
92. Schowner R, Elser I, Toth F, Robe E, Frey W, Buchmeiser MR. *Chem Eur J*, 2018; **24**:13336-13347.

93. Kim SY, Kim S, Park MJ. *Nat Commun*, 2010; **1**: 88.
94. Isoaari P, Srivastava V, Sillanpaa M. *Sci Total Environ*, 2019; **690**:604-619.
95. Tian F, Yang Y, Wang XL, An WL, Zhao X, Xu S, Wang YZ. *Mater Horiz*, 2019; **6**:1733-1739.
96. Tian F, Wang XL, Yang Y, An W, Zhao X, Xu S, Wang YZ. *ACS Sustainable Chem Eng*, 2020; **8**:2226-2235.
97. Liu X, Tian F, Zhao X, Du R, Xu S, Wang YZ. *Appl Surf Sci*, 2020; **529**:147151.
98. Liu X, Tian F, Zhao X, Du R, Xu S, Wang YZ. *Mater Horiz*, 2021; **8**:234-243.
99. Dang W, Kubouchi M, Sembokuya H, Tsuda K. *Polymer*, 2005; **46**:1905-1912.
100. Hanaoka T, Arao Y, Kayaki Y, Kuwata S, Kubouchi M. *Polym Degrad Stab*, 2021; **186**:109537.
101. Ma Y, Nutt S. *Polym Degrad Stab*, 2018; **153**:307-317.
102. Keith MJ, Román-Ramírez LA, Leeke G, Ingram A. *Polym Degrad Stab*, 2019; **161**: 225- 234.
103. Liu T, Zhang M, Guo X, Liu C, Liu T, Xin J, Zhang J. *Polym Degrad Stab*, 2017; **139**:20-27.
104. Lo JN, Nutt SR, Williams TJ. *ACS Sustainable Chem Eng*, 2018; **6**:7227-7231.
105. Kuang X, Zhou Y, Shi Q, Wang T, Qi HJ. *ACS Sustainable Chem Eng*, 2018; **6**:9189-9197.
106. Dupont J. 2011; **44(11)**: 1223-1231.
107. Hough-Troutman W L, Smiglak M, Griffin S, Reichert W M, Mirska I, Jodynis-Liebert J, Pernak J. *New Journal of Chemistry*, 2009; **33(1)**:26-33.
108. Berthod A, Carda-Broch S. *J Liq Chromatogr Relat Technol*, 2003; **26**:1493-1508.
109. Marszall MP, Kaliszan R. *Anal Chem*, 2007; **37**:127-140.
110. Merike V, Mihkel K, Mihkel K. *Electrophoresis*, 2002; **23**:426-430.
111. Yanes EG, Gratz SR, Baldwin MJ, Robison SE, Stalcup AM. *Anal Chem*, 2001; **73**:3838-3844.
112. López-Pastor M, Simonet BM, Lendl B, Valcárcel M. *Electrophoresis*, 2008; **29**:94- 107.
113. Yue ME, Shi YP. *J Sep Sci*, 2006; **29**: 272-276.
114. François y, Varenne A, Juillerat E, Servais AC, Chiap P, Gareil P. *J Chromatogr A*, 2007; **1138**:268-275.
115. Ventura SP, Fa ES, Quental MV, Mondal D, Freire MG, Coutinho JA. *Chem Rev*, 2017; **117**:6984.
116. Zhao CN, Zhang JJ, Li Y, Meng X, Li HB. *Molecules*, 2018; **23**:2498.
117. Chemat F, Vian MA, Cravotto G. *Int J Mol Sci*, 2012; **13**:8615.
118. Ibañez E, Cifuentes A. *Green Extraction Techniques: Principles, Advances and Applications*. Amsterdam: Elsevier; 2017.
119. Soquetta MB, Terra LM, Bastos CP. *CyTAJ Food*, 2018; **16**:400. DOI: 10.1080/19476337.2017.1411978.
120. Chemat F, Vian MA. *Alternative Solvents for Natural Products Extraction, Green Chemistry and Sustainable Technology*. Berlin Heidelberg: Springer Verlag; 2014. DOI: 10.1007/978-3-662-43628-8.
121. Torres-Valenzuela LS, Ballesteros-Gomez A, Rubio S. *Food Eng Rev*, 2020; **12**:83.
122. Nkuku CA, LeSuer RJ. *Journal of Physical Chemistry*, 2007; **11**:13271.
123. Zhao H, Baker GA. *Journal of Chemical Technology & Biotechnology*, 2013; **88**:3.
124. Lourus CLS, Cláudio AFM, Neves CMSS, Freire MG. *Int J Mol Sci*, 2010; **11**:1777-1791.
125. Kohno Y, Saita S, Murata K, Nakamura N. *Polym Chem*, 2011; **2**:862–867.
126. Zafarani-Moattar MT, Hamzehzadeh S. *Biotechnol Progr*, 2011; **27**:986-997.
127. Patel R, Meena K, Abbul BK. *Appl Biochem Biotechnol*, 2014; **172**:3701-3720.
128. Taha M, Almeida MR, Domingues P, Ventura SP. *Chem Eur J*, 2015; **21**:4781-4788.
129. Yamaguchi S, Yamamoto E, Mannen T, Nakamune T. *Biotechnol J*, 2013; **8**:17-31.
130. Pereira MM, Pedro SN, Quental MV, Lima AS. *J Biotechnol*, 2015; **206**:17-25.
131. Rogers TL, Nelsen AC, Sarkari M, Young TJ. *Pharm Res*, 2003; **20**:485-493.
132. Pei Y, Wang J, Wu K, Xuan X. *Sep Purif Technol*, 2009; **64**:288-295.
133. Erbedinger M, Mesiano AJ, Russel AJ. *Biotechnol Progr*, 2000; **16**:1129-1131. 134.
134. Guncheva M, Paunova K, Yancheva D, Svinyarov I, Bogdanov M. *J Mol Catal B Enzym*, 2015; **117**:62-68.
135. Li X, Zhang C, Li S, Huang H. *Ind Eng Chem Res*, 2015; **54**:8072-8079.
136. Olkiewicz M, Caporgno MP, Font J, Legrand J. *Green Chem*, 2015; **17**:2813-2824.
137. Munoz PA, Correa-Llantén DN, Blamey JM. *Lipids*, 2015; **50**:49-55.

138. Moniruzzaman M, Kamiya K, Goto M. *Langmuir*, 2009; **25**:977- 982.
139. Noritomi H, Suzuki K, Kikuta M, Kato S. *Biochem Eng J*, 2009; **47**:27-30.
140. Attri P, Venkatesu P, Kumar A. *Phys. Chem Chem Phys*, 2011; **13**:2788-2796.
141. Geng F, Zheng LQ, Yu L, Li GZ. *Process Biochem*, 2010; **45**:306-311.
142. Fujita K, MacFarlane DR, Forsyth M, Yoshizawa-Fujita M. *Biomacro-molecules*, 2007; **8**:2080-2086.
143. Noritomi H, Minamisawa K, Kamiya R, Kato S. *J Biomed Sci Eng*, 2011; **4**:94-99.
144. Park S, Kazlauskas RJ. *J Org Chem*, 2001; **66**:8395-8401.
145. Lozano P, de Diego T, Carrie D, Vaultier M, Iborra JL. *Biotechnol Lett*, 2001; **23**:1529-1533.
146. Kim KW, Song B, Choi MY, Kim MJ. *Org Lett*, 2001; **3**:1507-1509.
147. Schofer SH, Kaftzik N, Wasserscheid P, Kragl U. *Chem Commun*, 2001:425-426.
148. Itoh T, Akasaki E, Kudo K, Shirakami S. *Chem Lett*, 2001; **30**:262-263.
149. Lau RM, van Rantwijk F, Sheddon KR, Sheldon RA. *Org Lett*, 2000; **2**:4189-4191.
150. Nara SJ, Harjani JR, Salunkhe MM. *Tetrahedron Lett*, 2002; **43**:2979- 2982.
151. Nara SJ, Harjani JR, Salunkhe MM, Mane AT, Wadgaonkar PP. *Tetrahedron Lett*, 2003; **44**:1371-1373.
152. Erbedinger M, Mesiano AJ, Russell AJ. *Biotechnol Prog*, 2000; **16**:1129-1131.
153. Lozano P, de Diego T, Guegan JP, Vaultier M, Iborra JL. *Biotechnol Bioeng*, 2001; **75**:563-569.
154. Eckstein M, Sesing M, Kragl U, Adlercreutz P. *Biotechnol Lett*, 2002; **24**:867-87
155. Laszlo JA, Compton DL. *Biotechnol Bioeng*, 2001; **75**:181-186.
156. Lozano P, De Diego T, Carrie D, Vaultier M, Iborra JL. *J Mol Catal B Enzym*, 2003; **21**:9-13.
157. Persson M, Bornscheuer UT. *J Mol Catal B Enzym*, 2003; **22**:21-27.
158. Kaftzik N, Wasserscheid P, Kragl U. *Org Proc Res Dev*, 2002; **6**:553-557.
159. Howarth J, James P, Dai JF. *Tetrahedron Lett*, 2001; **42**:7517-7519.
160. Hinckley G, Mozhaev VV, Budde C, Khmelnitsky YL. *Biotechnol Lett.*, 2002; **24**:2083-2087.
161. Laszlo JA, Compton DL. *J Mol Catal B Enzym*, 2002; **18**:109-120.
162. Zhao H, Malhotra SV. *Biotechnol Lett*, 2002; **24**:1257-1260.
163. Jiang W, Zhu WS, Li HP, Wang X, Yin S, Chang YH, Li H. *Fuel*, 2015; **140**(3):590- 596.
164. Dharaskar SA, Wasewar KL, Varma MN, Shende DZ, Yoo CK. *Procedia Eng*, 2013; **51**:314-317.
165. Robert JM, Ashlie M, John N, John S, Jeff Y. *The Royal Society of Chemistry*, 1994; 3941-3994.
166. Wojciech C, Dwight R, Malcolm BJ. *Cellulose*, 2004; 403-411.
167. Tsiptsias C, Stefopoulos A, Kokkinomalis I, Papadopoulou L, Panayiotou C. *Green Technology*, 2008; 965-971.
168. Isogai A, Atalla RH. *Cellulose*, 1998; 309-319.
169. Kohler N, Sun C, Wang J, Zhang M. *Langmuir*, 2005; **21**(19):8858-8864.
170. Bertorelle F, Wilhelm C, Roger J, Gazeau F, Ménager C, Cabuil V. *Langmuir*, 2006; **22**(12):5385-5391.
171. Manojkumar K, Sivaramakrishna A, Vijayakrishna K. *Journal of Nanoparticle Research*, 2016; **18**(4):1-22.
172. Ding K, Miao Z, Liu Z, Zhang Z, Han B, An G, Miao S, Xie Y. *Journal of the American Chemical Society*, 2007; **129**(20):6362-6363.
173. Zheng W, Liu X, Yan Z, Zhu L. *ACS Nano*, 2008; **3**(1):115-122.
174. Alavi-Tabari SA, Khalilzadeh MA, Karimi-Maleh H. *Journal of Electroanalytical Chemistry*, 2018; **811**:84-88.
175. Tshemese Z, Mlowe S, Revaprasadu N, Deenadayalu N. *Materials Science in Semiconductor Processing*, 2017; **71**: 258-262.
176. Tshemese Z, Khan MD, Mlowe S, Revaprasadu N. *Materials Science and Engineering B*, 2018; **227**:116-121.
177. Sotolongo AC, Martinis EM, Wuilloud RG. *Analytical Methods*, 2018; **10**:338-346.
178. Pechtl MH, Campbell PS, Scholten JD, et al. *Nanoscale*, 2010; **2**(12):2601-2606.
179. Okoli CU, Kuttiyiel KA, Cole J, et al. *Ultrasonics Sonochemistry*, 2018; **41**:427-434.
180. Sajjadi H, Modaresi A, Magri P, et al. *Journal of Molecular Liquids*, 2013; **186**:1-6.
181. Biswas K. Rao CeNeR. *Chemistry A European Journal*, 2007; **13**(21):6123-6129.
182. Zhao XL, Wang CX, Hao XP, Yang JX, et al. *Materials Letters*, 2007; **61**(26):4791-4793.

183. Migowski P, Machado G, Texeira SR, et al. *Physical Chemistry Chemical Physics*, 2007; **9(34)**:4814-4821.
184. Redel E, Walter M, Thomann R, et al. *Chemistry A European Journal*, 2009; **15(39)**:10047-10059.
185. Fonseca GS, Machado G, Teixeira SR, Fecher GH, et al. *Journal of Colloid and Interface Science*, 2006; **301(1)**:193-204.
186. Holbrey JD, Plechkova NV, Seddon KR. In *Green Chemistry*; Royal Society of Chemistry: London, UK, 2006; **8**:411-414.
187. Abbott AP, Davies DL, Capper G, Rasheed RK, Tambyrajah V. *Ionic Liquids and Their Use*. WO2002026381A2, 4 April 2002.
188. Abbott AP, Davies DL, Capper G, Rasheed RK, Tambyrajah V. *Ionic Liquids and Their Use as Solvents*. WO2002026701A2, 4 April 2002.
189. Abbott AP, Capper G, Davies DL, Rasheed RK, Archer J, John C. *Trans Inst Me Finish*, 2004; **82**:14-17.
190. Abbott AP, Capper G, Davies DL, Rasheed RK. *Chem A Eur J*, 2004; **10**:3769-3774.
191. Professor Andrew P Abbott—University of Leicester, Available online: https://www2.le.ac.uk/departments/chemistry/people/academic-staff/andrew_p_abbott, accessed on 10 June 2020.
192. Runge W. *Supplement to the Treatise. Technology Entrepreneurship: A Treatise on Entrepreneurs and Entrepreneurship for and in Technology Ventures*. German: Karlsruhe, KIT Scientific Publishing; 2014.
193. NOHMs Technologies Inc, Available online: <http://www.nohms.com/technology/>, accessed on 10 June 2020.
194. Moganty SS, Lee J. Hybrid Ionic Liquid Electrolytes. US20160164137A1, 9 June 2016.
195. Zyga L. Metal-Air Battery Could Store 11 Times More Energy than Lithium-Ion. *Phys Org*, 2009. Available online: <https://phys.org/news/2009-11-metal-air-battery-energy-lithium-ion.html>, accessed on 10 June 2020.
196. Friesen CA, Wolfe D, Johnson PB. Metal-Air Cell with Ion Exchange Material. WO2012174558A1, 20 December 2012.
197. Tullo AH. *CEn Glob Enterp*, 2017; **95**:21-22.
198. Friesen CA, Krishnan R, Tang T, Wolfe D. Metal-Air Cell with Tuned Hydrophobicity. WO2011159391A1, 22 December 2011.
199. Vagt U. *CHEManager Eur*, 2008; **11**:8.
200. Kalb RS. Toward Industrialization of Ionic Liquids. In: Shiflett MB, Ed. In *Commercial Applications of Ionic Liquids*. Switzerland: Springer Cham; 2020. p. 261-282.
201. ZapGo Ltd. Carbon-Ion™: A new, safer and faster charging category of rechargeable energy storage devices. In Brochure; ZapGo: Oxford, UK, 2016; 1-16.
202. IoLiTec, Electrolytes for Metal Deposition, Available online: <https://iolitec.de/node/652>, accessed on 10 June 2020.
203. Meara O, Alemany M, Maase A, Vagt M, Malkowsky UI. *Met Finish*, 2009; **107**:38-39.
204. Alemany A, Malkowsky I, Vagt U, Maase MO, Meara M. *Plat Surf Finish*, 2010; **97**:34-37.
205. Freydina E, Abbott JG, Lund AC, Hilty RD, et al. Nanostructured Aluminum Alloys for Improved Hardness. US10590558B2, 17 March 2020.
206. CORDIS European Commission. Final Report Summary—SCAIL-UP (Scaling-up of the Aluminium Plating Process from Ionic Liquids), 2017. Available online: <https://cordis.europa.eu/project/id/608698/reporting>, accessed on 25 August 2020.
207. Tullo AH. *Chem Eng News*, 2020; **98**:24.
208. McCoy M. *Chem Eng News*, 2016; **94**:16.
209. Timken HK, Luo H, Chang BK, Carter E, Cole M. ISOALKY™ Technology: Next-Generation Alkylate Gasoline Manufacturing Process Technology Using Ionic Liquid Catalyst. In: Shiflett MB, Ed. In *Commercial Applications of Ionic Liquids*. Switzerland: Springer Cham; 2020. p. 33-47.
210. Seddon K. *Green Chem*, 1999; **1**:G58.
211. Timken HKC, Elomari S, Trumbull S, Cleverdon R. Integrated Alkylation Process Using Ionic Liquid Catalysts. WO2006073749A2, 13 July 2006.
212. Elomari S, Trumbull S, Timken HKC, Cleverdon R. Alkylation Process Using Chloroaluminate Ionic Liquid Catalysts. WO2006068983A2, 29 June 2006.
213. Martins SC, Nafis DA, Bhattacharyya A. Alkylation Process Using Phosphonium-Based Ionic Liquids. US9156028B2, 13 October 2013.
214. Holbrey JD, Plechkova NV, Seddon KR. Recalling COIL. In *Green Chemistry*; Royal Society of Chemistry: London, UK, 2006; **8**:411-414.
215. Freemantle M. *Chem Eng News*, 2005; **83**:33-38.

216. Abai M, Atkins MP, Hassan A, Holbrey JD, et al. *Dalt Trans*, 2015; **44**:8617-8624.
217. Boada R, Cibin G, Coleman F, et al. *Dalt Trans*, 2016; **45**:18946-18953.
218. Rogers RD, Holbrey J, Rodriguez H. Process for Removing Metals from Hydrocarbons. WO2010116165A2, 14 October 2010.
219. Abai M, Atkins MP, Cheun KY, et al. Process for Removing Metals from Hydro-carbons. WO2012046057A2, 12 April 2012.
220. Cheun KY, Holbrey JD, Atkins MP. Process for Removing Heavy Metals from Hydrocarbons. WO2016139280A1, 9 September 2016.
221. Werner S, Szesni N, Kaiser M, Haumann M, Wasserscheid P. *Chem Eng Technol*, 2012; **35**:1962-1967.
222. Kalb RS. Toward Industrialization of Ionic Liquids. In: Shiflett MB, editor. In Commercial Applications of Ionic Liquids. Switzerland: Springer Cham; 2020. p. 261–282.
223. Clariant Specialty Chemicals, Available online: <https://www.clariant.com/en/Corporate>, accessed on 28 August 2020.
224. Szesni N, Fischer R, Hagemeyer A, Grossmann F, Hou HC, Boyer J, Sun M, Urbancic M, Lugmair C, Lowe DM. Catalyst Composition for Selective Hydrogenation with Improved Characteristics. WO2013057244A1, 25 April 2013.
225. Barth T, Korth W, Jess A. *Chem Eng Technol*, 2017; **40**:395-404.
226. 3MTM Antistatic Additives. Ionic Liquids and Solids; 3M: Saint Paul, MI, USA; 2016.
227. BASF. Basionics-High-Performance Antistatic Additives Brochure; BASF: Ludwigshafen, Germany; 2016.
228. Evonik Industries. Press Release, New Tego Products Increase Conductivity and Electrostatic Charge Dissipation of Paints and Coatings 2009, Available online: <https://corporate.evonik.com/en/new-tego-products-increase-conductivity-and-electrostatic-charge-dissipation-of-paints-and-coatings-101049.html>, accessed on 19 August 2020.
229. Koei Chemical Co., L, Ionic Liquids-Type Antistatic Agents for Resins. Available online: https://www.koeichem.com/en/en_product/ion/antistatic.html, accessed on 22 August 2020.
230. Hoff A, Jost C, Prodi-Schwab A, Weyershausen B, Schmidt FG. *Elem Degussa Sci Newsl*, 2004; **9**:10-15.
231. Weyershausen B, Lehmann K. *Green Chem.*, 2005; **7**:15-19.
232. Iolitec, Ionic Liquids Technologies, Available online: <https://iolitec.de/>, accessed on 23 July 2020.
233. CORDIS European Commission, Final Report Summary—IOLICAP (Novel IONic Liquid and Supported Ionic Liquid Solvents for Reversible CAPture of CO₂), Available online: <https://cordis.europa.eu/project/id/283077/reporting>, accessed on 17 August 2020.
234. Natural Fiber Welding, Inc. Available online: <https://naturalfiberwelding.com/>, accessed on 25 August 2020.
235. DeLong HC, Trulove PC, Haverhals LM, Reichert WM. Natural Fiber Welding. US8202379B1, 19 June 2012.
236. Haverhals LM, Reichert WM, De Long HC, Trulove PC. *Mater Eng*, 2010; **295**:425-430.
237. Haverhals LM, Durkin DP, Trulove PC. Natural Fiber Welding. In: Shiflett MB, Ed. In Commercial Applications of Ionic Liquids. Switzerland: Springer Cham; 2020. p. 211-226.
238. Geyer R, Jambeck JR, Law KL. *Sci Adv*, 2017; **3**:1-5.
239. Peterson ML, Hickey MB, Zaworotko MJ, Almarsson O. *J Pharm Pharm Sci*, 2006; **9**:317-326.
240. Morissette SL, Almarsson Ö, Peterson ML, et al. *Adv. Drug Deliv Rev*, 2004; **56**:275–300.
241. Singhal D, Curatolo W. *Adv. Drug Deliv Rev*, 2004; **56**:335-347.
242. Datta S, Grant D. *Nat Rev Drug Discov*, 2004; **3**:42-57.
243. Rodríguez H, Bica K, Rogers RD. *Trop J Pharm Res*, 2008; **7**:1011-1012.
244. Yang Z, Pan W. *Enzym Microb Technol*, 2005; **37**:19-28.
245. Mann JP, McCluskey A, Atkin R. *Green Chem*, 2009; **11**:785-792.
246. Byrne N, Angell CA. *J Mol Biol*, 2008; **378**:707-714.
247. Fujita K, MacFarlane D, Forsyth M, et al. *Biomacromolecules*, 2007; **8**:2080- 2086.
248. Baker SN, McCleskey TM, Pandey S, Baker GA. *Chem Commun*, 2004; 940-941.
249. Myles L, Gore R, Špulák M, Gathergood N, Cannon SJ. *Green Chem*, 2010; **12**:1157-1162.

250. Hossain MI, El-Harbawi M, Alitheen NBM, Noaman YA, Lévêque JM, Yin CY. *Ecotoxicol Environ Saf*, 2013; **87**:65-69.
251. Coleman D, Špulák M, Garcia MT, Gathergood N. *Green Chem*, 2012; **14**:1350-1356.
252. Biczak R, Bałczewski P, Bachowska B, et al. *Elem*, 2013; **188**:459-461.
253. Malhorta SV, Kumar VA. *Bioorg. Med. Chem Lett*, 2010; **20**:581-585.
254. Terrecilla JS, García J, Rojo E, Rodríguez F. *J Hazard Mater*, 2009; **164**:182-194.
255. Allen D, Sinclair-Rosselot K. *Pollution Prevention for Chemical Processes*; John Wiley: New York, NY, USA; 1997.
256. Hough-Troutman WL, Smiglak M, Griffin S, et al. *New J Chem*, 2009; **33**:26-33.
257. Demberelnyamba D, Kim KS, Choi SJ, et al. *Bioorg Med Chem*, 2004; **12**:853-857.
258. Carson L, Chau PKW, Earle MJ, et al. *Green Chem*, 2009; **11**:492-497.
259. Stoimenovski J, MacFarlane DR, Bica K, Rogers RD. *Pharm Res*, 2010; **27**:521-526.
260. Hough W, Smiglak M, Rodríguez H, et al. *New J Chem*, 2007; **31**:1429-1436.
261. Matzke M, Stolte S, Thiele K, et al. *Green Chem*, 2007; **9**:1198-1207.
262. Yu M, Li SM, Li XY, Zhang BJ, Wang JJ. *Ecotoxicol Environ Saf*, 2008; **71**:903-908.
263. Murshe SMS, Nieto de Castro CA. *Nanofluids: Synthesis, Properties and Applications*; Nova Science Publishers Inc: New York, NY, USA; 2014.
264. Nieto de Castro CA, Vieira SIC, Lourenço MJ, Murshed SM. *J Nanofluids*, 2017; **6**:804-811.
265. Cacia K, Murshed SMS, Pabón E, Buitrago R. *J Therm Anal Calorim*, 2020; **140**:109-114.
266. Bridges NJ, Visser AE, Fox EB. *Energ Fuel*, 2011; **25**:4862-4864.
267. Kanakubo M, Harris KR. *J Chem Eng Data*, 2015; **60**:1408-1418.
268. Liu H, Maginn E, Visser AE, Bridges NJ, Fox EB. *Ind Eng Chem Res*, 2012; **51**:7242-7254.
269. Gardas RL, Coutinho JAP. *Ind Eng Chem Res*, 2008; **47**:5751- 5757.
270. Papovic S, Bešter-Roga M, Vraneš M, Gadzuric S. *J Chem Thermodyn*, 2016; **99**:1-10.
271. Safarov J, Hamidova R, Zepik S, et al. *J Mol Liq*, 2013; **187**:137-156.
272. Nieto de Castro CA, Murshed SMS, Lourenço MJV, Santos FJV, Lopes MLM, Franç JMP. *Int J Therm Sci*, 2012; **62**:34-39.
273. Páez MS, Vega YA, Romero CM. *J Mol Liq*, 2017; **243**:78-84.
274. Nieto de Castro CA, Lourenço MJV, Ribeiro APC, Langa E, et al. *J Chem Eng Data*, 2010; **55**:65-69.
275. Paulechka YU, Kabo AG, Blokhin AV, Kabo GJ, Shevelyova MP. *J Chem Eng Data*, 2010; **55**:2719-2724.
276. Ribeiro APC, Vieira SIC, França JMP, Queirós CS, Langa E, Lourenço MJV, Murshed SMS, Nieto de Castro CA. *Thermal properties of ionic liquids and ionanofluids*. In *Ionic Liquids: Theory, Properties, New Approaches*. Kokorin A, Ed. InTech: Rijeka Croatia; 2011.
277. Li JG, Hu YF, Ling S, Zhang JZ. *J. Chem Eng Data*, 2011; **56**:3068-3072.
278. Oster K, Hardacre C, Jacquemin J, Ribeiro APC, Elsinawi. *Aust J Chem*, 2018; **72**:21-33.
279. Navarro P, Larriba M, Rojo E, García J, Rodríguez F. *J Chem Eng Data*, 2013; **58**:2187-2193.
280. Fröba AP, Kremer H, Leipertz A. *J Phys Chem B*, 2008; **112**:12420-12430.
281. França JMP, Vieira SIC, Lourenço MJV, Murshed SMS, Nieto de Castro CA. *J. Chem Eng Data* 2013; **58**:467-476.
282. Ficke LE, Rodríguez H, Brennecke JF. *J Chem Eng Data*, 2008; **53**:2112-2119.
283. Franca JMP. *Solid-Liquid Interactions in Ionanofluids. Experiments and Molecular Simulation*. Ph.D. Thesis, Universidade de Lisboa, Lisboa, Portugal, Université Clermont, Auvergne, France; 2017.
284. Jó'zwiak B, Dzido G, Zorebski E, et al. *ACS Appl Mater Interfaces*, 2020; **12**:38113-38123.
285. Wu JY, Chen YP, Su CS. *Chem Eng*, 2014; **45**:2205-2211.
286. Vallejo JP, Liñeira del Río JM, Fernández J, Lugo L. *J Mol Liq*, 2020; **319**:114335.
287. Chereches EI, Prado JI, Chereches M, Minea AA, Lugo L. *J Mol Liq*, 2019; **291**:111332.
288. Dakkach M, Gaciño FM, Guimarey MJG, et al. *J Chem Thermodyn*, 2018; **218**:27-38.
289. Massel M, Revelli AL, Paharik E, Rauh M, Mark LO, Brennecke JF. *J Chem Eng Data*, 2015; **60**:65-73.
290. Paul TC, Morshed AKMM, Fox EB, Khan JA. *Appl Therm Eng*, 2017; **110**:1-9.

291. Zarrougui R, Dhahbi M, Lemordant D. *J Solut Chem*, 2010; **39**:921-942.
292. Wang F, Han L, Zhang Z, Fang X, Shi J, Ma W. *Nanoscale Res Lett*, 2012; **7**:314-319.
293. Liu J, Wang F, Zhang L, Fang X, Zhang Z. *Renew Energy*, 2014; **63**:519-523.
294. Fillion JJ, Brennecke JF. *J Chem Eng Data*, 2017; **62**:1884-1901.
295. Ferreira AGM, Simoes PN, Ferreira AF, Fonseca MA, Oliveira MSA, Trino ASM. *J Chem Thermodyn*, 2013; **64**:80-92.
296. Xie H, Zhao Z, Zhao J, Gao H. *Chin J Chem Eng*, 2016; **24**:331-338.
297. Ficke LE, Novak RR, Brennecke JF. *J Chem Eng Data*, 2010; **55**:4946-4950.
298. Patil VS. Ionic Liquids with and without Transition Metal Nanoparticles for Advanced Heat Transfer Fluids and Catalysis Applications. Ph.D. Thesis, Universitat Rovira I Virgili, Tarragona, Spain; 2017.
299. Zhang M, Kamavaram V, Reddy RG. *J Phase Equilib Diff*, 2005; **26**:124-130.
300. Pamies R, Aviles MD, Arias-Pardilla J, Espinosa T, Carrion FJ, Sanes J, Bermúdez MD. *Tribol Int*, 2018; **122**:200-209.
301. Stoppa A, Buchner R, Hefter G. *J Mol Liq*, 2010; **153**:46-51.
302. Paulechka YU, Blokhin AV. *Thermodynamics*, 2014; **79**:94-99.