


 Cite this: *New J. Chem.*, 2021, 45, 53

Effects of repeat unit charge density on the physical and electrochemical properties of novel heterocationic poly(ionic liquid)s†

 Merlin Cotessat,^a Dimitri Flachard,^b Daniil Nosov,^{id}^a Elena I. Lozinskaya,^{id}^c Denis O. Ponkratov,^c Daniel F. Schmidt,^a Eric Drockenmuller^{id}*^a and Alexander S. Shaplov^{id}*^a

We report the synthesis and structure/property correlations of a series of eight poly(ionic liquid)s (PILs) obtained from sequential AA + BB polyaddition by copper(i)-catalyzed azide–alkyne cycloaddition and the subsequent *N*-alkylation reaction. The different repeat units contain one to four ion pairs, with one to four bis(trifluorosulfonyl)imide (TFSI) anions and one or two types of ammonium, imidazolium or 1,2,3-triazolium counter-cations. Their physical, ion conducting and electrochemical properties are discussed based on the repeat unit charge density and structure of the cationic moieties. The comparison of several pairs of polyelectrolytes revealed that ionic conductivity is dependent on (1) the ratio between the number of charge carriers per monomer unit and the number of surrounding atoms/groups that can solvate the ions and (2) the nature of the cation. The highest conductivity ($1.8 \times 10^{-5} \text{ S cm}^{-1}$ at 25 °C) was reached when PILs contain two 1,2,3-triazolium cations that are separated by an oxyethylene spacer. The incorporation of an additional type of cation (either imidazolium or ammonium with 1,2,3-triazolium) in one PIL allows its cathodic limit to be increased up to -0.4 V vs. Li^+/Li (70 °C) and the oxidation instability of 1,2,3-triazolium cations to be overcome.

 Received 17th August 2020,
 Accepted 20th October 2020

DOI: 10.1039/d0nj04143b

rsc.li/njc

Introduction

In the last decade, intensive efforts focused on flexible, lightweight and safe energy storage devices have been stimulated by the increased demand of next-generation portable and flexible electronics, wearable devices and electric cars. Both flexible solid-state supercapacitors (SCs)^{1,2} and solid-state Li batteries^{3–6} have attracted increased interest given the convincing advantages of transitioning from volatile, flammable and toxic liquid electrolytes to lightweight, non-volatile polymer electrolytes. In comparison with the existing technologies, solid-state SCs and Li batteries are smaller in size, light weight, and easy to handle, and offer a broader range of operating temperatures, and greater safety and reliability.^{1,3,4,7,8} The

development of safe, efficient solid-state SCs and lithium batteries requires solid polymer electrolytes (SPEs) with high ionic conductivity ($>10^{-4} \text{ S cm}^{-1}$ at 25 °C) as well as good electrochemical and thermal stabilities.^{3,9}

Among promising candidates for SPEs, a new class of polyelectrolytes, namely polymeric analogs of ionic liquids, a.k.a. poly(ionic liquid)s (PILs), has recently gained significant attention.^{10–17} PILs can be considered as the macromolecular analogs of ionic liquids (ILs), in the best cases combining the advantages of polymers (viscoelasticity, processability, film-forming properties, *etc.*) and ILs (high thermal and electrochemical stabilities, enhanced ionic conductivity, *etc.*).^{10,11,15} The use of PILs in SPEs instead of common neutral polymers offers several advantages. Firstly, PILs being polyelectrolytes, impart additional charges to the system. Secondly, when ILs are added to PILs, the optimal homogeneous distribution of charges can be reached at the molecular level due to strong electrostatic interactions, thus minimizing phase separation, IL leakage and providing the possibility of increasing the IL content up to 70–90 wt%.^{9,18}

Nowadays a broad variety of PILs are prepared using countless combinations of cations (ammonium, pyridinium, imidazolium, phosphonium, 1,2,3- and 1,2,4-triazoliums, *etc.*) and anions (halides, perfluorinated sulfonimides, *etc.*).^{10,19–22} An analysis of

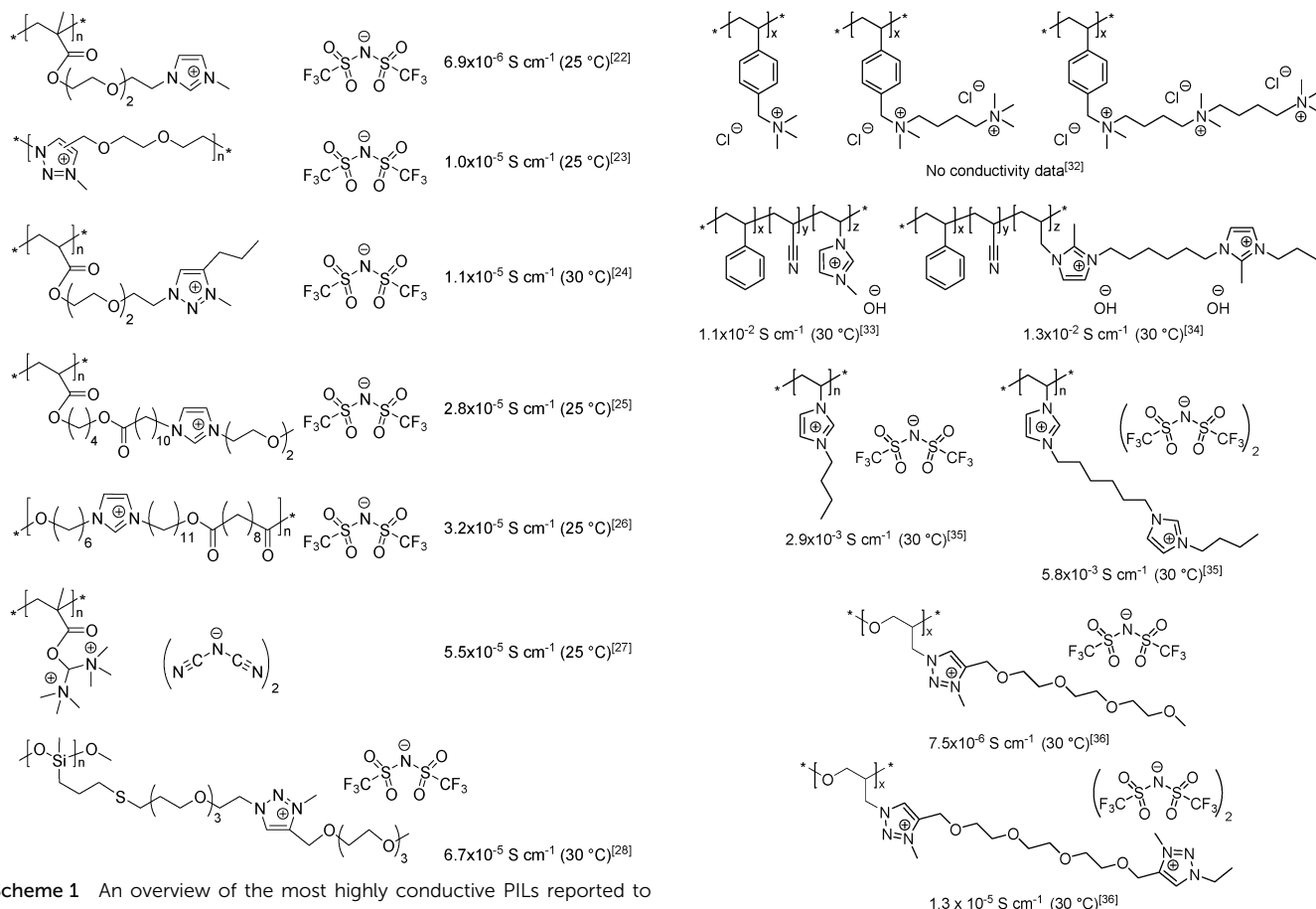
^a Luxembourg Institute of Science and Technology (LIST),
 5 avenue des Hauts-Fourneaux, L-4362 Esch-sur-Alzette, Luxembourg.
 E-mail: alexander.shaplov@list.lu

^b Univ. Lyon, Université Lyon 1, CNRS, Ingénierie des Matériaux Polymères,
 UMR 5223, F-69003, Lyon, France. E-mail: eric.drockenmuller@univ-lyon1.fr

^c A. N. Nesmeyanov Institute of Organoelement Compounds Russian Academy of
 Sciences (INEOS RAS), Vavilov str. 28, 119991, Moscow, Russia

† Electronic supplementary information (ESI) available: ¹H, ¹³C and ¹⁹F NMR and FT-IR spectra, and DSC and TGA traces of ionic monomers and PILs. See DOI: 10.1039/d0nj04143b

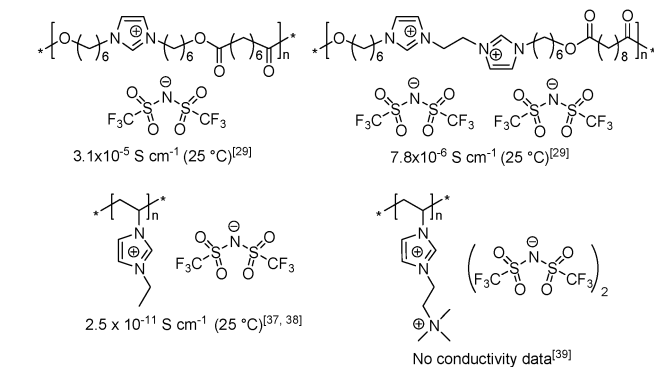




Scheme 1 An overview of the most highly conductive PILs reported to date.

the published data allows the identification of the main factors influencing the bulk conductivity of PILs: the structure of the cation and anion, the length and type of the spacer between the polymer backbone and chemically grafted ion, the glass-transition temperature and the polymerization degree.^{3,10–17,23,24} It should be noted that all of the aforementioned factors are interrelated and that it is their combination that governs the ionic conductivity of PILs. The structures of the most highly conductive PILs synthesized to date^{25–31} are listed in Scheme 1; the maximum bulk ionic conductivity reported in a neat, well-purified linear PIL is $6.7 \times 10^{-5} \text{ S cm}^{-1}$ at 30 °C under anhydrous conditions.³¹

Although attempts to enhance the ionic conductivity of PILs have been made by varying the anion and cation structures, the length of the spacer and the placement of the ions in the polymer backbone *vs.* in the side chain, the influence of repeat unit charge density on the bulk conductivity of PILs has been scarcely addressed.^{29,32–39} As seen in Scheme 2, the maximum number of ion pairs per monomer unit reported to date is three,³² and the structure/conductivity relationships remain sometimes contradictory. For instance, comparable hydroxide conductivities were reported for imidazolium-based PILs having cationic or dicationic repeat units and hydroxide counter-anions.^{33,34} Conversely, Fan *et al.*³⁵ and Drockenmüller *et al.*³⁶ observed a *ca.* 2 fold increase in ionic conductivity for PILs



Scheme 2 Examples of the repeat unit charge density influence on the ionic conductivity of PILs.

carrying one and two pendent imidazolium or 1,2,3-triazolium cations in each repeating unit (Scheme 2). In total contradiction, Gibson *et al.* observed a fourfold decrease in ionic conductivity for polyesters when the number of main-chain imidazolium cations was increased from one to two.²⁹ Finally, while extremely low ionic conductivity was reported for poly(1-vinyl-3-ethylimidazolium) bis(trifluoromethylsulfonyl)-imide,^{37,38} the ionic conductivity of the only example of heterodicationic PIL reported so far (*i.e.* poly(*N,N,N*-trimethyl)-*N*-(1-vinylimidazolium-3-ethyl)-ammonium bis(trifluoromethyl-sulfonyl)imide³⁹) has not been reported.



To fill this gap in fundamental knowledge concerning the influence of repeat unit charge density on the bulk ionic conductivity of SPEs, in this work we report the synthesis and comparison of eight PILs obtained from sequential AA + BB polyaddition by copper(I)-catalyzed azide-alkyne cycloaddition (CuAAC) using ammonium or imidazolium dialkynes with α,ω -diazido-tetra(ethylene glycol) and subsequent *N*-alkylation of the 1,2,3-triazole groups. This versatile and general approach afforded the preparation of a series of PILs carrying TFSI counter-anions and differing by the chemical nature and number of cations in the repeat unit (Scheme 4). Their physical and electrochemical properties are discussed based on the repeat unit charge density as well as on the combination and chemical nature of the cations. These novel PILs exhibit conductivity up to $1.6 \times 10^{-5} \text{ S cm}^{-1}$ at 25 °C as well as electrochemical stability up to 3.6 V vs. Li⁺/Li.

Results and discussion

The analysis of Schemes 1 and 2 led us to the conclusion that the synthesis of (meth)acrylate or siloxane monomers with multiple ion pairs would require advanced multistep synthetic procedures. In contrast, synthesis of 1,2,3-triazolium-based PILs (TPILs) through AA + BB CuAAC polyaddition and subsequent quantitative *N*-alkylation of poly(1,2,3-triazole) intermediates allows unprecedented opportunities for macromolecular design taking advantage of the appealing attributes of the “click chemistry” philosophy.¹⁴ Such an approach has been previously applied to the synthesis of aliphatic, aromatic, and ethylene glycol-based TPILs.^{14,40,41} Thus, to fulfill the purpose of this study the synthetic platform towards TPILs was selected as the most favorable approach for the synthesis and investigation of a library of monocationic and heterodicationic PILs with tunable repeat unit charge density (Scheme 4 and Table 1). 1,11-Diazido-3,6,9-trioxaundecane (**6**) was chosen as the diazide monomer due to its easy accessibility and the presence of oxyethylene units that on one hand impart flexibility to the polymer chain and on another improve the solubility of ionic species, facilitate their dissociation, and promote an increase in the ionic conductivity of the resulting PILs.¹⁰ A TFSI ion was selected as an anion providing PILs both with high ionic conductivity (Scheme 1) and electrochemical stability.⁶

Synthesis of dialkyne monomers

The first step of our investigation started with the design of dialkyne monomers. As first candidates *N,N*-dimethyl-*N,N*-dipropargylammonium bis(trifluoromethylsulfonyl)imide (**1**) and *N,N*-dibutyl-*N,N*-dipropargylammonium bis(trifluoromethylsulfonyl)imide (**2**) each having one ammonium cation were prepared as reported previously.⁴²

Dipropargyl monomer **3** having two ammonium cations and a flexible *n*-hexyl spacer in between was designed in order to increase the number of cations and to enable sufficient ion motion (Scheme 3). The synthetic strategy for the preparation of **3** included *N*-alkylation of *N,N,N',N'*-tetramethyl-1,6-hexanediamine

by propargyl bromide and an ion exchange reaction with excess LiTFSI. As a result, monomer **3** was obtained as a white crystalline solid showing a relatively low melting point ($T_m = 68\text{--}70 \text{ }^\circ\text{C}$, Fig. S15 in the ESI[†]). Monomer **4** was further developed to contain two imidazolium cations and by analogy with **3** has the same linear *n*-hexyl spacer (Scheme 3). For monomer **4** the reaction pathway contained three steps: (1) synthesis of *N*-propargyl imidazole through *N*-alkylation of imidazole with propargyl tosylate, (2) *N*-alkylation of *N*-propargyl imidazole by 1,6-dibromohexane and (3) ion exchange reaction with an excess of LiTFSI. After isolation and purification monomer **4** was obtained as a slightly viscous yellow oil. DSC analysis confirmed the glassy behavior of **4** with a single glass transition temperature ($T_g = -56 \text{ }^\circ\text{C}$) and the absence of any crystallization or melting processes (Fig. S16, ESI[†]).

The structure and purity of monomers **3** and **4** were proved by ¹H, ¹³C and ¹⁹F NMR spectroscopy, FTIR spectroscopy and elemental analysis (Fig. S1–S6, ESI[†]). The FTIR spectra of monomers **3** and **4** show absorption bands at ~ 3050 , 2990–2960, 2930 and 2880–2850 cm^{-1} that were assigned to CH₂ stretching (Fig. S3 and S6, ESI[†]). The absorption bands at 3310–3250 (ν_{CH}), 2135 ($\nu_{\text{C}=\text{C}}$) and ~ 615 (δ_{CH}) cm^{-1} were identified as characteristic bands of alkyne groups. The characteristic bands of the TFSI anion were observed at $\sim 1350\text{--}1340$ (asymmetric S=O), $\sim 1200\text{--}1190$ (CF), $\sim 1150\text{--}1140$ (symmetric S=O) and $\sim 1060\text{--}1050$ (CF) cm^{-1} . Finally, ¹⁹F NMR spectra of **3** and **4** contain a single signal at -78.7 ppm assigned to trifluoromethyl groups (Fig. S2 and S5, ESI[†]).

Synthesis of PILs

Both ionic dialkynes **1–4** and neutral dialkyne **5** were first polymerized with 1,11-diazido-3,6,9-trioxaundecane (**6**) by AA + BB CuAAC polyaddition in *N,N*-dimethylformamide (DMF) (Scheme 4). In order to obtain high molar mass PILs, the previously optimized conditions involving CuIP(OEt)₃ as a catalyst and a monomer concentration of 20 wt% were used.^{26,44,45} The change from CH₂Cl₂ to more polar DMF was governed by the solubility of the formed polymers. The application of the abovementioned reaction conditions to the polyaddition of dialkynes **1–4** with diazide **6** resulted in the preparation of PILs having one (Scheme 4, polymers **7** and **8**) or two (Scheme 4, polymers **9** and **10**) ion pairs per repeat unit.

The molar mass of the obtained polymers was estimated by size exclusion chromatography (SEC). To suppress the insufficiently charged screening or the so-called “polyelectrolyte effect” during SEC studies, an electrolyte, namely LiTFSI, was added to the polymer solution in DMF. It was revealed that in all cases CuAAC polyaddition resulted in the formation of high molar mass polymers, with number average molar masses (M_n) ranging from 50 to 120 kg mol^{-1} for PILs **8–11** (Table 1). PIL **10**, based on bisimidazolium monomer **4**, showed the highest M_n value most probably due to its better solubility (Table 1, polymer **10**).

To increase the repeat unit charge density poly(1,2,3-triazole)s **8–11** were further subjected to *N*-alkylation reaction using iodomethane and ion exchange reaction with LiTFSI to obtain the corresponding 1,2,3-triazolium-containing analogues **12–15** (Scheme 4). PIL **7** was not considered further due



Table 1 Physical properties of poly(1,2,3-triazole)s and poly(1,2,3-triazolium)s

Polymer Structure	No.	M_n^a (kDa)	D^a	T_g^b (°C)	T_{onset}^c (°C)	σ_{DC} at 25 °C (S cm ⁻¹)	Electro-chemical window ^d (V)	Charge density (mmol g ⁻¹)
	7	50.5	2.79	13	235	9.8×10^{-10}	— ^e	1.55
	8	62.8	1.94	8	220	2.8×10^{-9}	4.0	1.37
	12	19.9	1.06	15	175	4.9×10^{-10}	3.3	2.27
	9	57.4	1.77	3	270	5.8×10^{-8}	4.3	1.90
	13	56.2	1.22	8	230	8.3×10^{-9}	3.3	2.43
	10	119.0	1.82	-11	290	6.0×10^{-7}	3.9	1.82
	14	121.9	1.48	-1	265	4.2×10^{-7}	3.1	2.37
	15	71.7	2.59	-32	280	1.8×10^{-5}	4.8	2.06
	16 ^f	—	—	-35 ⁴³	—	$9.0 \times 10^{-6.43}$	—	1.97

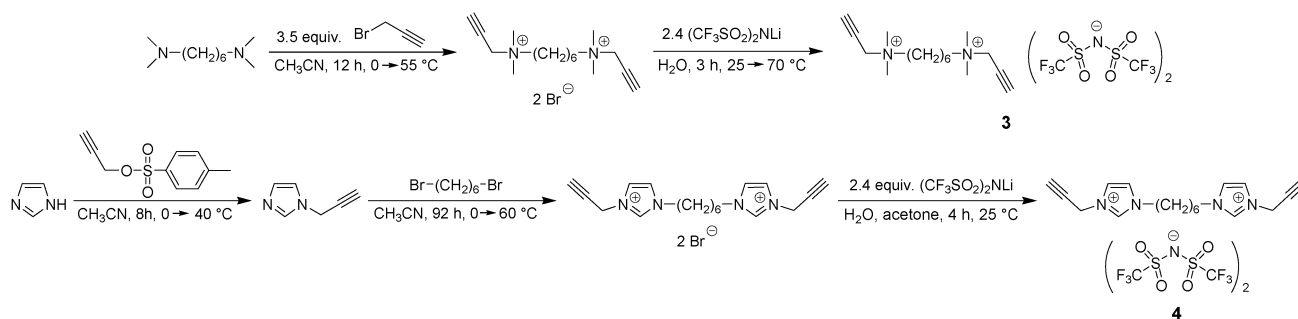
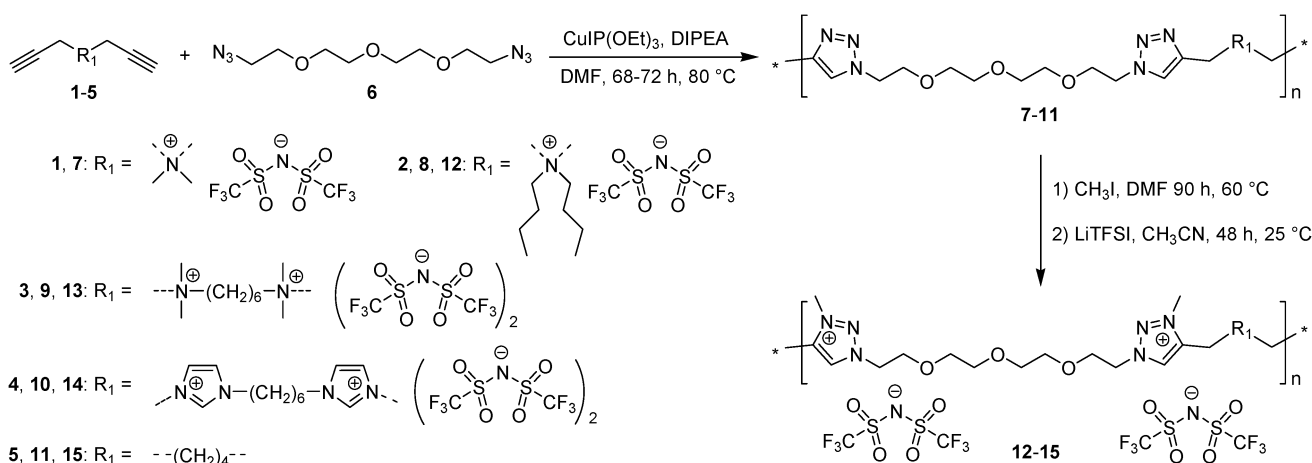
^a Obtained by GPC in 0.1 M LiTFSI in DMF at 50 °C with polystyrene standard calibration. ^b Obtained by DSC. ^c Obtained by TGA. ^d Obtained by cyclic voltammetry at 70 °C (stainless steel as a working electrode and Li foil as counter and reference electrodes, scan rate 0.5 mV s⁻¹). ^e Not determined due to the very low conductivity. ^f For comparison.

to its limited solubility that prevented the quantitative *N*-alkylation of 1,2,3-triazole groups. It should be mentioned that the previously developed single step method involving *N*-alkylation of poly(1,2,3-triazole)s **8–11** by *N*-methyl bis(trifluoromethylsulfonyl)imide²⁶ could not be performed quantitatively even after 72 h at 90 °C. Probably this can be explained by steric hindrance neighboring the 1,2,3-triazole rings due to the near proximity of an ammonium or imidazolium cation and a bulky

TFSI anion. Optimization of the reaction duration and the amount of CH₃I allowed achieving quantitative *N*-alkylation of the 1,2,3-triazole groups as confirmed by ¹H NMR. Further ion exchange reaction with excess LiTFSI in aqueous medium led to the preparation of polymers **12–15** having from two to four ion pairs per repeat unit.

PILs **8–10** and **12–15** are amorphous and readily soluble in acetone, acetonitrile and polar aprotic solvents (*i.e.* DMF,



Scheme 3 Synthesis of dicationic dipropargyl monomers **3** and **4**.Scheme 4 Synthetic pathway for the preparation of cationic PILs **7-10** and **15** and heterocationic PILs **12-14**.

DMSO, DMAc and NMP). Their structure, composition and purity were supported by ^1H , ^{13}C and ^{19}F NMR and FTIR spectroscopy (see *e.g.* Fig. S7–S14, ESI †) as well as by elemental analysis. ^1H NMR corroborated the quantitative *N*-alkylation of the 1,2,3-triazole groups in **12–15** by: (1) the appearance of the *N*-3 methyl signal at 4.34 ppm; (2) the shift from 4.55 to 4.83 ppm of the *N*-1 methylene signal; and (3) the shift from 8.22 to 8.97 ppm for the signal of the 1,2,3-triazole and 1,2,3-triazolium protons, respectively. The quantitative formation of 1,2,3-triazolium rings in **12–15** was also supported by elemental analysis. Finally, FTIR spectra of **7–10** and **12–15** showed the presence of aliphatic C–H groups (2930 and 2880 cm^{-1}) and the bands related to the TFSI anion (1346 , 1328 , 1175 , 1131 and 1050 cm^{-1}).

Thermal properties of PILs

Thermal properties of PILs **7–10** and **12–15** were assessed *via* differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA). DSC analysis revealed several interconnections between T_g and the chemical structure of PILs (*e.g.* Fig. S17 and S18 (ESI †) for polymers **10** and **14**). Firstly, the increase in the length of side chains in the ammonium cations affords a decrease in T_g (Table 1, polymers **7** and **8**). This tendency fully correlates with the melting point variation with the

carbon number in the side alkyl chain for 1-methyl-3-alkyl imidazolium ILS 46 and with the size of the side chain in 1,2,3-triazolium PILs. 43 The separation of the ammonium cations by a flexible aliphatic spacer increases the flexibility of the polymer backbone and thus reduces T_g (Table 1, polymers **7** and **9**). The transition from ammonium to imidazolium cations results in the lowering of PIL heat resistance (Table 1, polymers **9** and **10**, **13** and **14**), which was found to be in full agreement with data reported for (meth)acrylate-based PILs. 47,48 The correlation between repeat unit charge density and T_g can be summarized as follows:

$$T_g \text{ 7 (1 charge)} > T_g \text{ 13 (4 charges)} > T_g \text{ 9 (2 charges)}$$

$$T_g \text{ 12 (3 charges)} > T_g \text{ 8 (1 charge)}$$

Thus, it can be concluded that with the exception of polymer **7**, which does not contain a flexible spacer between ammonium and 1,2,3-triazole groups, the increase in repeat unit charge density induces a rise in T_g . This reduction of polymer chain mobility is most likely due to electrostatic interactions.

The thermal stability of PILs in air was found to be strongly dependent on the cation structure (Table 1 and Fig. S19, ESI †). The overall evolution of the thermal stability of PILs



(T_{onset} values) according to the cation's nature can be summarized as follows:

$$T_{\text{onset}} \mathbf{10} \text{ (imidazolium cations, } 290 \text{ }^\circ\text{C)} > T_{\text{onset}} \mathbf{15} \\ \text{(1,2,3-triazolium cations, } 280 \text{ }^\circ\text{C)} > T_{\text{onset}} \mathbf{9} \\ \text{(ammonium cations, } 270 \text{ }^\circ\text{C)}$$

As generally observed for ILs⁴⁹ and methacrylate PILs,⁴⁷ the imidazolium PIL **10** demonstrates higher thermal stability than PILs **7**, **8** and **9** having ammonium cations (Table 1). The transition to 1,2,3-triazolium cations in polymer **15** results in the lowering of the T_{onset} value in comparison with imidazolium analog **10**. The thermal stability of PILs decreased following the orders below with respect to the repeat unit charge density:

$$T_{\text{onset}} \mathbf{9} \text{ (2 charges, } 270 \text{ }^\circ\text{C)} > T_{\text{onset}} \mathbf{7} \text{ (1 charge, } 235 \text{ }^\circ\text{C)} \\ \approx T_{\text{onset}} \mathbf{13} \text{ (4 charges, } 230 \text{ }^\circ\text{C)}$$

$$T_{\text{onset}} \mathbf{8} \text{ (1 charge, } 220 \text{ }^\circ\text{C)} > T_{\text{onset}} \mathbf{12} \text{ (3 charge, } 175 \text{ }^\circ\text{C)}$$

$$T_{\text{onset}} \mathbf{10} \text{ (2 charges, } 290 \text{ }^\circ\text{C)} > T_{\text{onset}} \mathbf{14} \text{ (4 charges, } 265 \text{ }^\circ\text{C)}$$

Analyzing the thermal stability data mentioned above, it is possible to summarize that the increase in the repeat unit charge density of PILs affords a decrease in their decomposition temperatures. This is mainly due to the limited thermal stability of the 1,2,3-triazolium cation towards de-*N*-alkylation.⁵⁰

Electrochemical properties of PILs

Ionic conductivities of polymer electrolytes were investigated as a function of temperature by electrochemical impedance spectroscopy (Table 1 and Fig. 1). Depending on the chemical nature of the cations and the repeat unit charge density the σ values determined at 25 °C under anhydrous conditions for synthesized PILs varied from 9.8×10^{-10} to 1.8×10^{-5} S cm⁻¹.

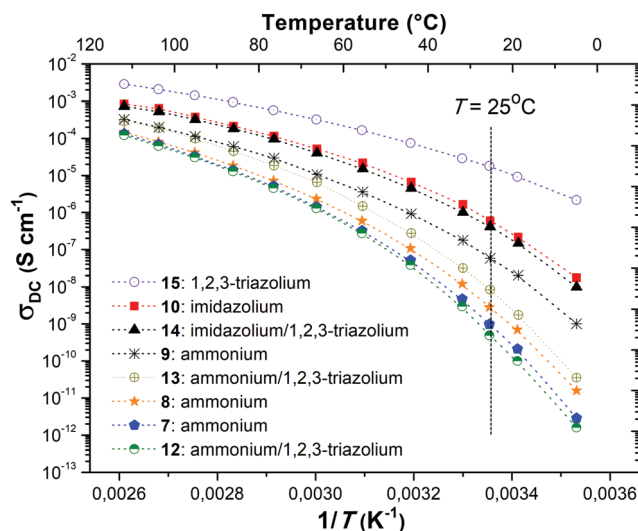


Fig. 1 Evolution of ionic conductivity (σ_{DC}) with inverse temperature for PILs **7–10** and **12–15**.

The influence of the cation chemical structure on the ionic conductivity of monocationic PILs can be ranked as follows:

$$\sigma \mathbf{15} \text{ (1,2,3-triazolium, } 1.6 \times 10^{-5} \text{ S cm}^{-1}) \gg \sigma \mathbf{10} \text{ (imidazolium, } 6.0 \times 10^{-7} \text{ S cm}^{-1}) > \sigma \mathbf{9} \text{ (ammonium, } 5.8 \times 10^{-8} \text{ S cm}^{-1})$$

This order fully correlates with the previously reported influence of cation structures on the conductivity of ILs.^{51,52} The insertion of side “dangling” chains decreases T_g and improves ionic conductivity as it can be concluded from the comparison of PILs **7** and **8** (Table 1). Such an increase in σ can be explained by the decrease in packing density of the corresponding polyelectrolyte.¹⁰ The separation of cations by an aliphatic spacer in **9** increases the solvation of ions and thus influences their dissociation and improves ionic conductivity (Table 1, polymers **7** and **9**). Finally, the influence of repeat unit charge density on the ionic conductivity of PILs can be ranked in the following decreasing orders:

$$\sigma \mathbf{9} \text{ (2 charges, } 5.8 \times 10^{-8} \text{ S cm}^{-1}) > \sigma \mathbf{13} \text{ (4 charges, } 8.3 \times 10^{-9} \text{ S cm}^{-1}) > \sigma \mathbf{7} \text{ (1 charge, } 9.8 \times 10^{-10} \text{ S cm}^{-1})$$

$$\sigma \mathbf{10} \text{ (2 charges, } 6.0 \times 10^{-7} \text{ S cm}^{-1}) > \sigma \mathbf{14} \text{ (4 charges, } 4.2 \times 10^{-7} \text{ S cm}^{-1})$$

$$\sigma \mathbf{15} \text{ (2 charges, } 1.8 \times 10^{-5} \text{ S cm}^{-1}) > \sigma \mathbf{16} \text{ (1 charge, } 9.6 \times 10^{-6} \text{ S cm}^{-1})$$

Taking these data into account it is possible to conclude that in the present work the repeat unit charge density vs. conductivity dependence passes through the maximum at PILs carrying two ion pairs. There are several possible explanations of the observed phenomenon. Possibly, the initial increase in the number of cations leads to the increase in the repeat unit charge density or in the concentration of free anions and results in the conductivity magnification. However, at some point in a solid polymer, only the surrounding atoms/groups can solvate the ions, the ratio between the number of charges and the solvating groups becomes too low for efficient dissociation and the conductivity starts to decrease. Following this assumption, the dilution of ions by incorporation of an aliphatic spacer leads to an immediate rise in ionic conductivity (Table 1, polymers **7** and **9**). Another explanation can be connected with the highly documented influence of the PIL structure/composition on T_g .¹⁰ As the increase in repeat unit charge density generally induces a rise of T_g (Table 1 and Fig. S20, ESI[†]) and as ionic conductivity has an inverse relationship with the latter,¹⁰ it becomes obvious that the increase in the number of charges will reduce the ionic conductivity of PILs. In accordance with this assumption the transfer from monocationic or dicationic PILs (Table 1, polymers **8**, **9**, and **10**) to tricationic or tetracationic ones (Table 1, polymers **12**, **13**, and **14**) is accompanied by an increase in T_g and as a result in a decrease in σ . Finally, it seems obvious that 1,2,3-triazolium being the most ion conducting cation



studied herein and in the literature,³¹ its presence systematically enhances σ , except for PIL **12**, compared to the corresponding poly(1,2,3-triazole) analogue **8**.

Additionally, the charge density of PILs was calculated as the number of charges per gram of polymeric material (Table 1 and Fig. S21, ESI†) and its influence on the ionic conductivity of PILs was estimated by the following decreasing orders:

$$\sigma \mathbf{9} (1.90 \text{ mmol g}^{-1}, 5.8 \times 10^{-8} \text{ S cm}^{-1}) > \sigma \mathbf{13} (2.43 \text{ mmol g}^{-1}, 8.3 \times 10^{-9} \text{ S cm}^{-1}) > \sigma \mathbf{7} (1.55 \text{ mmol g}^{-1}, 9.8 \times 10^{-10} \text{ S cm}^{-1})$$

$$\sigma \mathbf{10} (1.82 \text{ mmol g}^{-1}, 6.0 \times 10^{-7} \text{ S cm}^{-1}) > \sigma \mathbf{14} (2.37 \text{ mmol g}^{-1}, 4.2 \times 10^{-7} \text{ S cm}^{-1})$$

$$\sigma \mathbf{15} (2.06 \text{ mmol g}^{-1}, 1.8 \times 10^{-5} \text{ S cm}^{-1}) > \sigma \mathbf{16} (1.97 \text{ mmol g}^{-1}, 9.6 \times 10^{-6} \text{ S cm}^{-1})$$

Similarly to repeat unit charge density, the charge density *vs.* conductivity dependence passes through the maximum at an average value of 1.92 mmol g⁻¹. It can also be concluded that the ionic conductivity of PILs is mainly ruled by segmental motion and T_g of polyelectrolytes rather than by their charge density.

The conductivity of PILs as a function of inverse temperature between 10 and 110 °C is shown in Fig. 1. Independently of the repeat unit charge density the ionic conductivity of all studied PILs increased with increasing temperature and reached a level ranging from *ca.* 1×10^{-4} to 3×10^{-3} S cm⁻¹ at 110 °C. The temperature dependence of conductivity did not follow linear Arrhenius behavior indicating that anion diffusion results from various mechanisms: hopping of TFSI anions along a positively charged polymer backbone and local segmental motion of polymer chains.

Besides ionic conductivity another important property that should be considered for practical applications of PILs in *e.g.* the field of energy is their electrochemical stability. The electrochemical stability towards oxidative processes at the interface with the cathode is particularly important for battery applications.^{3,53} The electrochemical stability limits of PILs were assessed *via* cyclic voltammetry. As reasonable for electrochemical applications the ionic conductivity ($>10^{-5}$ S cm⁻¹) for PILs studied in this work was achieved only at 70 °C (Fig. 1); the latter was chosen as the experimental temperature for ESW studies. It should be noted that the selected temperature exceeds the T_g of all PILs and therefore during measurements they were in a rubber-like elastic state. Fig. 2 shows the anodic and cathodic scans of most ionic conductive polymers **9**, **10**, **13**, **14** and **15** at 70 °C. For all PILs two reversible redox peaks at -0.45 V and between -0.05 and 0.30 V *vs.* Li⁺/Li were observed that are associated with the reversible lithium plating and stripping at the copper electrode surface. Generally plating/stripping depends on the number of Li⁺ ions that are able to reach the surface of the Cu electrode. In its turn, this process is primarily affected by the lithium ion transport⁵⁴ and secondarily by the overall conductivity, which instead

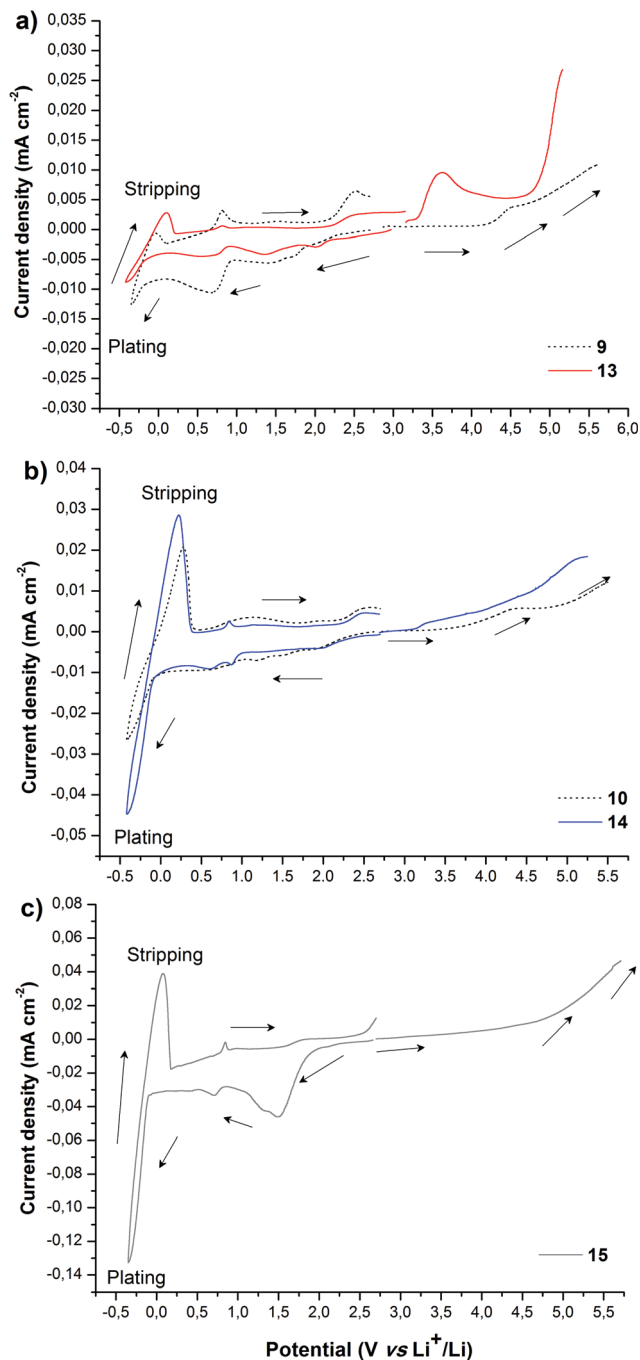


Fig. 2 Electrochemical stability windows obtained by CV for (a) PILs **9** and **13**, (b) PILs **10** and **14** and (c) PIL **15** at 70 °C (stainless steel as a working electrode and Li foil as counter and reference electrodes, scan rate 0.5 mV s⁻¹).

considers the movement of all charges (anions and cations). In the studied systems the intensity of the peaks was directly dependent on the ionic transport in polymers. The higher is the overall conductivity of PIL and, respectively, the ion movement, the more intense were the lithium plating and stripping processes.

Fig. 2 suggests that the cathodic limit potential is mainly determined by the reduction of the respective cations.



The irreversible oxidation peak at 1.5 V vs. Li^+/Li found for polymer **15** (Fig. 2c) can be attributed to the partial degradation of the 1,2,3-triazolium cations as was observed previously.²⁶ The increased cathodic stability up to -0.4 V vs. Li^+/Li and observation for Li stripping and plating in comparison with previously reported 1,2,3-triazolium-based PILs²⁶ may on one hand arise from the presence of the oxyethylene chain that could contribute to preventing the full reduction of the 1,2,3-triazolium cation. A similar influence of $\text{CH}_2\text{CH}_2\text{O}$ groups on the cathodic limit of 1,2,3-triazolate anions was recently reported.⁵⁵ On another hand, the cathodic stability of 1,2,3-triazolium-based PILs can be improved by the presence of LiTFSI as was noted formerly for pyrrolidinium ILs and PILs.^{6,56} It was previously observed that 1,2,3-triazolium cations exhibit less cathodic stability than quaternary ammonium and imidazolium cations.⁵⁷ Following these observations for ILs, imidazolium-based polymer **10** shows better cathodic stability in comparison with **15** and reveals no oxidation peaks until Li plating at -0.41 V (Fig. 2b and c). An ammonium-based polymer **9** demonstrates one additional marked oxidation peak at ca. 1.5 V (Fig. 2a). Although in PIL **9** the ammonium cation is separated from the 1,2,3-triazole ring by a methylene group, these peaks can probably be attributed to the cleavage of the ring by Dimroth rearrangement.⁵⁸ Interestingly, the presence of ammonium and imidazolium cations in heterocationic PILs **13** and **14** fully suppresses the reduction of 1,2,3-triazolium cations, thus increasing the electrochemical stability of PILs (Fig. 2a and b). Additionally, the couple of reversible peaks at 0.7 and 0.9 V versus Li^+/Li may be attributed to a redox process involving oxide species at the electrode surface. The other couple of small peaks observable in the potential range between 1.5 and 2.7 V is most likely associated with the multi-step decomposition of some impurity traces in PILs such as solvents. Indeed, these peaks disappear in the subsequent voltammetric cycles, which clearly states for the non-reproducibility of the processes and their insignificance for the overall performance of the system.

The anodic limit potential of cationic PILs is commonly determined by the oxidation of the anion. As all studied PILs contain the same TFSI anion, their anodic stability was expected to be above 4.0 V vs. Li^+/Li .^{6,39} Indeed, polymers **9** and **15** demonstrate electrochemical stability windows up to 4.3 and 4.8 V, respectively (Fig. 2a and c). In accordance with what was observed previously for ILs,⁵¹ imidazolium-based polymer **10** shows a slightly reduced anodic limit equal to 3.9 V vs. Li^+/Li (Fig. 2b). Unexpectedly, some irreversible reduction peaks at 3.6 and 3.3 V were found for polymers **13** and **14** having mixed cations (Fig. 2a and b). The authors were not able to find reasonable explanation for this phenomenon as usually the mixtures of Li salts either in EC-EMC solution⁵⁹ or in polymer electrolyte⁶⁰ show an improved stability in comparison with systems containing single salts. Overall, except for **13** and **15**, the breakdown voltage accounting for the full degradation of polyelectrolyte was not clearly observed. This can be attributed to the diffusion resistance and slow process kinetics in polymer electrolytes.^{61,62}

Experimental

Iodomethane (>99%, Aldrich), deca-1,9-diyne (**5**, 99%, Aldrich), 1,11-diazido-3,6,9-trioxoundecane (**6**, 99%, Aldrich), *N,N,N',N'*-tetramethyl-1,6-hexanediamine (99%, Aldrich), propargyl bromide (>97.0%, TCI chemicals), diisopropylethylamine (DIPEA, 99%, Aldrich), *N,N,N',N',N'*-pentamethyldiethylenetriamine (PMDETA, 99%, Aldrich), copper(i) iodide triethylphosphite ($\text{CuIP}(\text{OEt})_3$, 97%, Aldrich), lithium bis(trifluoromethylsulfonyl)imide (LiTFSI, 99%, Aldrich), acetonitrile (CH_3CN , HPLC grade 99%, Acros), dimethyl formamide (DMF, 99.8%, anhydrous, Acros), lithium metal foil (Chemetall Foote Corporation) and activated decolorizing charcoal (Acros) were used without further purification.

Imidazole (>99%, Aldrich) was purified by vacuum sublimation at $80\text{ }^\circ\text{C}/10^{-2}$ mmHg. 1,6-Dibromohexane (96%, Aldrich) was distilled under vacuum. Propargyl tosylate was prepared by the reaction of tosyl chloride with an excess of propargyl alcohol in accordance with the well-established procedure.⁶³

Ionic liquid monomers (ILMs) *N,N*-dimethyl-*N,N*-dipropargylammonium bis(trifluoromethylsulfonyl)imide (**1**) and *N,N*-dibutyl-*N,N*-dipropargylammonium bis(trifluoromethylsulfonyl)imide (**2**) were synthesized as described previously.⁴²

Synthesis of *N,N,N',N'*-tetramethyl-*N,N'*-dipropargyl-hexane-1,6-diammonium bis(trifluoromethylsulfonyl)imide (**3**)

Monomer **3** was synthesized in two steps: (1) *N*-alkylation of *N,N,N',N'*-tetramethyl-1,6-hexanediamine by propargyl bromide and (2) ion exchange reaction with an excess of LiTFSI.

A solution of propargyl bromide (9.13 g, 76.8 mmol) in anhydrous CH_3CN (10 mL) was added dropwise to a solution of *N,N,N',N'*-tetramethyl-1,6-hexanediamine (3.78 g, 21.9 mmol) in anhydrous CH_3CN (20 mL) maintained at $0-5\text{ }^\circ\text{C}$ under an inert atmosphere. After stirring for 5 minutes the start of the precipitation of a beige powder was observed. Stirring was continued for 1 h at $0-5\text{ }^\circ\text{C}$, then for 1 h at $25\text{ }^\circ\text{C}$ and finally for 10 h at $55\text{ }^\circ\text{C}$. The reaction mixture was centrifuged for 5 min at 5000 rpm, the mother liquor was decanted and a fresh portion of CH_3CN was added to the precipitate. After intense shaking the suspension was again centrifuged and this procedure was repeated three times. *N,N,N',N'*-Tetramethyl-*N,N'*-dipropargyl-hexane-1,6-diammonium bromide was isolated by filtration as a white-beige powder and dried for 8 h at $70\text{ }^\circ\text{C}/1$ mmHg (hereinafter with a special flask filled with P_2O_5 and introduced into the vacuum line). Yield: 8.64 g (96%); m.p. = $204-206\text{ }^\circ\text{C}$.

Lithium bis(trifluoromethylsulfonyl)imide (3.53 g, 12.2 mmol) was dissolved in H_2O (8 mL) and added dropwise to a solution of *N,N,N',N'*-tetramethyl-*N,N'*-dipropargyl-hexane-1,6-diammonium bromide (2.10 g, 5.12 mmol) in H_2O (20 mL) at room temperature. The mixture was stirred for 2 h at $25\text{ }^\circ\text{C}$ and the precipitation of a yellow powder was observed. The temperature was raised to $70\text{ }^\circ\text{C}$ and stirring was continued for 1 h. After cooling down to room temperature the upper aqueous layer was decanted and the residual powder was redissolved in 80 mL of



dichloromethane (CH₂Cl₂). The CH₂Cl₂ solution was washed with water (3 × 30 mL) and dried over anhydrous MgSO₄. The magnesium sulfate was filtered off, the decolorizing charcoal was added to the solution and the suspension was stirred at room temperature for another 2 h. The charcoal was filtered off and CH₂Cl₂ was stripped off under reduced pressure. Bispropargyl monomer **3** was obtained as a white crystalline powder, which was dried at 50 °C/1 mmHg for 4 h. Yield: 3.45 g (83%); m.p. = 68–70 °C (68 °C by DSC); cryst.p. = 14 °C (DSC); *T*_g = –45 °C (DSC); ¹H NMR (300 MHz, DMSO-*d*₆, δ, ppm): 4.35 (d, 4H, *J*_{HH} = 3.0 Hz, CH₂C≡CH), 4.02 (t, 2H, *J*_{HH} = 3.0 Hz, C≡CH), 3.33 (m, 4H, NCH₂CH₂), 3.07 (s, 12H, NCH₃), 1.71 (m, 4H, NCH₂CH₂), 1.33 (m, 4H, NCH₂CH₂CH₂); ¹³C NMR (75.5 MHz, DMSO-*d*₆, δ, ppm): 125.8–113.0 (q, ¹*J*_{CF} = 322 Hz, CF₃), 82.8, 72.2, 62.9, 53.2, 49.8, 25.1, 21.6; ¹⁹F NMR (282 MHz, DMSO-*d*₆, δ, ppm): –78.7 (s, CF₃); IR (KBr pellet): 3305 (m, ν_{C≡H}), 3251 (m, ν_{C≡H}), 3049 (w, ν_{CH}), 2996 (w, ν_{CH}), 2966 (m, ν_{CH}), 2626 (w, ν_{CH}), 2867 (w, ν_{CH}), 2135 (w, ν_{C≡H}), 1484 (m), 1423 (w), 1348 (vs, ν_{as=O}), 1196 (vs, ν_{CF}), 1141 (s, ν_{s=O}), 1054 (vs, ν_{CF}), 969 (w), 890 (m), 796 (m), 766 (w), 741 (m), 688 (w), 654 (w), 615 (vs, δ_{C≡H}), 570 (s), 514 (s) cm^{–1}; anal. calcd for C₂₀H₃₀F₁₂N₄O₈S₄ (810.7), %: C 29.63; H 3.73; N 6.91; found, %: C 29.48; H 3.66; N 6.90.

Synthesis of 1,1'-(hexane-1,6-diyl)bis(3-propargyl-imidazolium) bis(trifluoromethylsulfonyl)imide (4)

Monomer **4** was prepared *via* a three step reaction: (1) synthesis of *N*-propargyl imidazole, (2) *N*-alkylation of *N*-propargyl imidazole by 1,6-dibromohexane and (3) ion exchange reaction with an excess of LiTFSI.

A solution of propargyl tosylate (48.08 g, 228.7 mmol) in anhydrous CH₃CN (100 mL) was added dropwise to a solution of imidazole (34.2 g, 503 mmol) in anhydrous CH₃CN (140 mL) at 0–5 °C under an inert atmosphere. Stirring was continued for 1 h at 0–5 °C and then for 2 h at 40 °C. CH₃CN was evaporated under reduced pressure at room temperature and the residual oil was distilled under vacuum (bp = 62 °C/0.08 mmHg). The obtained colorless oil still contained 10 mol% of imidazole (¹H NMR). To remove impurity 1.5 mL of 37 wt% HCl was added to the oil with vigorous stirring at room temperature. Stirring was continued for 15 minutes and CH₂Cl₂ (100 mL) was added to the solution. The organic layer was separated, washed with H₂O (3 × 5 mL) and dried over anhydrous MgSO₄. The magnesium sulfate was filtered off and CH₂Cl₂ was evaporated under reduced pressure. *N*-Propargyl imidazole was obtained as a yellow oil, which was dried at 25 °C/1 mmHg for 4 h. Yield: 9.78 g (40%); ¹H NMR (400 MHz, DMSO-*d*₆, δ, ppm): 7.70 (s, 1H, NCHN), 7.22 (s, 1H, NCH=CHN), 6.95 (s, 1H, NCH=CHN), 4.93 (m, 2H, CH₂C≡CH–), 3.52 (m, 2H, CH₂C≡CH); ¹³C NMR (100 MHz, DMSO-*d*₆, δ, ppm): 137.1 (C₂im), 128.9 (C₄im), 119.5 (C₅im), 78.7 (CH₂C≡CH), 76.2 (CH₂C≡CH–), 35.6 (CH₂N); IR (KBr pellet): 3290 (vs, ν_{C≡H}), 3191 (s, ν_{CH} imidazole), 3112 (vs, ν_{CH} ν_{CH} imidazole), 2961 (m, ν_{CH}), 2927 (m, ν_{CH}), 2121 (s, ν_{C≡H}), 1670 (w), 1637 (w), 1594 (m), 1505 (vs), 1435 (s), 1392 (s), 1348 (s), 1282 (vs), 1231 (vs), 1107 (s), 1075 (vs), 1029 (s), 944 (m),

907 (s), 820 (vs), 736 (vs, δ_{CH} imidazole), 660 (vs), 616 (vs, δ_{C≡H}), 458 (w) cm^{–1}.

A solution of 1,6-dibromohexane (4.50 g, 18.5 mmol) in anhydrous CH₃CN (25 mL) was added dropwise to a solution of *N*-propargyl imidazole (4.29 g, 40.6 mmol) in anhydrous CH₃CN (25 mL) maintained at 0–5 °C under an inert atmosphere. The stirring was continued for 1 h at 0–5 °C, 2 h at 25 °C, 66 h at 45 °C and 23 h at 60 °C. The precipitation of the light-brown oil was observed and the emulsion was placed in the fridge (5 °C) for 48 h, whereupon the crystallization occurred. The light-brown crystals were filtered, washed with Et₂O, then thrice with CH₂Cl₂, filtered again and smashed into powder with a mortar. 1,1'-(Hexane-1,6-diyl)bis(3-propargyl-imidazolium) bromide was crystallized from a CH₃CN/H₂O mixture (150/11 mL) as light-yellow crystals that were dried at 60 °C/1 mmHg for 12 h. Yield: 6.20 g (73%); m.p. = 143–144 °C (139 °C by DSC).

Lithium bis(trifluoromethylsulfonyl)imide (11.30 g, 39.3 mmol) was dissolved in H₂O (25 mL) and added dropwise to a solution of 1,1'-(hexane-1,6-diyl)bis(3-propargyl-imidazolium) bromide (5.98 g, 13.1 mmol) in H₂O (30 mL) at room temperature. The mixture was stirred for 2 h at 25 °C and the precipitation of a yellow powder was observed. The yellow powder was filtered and dissolved in acetone (60 mL). A solution of LiTFSI (6.50 g, 22.6 mmol) in acetone (15 mL) was added dropwise under stirring to the obtained solution. The reaction continued overnight at room temperature, whereupon the solvent was evaporated and the residual orange oil was extensively washed with H₂O (5 × 20 mL). The oil was dissolved in CH₃CN and dried over anhydrous MgSO₄. The magnesium sulfate was filtered off, the decolorizing charcoal was added to the solution and the suspension was stirred at room temperature for another 2 h. The charcoal was filtered off and CH₃CN was evaporated under reduced pressure. Monomer **4** was obtained as a slightly viscous yellow oil, which was dried at 55 °C/1 mmHg for 12 h. Yield: 7.09 g (63%); *T*_g = –56 °C (DSC); ¹H NMR (600 MHz, DMSO-*d*₆, δ, ppm): 9.26 (s, 2H, NCHN), 7.81 (s, 4H, NCH=CHN), 5.18 (m, 4H, CH₂C≡CH–), 4.19 (t, 4H, *J*_{HH} = 7.2 Hz, CH₂N), 3.81 (m, 2H, CH₂C≡CH–), 1.80 (t, 4H, *J*_{HH} = 7.2 Hz, NCH₂CH₂), 1.28 (m, 4H, NCH₂CH₂CH₂); ¹³C NMR (151 MHz, DMSO-*d*₆, δ, ppm): 136.1 (C₂im), 122.8 (C₄im), 122.4 (C₅im), 122.7–116.3 (q, ¹*J*_{CF} = 322 Hz, CF₃), 78.9 (CH₂C≡CH–), 75.9 (CH₂C≡CH–), 48.9 (NCH₂), 38.6 (CH₂C≡CH), 29.1 (NCH₂CH₂), 24.9 (NCH₂CH₂CH₂); ¹⁹F NMR (565 MHz, DMSO-*d*₆, δ, ppm): –78.8 (s, CF₃); IR (KBr pellet): 3277 (m, ν_{C≡H}), 3151 (s, ν_{CH} imidazolium), 2949 (m, ν_{CH}), 2868 (w, ν_{CH}), 2138 (w, ν_{C≡H}), 1564 (s), 1475 (w), 1444 (w), 1350 (vs, ν_{as=O}), 1330 (vs), 1193 (vs, ν_{CF}), 1137 (vs, ν_{s=O}), 1055 (vs, ν_{CF}), 955 (w), 842 (m), 790 (s), 740 (s, δ_{CH} imidazolium), 654 (m), 614 (vs, δ_{C≡H}), 570 (vs), 513 (s), 406 (m) cm^{–1}; anal. calcd for C₂₂H₂₄F₁₂N₆O₈S₄ (856.7), %: C 30.84; H 2.82; F 26.61; found, %: C 31.18; H 2.80; F 26.69.

CuAAC polyaddition of bisalkynes with 1,11-diazido-3,6,9-trioxundecane (6). Synthesis of poly(1,2,3-triazole)s 7–11

Polymers **7–11** were synthesized *via* AA + BB CuAAC polyaddition and a typical procedure is detailed below by the example of poly(1,2,3-triazole) **10** preparation.



Dialkyne **4** (1.01 g, 1.18 mmol), diazide **6** (0.29 g, 1.18 mmol) and *N,N*-diisopropylethylamine (0.31 g, 2.36 mmol) were dissolved in anhydrous DMF (6 mL) under an inert atmosphere at room temperature. Afterwards CuIP(OEt)₃ (42.2 mg, 0.118 mmol) was added in one portion to the transparent yellow solution and stirring was continued in the dark for 68 h at 80 °C. The resulting brown solution was precipitated into excess of diethyl ether and the brown oil was centrifuged (10 000 rpm, 5 min). The oil was dissolved in CH₃CN (5 mL) and precipitated twice into an excess of H₂O containing 2–3 drops of PMEDTA and then thrice in H₂O. Poly(1,2,3-triazole) **10** was recovered as a dark brown gummy material and was dried at 80 °C/1 mmHg for 12 h. Yield: 0.82 g (63%); *T*_g = –12 °C (DSC); *M*_n = 119.3 kDa, *M*_w/*M*_n = 1.82 (GPC); ¹H NMR (600 MHz, DMSO-*d*₆, δ, ppm): 9.26 (s, 2H, imNCHN), 8.22 (s, 2H, CCHNtriazole), 7.78 (s, 2H, imNCH=CHN), 7.76 (s, 2H, imNCH=CHN), 5.53 (s, 4H, imNCH₂triazole), 4.55 (t, 4H, *J*_{HH} = 4.8 Hz, NCH₂CH₂O), 4.17 (t, 4H, *J*_{HH} = 7.8 Hz, imNCH₂CH₂), 3.82 (t, 4H, *J*_{HH} = 5.4 Hz, NCH₂CH₂O), 3.52 (m, 4H, NCH₂CH₂OCH₂CH₂O), 3.46 (m, 4H, NCH₂CH₂OCH₂CH₂O), 1.79 (m, 4H, imNCH₂CH₂), 1.28 (m, 4H, imNCH₂CH₂CH₂); ¹³C NMR (151 MHz, DMSO-*d*₆, δ, ppm): 140.2 (CCHNtriazole), 136.1 (C₂im), 124.9 (CCHNtriazole), 122.6 (C₄im and C₅im), 123.7–116.3 (q, ¹*J*_{CF} = 322 Hz, CF₃), 69.6, 69.5 (NCH₂CH₂OCH₂CH₂OCH₂CH₂), 68.6 (NCH₂CH₂O), 49.6 (NCH₂CH₂O), 48.9 (imNCH₂CH₂), 43.8 (imNCH₂triazole), 29.1 (imNCH₂CH₂), 24.9 (imNCH₂CH₂CH₂); ¹⁹F NMR (565 MHz, DMSO-*d*₆, δ, ppm): –78.8 (s, CF₃); IR (ATR mode): 3152 (m, ν_{CH} imidazolium), 3113 (w, ν_{CH} triazole), 3096 (w, ν_{CH} triazole), 2934 (w, ν_{CH}), 2916 (w, ν_{CH}), 2868 (m, ν_{CH}), 1697 (w), 1598 (w), 1563 (m), 1515 (w), 1449 (m), 1348 (s, ν_{as}=O), 1329 (s), 1180 (vs, ν_{CF}), 1132 (vs, ν_{ss}=O), 1051 (vs, ν_{CF}), 833 (m), 740 (s, δ_{CH} imidazolium), 652 (m), 613 (vs, δ_{C≡H}), 599 (s), 570 (vs), 531 (w), 512 (vs) cm^{–1}; Anal. calcd for C₃₁H₄₄F₁₂N₁₂O₁₁S₄ (1117.0), %: C 33.33; H 3.97; N 15.05; found, %: C 33.19; H 3.96; N 14.94.

Synthesis of poly(1,2,3-triazolium)s 12–15 by *N*-alkylation and ion exchange reactions

Polymers **12–15** were synthesized in two steps: *N*-alkylation of the 1,2,3-triazole groups of polymers **8–11** by an excess of CH₃I and subsequent ion exchange reaction with LiTFSI. A typical procedure is given below by the example of PIL **14** preparation.

Iodomethane (0.93 g, 6.54 mmol) was added to a solution of poly(1,2,3-triazole) **10** (0.72 g, 0.654 mmol) in anhydrous DMF (10 mL) at room temperature under an inert atmosphere. Stirring was continued at 60 °C for 90 h, whereupon additional CH₃I (0.50 g, 3.52 mmol) was added and the solution was further stirred at 80 °C for 72 h. The crude reaction mixture was diluted with DMF and precipitated two times in acetone to yield a dark brown viscous oil which was separated by centrifugation (10 000 rpm, 10 min).

The obtained oil was redissolved in H₂O (20 mL) and an aqueous solution of LiTFSI (0.69 g, 2.4 mmol in 5 mL of H₂O) was added dropwise under vigorous stirring at room temperature. The mixture was stirred for 12 h at 40 °C and the precipitation of a viscous brown oil was observed. The aqueous layer was decanted and the oil was washed with H₂O (3 × 20 mL), dissolved in acetone (6 mL) and precipitated again into an excess

of H₂O. Polymer **14** was recovered as a brown gummy solid after centrifugation (10 000 rpm, 10 min) and drying at 80 °C/1 mmHg for 12 h. Yield: 0.67 g (69%); *T*_g = –1 °C (DSC); *M*_n = 121.9 kDa, *M*_w/*M*_n = 1.48 (GPC); ¹H NMR (600 MHz, DMSO-*d*₆, δ, ppm): 9.30 (s, 2H, imNCHN), 8.97 (s, 2H, CCHNtriazolium), 7.85 (m, 4H, imNCH=CHN), 5.84 (s, 4H, imNCH₂triazolium), 4.83 (t, 4H, *J*_{HH} = 4.8 Hz, NCH₂CH₂O), 4.34 (s, 6H, triazolium NCH₃), 4.18 (t, 4H, *J*_{HH} = 7.8 Hz, imNCH₂CH₂), 3.92 (t, 4H, *J*_{HH} = 5.4 Hz, NCH₂CH₂O), 3.58 (m, 4H, NCH₂CH₂OCH₂CH₂O), 3.51 (m, 4H, NCH₂CH₂OCH₂CH₂O), 1.82 (m, 4H, imNCH₂CH₂), 1.35 (m, 4H, imNCH₂CH₂CH₂); ¹³C NMR (151 MHz, DMSO-*d*₆, δ, ppm): 137.2 (C₂im), 131.4 (CCHNtriazolium), 122.9 (C₄im), 122.7 (C₅im), 122.6–116.3 (q, ¹*J*_{CF} = 322 Hz, CF₃), 69.5 (NCH₂CH₂OCH₂CH₂OCH₂CH₂), 67.4 (NCH₂CH₂), 53.4 (NCH₂CH₂O), 49.1 (imNCH₂CH₂), 40.3 (imNCH₂triazolium), 38.4 (NCH₃), 29.2 (imNCH₂CH₂), 25.1 (imNCH₂CH₂CH₂); ¹⁹F NMR (565 MHz, DMSO-*d*₆, δ, ppm): –78.8 (s, CF₃); IR (ATR mode): 3151 (w, ν_{CH} imidazolium), 3119 (w, ν_{CH} triazolium), 3089 (w, ν_{CH} triazolium), 2929 (w, ν_{CH}), 2875 (m, ν_{CH}), 1561 (m), 1457 (m), 1346 (s, ν_{as}=O), 1328 (s), 1175 (vs, ν_{CF}), 1131 (vs, ν_{ss}=O), 1050 (vs, ν_{CF}), 935 (w), 843 (m), 791 (s), 741 (s, δ_{CH} imidazolium), 692 (m, δ_{CH} triazolium), 652 (s), 611 (vs, δ_{C≡H}), 598 (s), 569 (vs), 505 (vs) cm^{–1}; anal. calcd for C₃₇H₅₀F₂₄N₁₄O₁₉S₈ (1707.4), %: C 26.03; H 2.95; N 11.49; found, %: C 26.07; H 2.91; N 11.51.

Methods

NMR spectra were recorded on AMX-400, Avance II 500 MHz and Avance III HD 600 MHz spectrometers (Bruker) at 25 °C in the indicated deuterated solvents and are listed in ppm. The signal corresponding to the residual protons or carbons of the deuterated solvent was used as an internal standard for ¹H and ¹³C NMR, respectively. C₆F₆ was utilized as an external standard for ¹⁹F NMR. IR spectra were acquired on a Nicolet Magna-750 Fourier IR-spectrometer using KBr pellets or on a Bruker Tensor 27 Fourier IR-spectrometer using ATR technology (128 scans, resolution is 2 cm^{–1}) and Spectragryph optical spectroscopy software.⁶⁴

A 1200 Infinity gel permeation chromatograph (GPC, Agilent Technologies) was used to determine *M*_n, *M*_w and *M*_w/*M*_n of PILs. The chromatograph was equipped with an integrated IR detector, a PL PolarGel-M column and a PL PolarGel-M guard column (Agilent Technologies). The eluent was a 0.1 M solution of LiTFSI in DMF and the flow rate was 1.0 mL min^{–1} at 50 °C. Polystyrene standards (EasiVial PS-M, Agilent Technologies, *M*_p = 162–500 × 10³ g mol^{–1}) were used to perform calibration.

For Differential Scanning Calorimetry (DSC) measurements all samples (ionic monomers and PILs) were hermetically sealed in aluminum pans inside an argon-filled glove-box (MBRAUN MB-Labstar, H₂O and O₂ content < 0.5 ppm). DSC experiments were performed on a DSC3 + STARE System differential calorimeter (Mettler Toledo). For PILs DSC measurements were conducted with a heating rate of 10 °C min^{–1} in the range of –80 to 150 °C, while for ionic monomers a very low rate of 3 °C min^{–1} was applied as recommended for highly viscous ionic liquids.⁶⁵ Glass transition temperatures (*T*_g) were determined during the second heating cycle.



Thermal gravimetric analysis (TGA) was carried out in air on a TGA2 STARE System (Mettler Toledo) applying a heating rate of $5\text{ }^{\circ}\text{C min}^{-1}$.

Ionic conductivity (σ_{DC}) was determined by electrochemical impedance spectroscopy (EIS or complex alternating-current (AC) impedance spectroscopy) with a VSP potentiostat/galvanostat (Bio-Logic Science Instruments). To avoid any influence of moisture/humidity on the ionic conductivity of polymer electrolytes, the latter were preliminary dried at $60\text{ }^{\circ}\text{C}/1\text{ mmHg}$ for 12 h in a B-585 oven (Buchi Glass Drying Oven, Switzerland) filled with P_2O_5 and were transferred under vacuum inside an argon-filled glovebox. Polymers were sandwiched between two stainless steel electrodes. The distance between the electrodes was kept equal to $250\text{ }\mu\text{m}$ using a Teflon spacer ring with an inner area of 0.5 cm^2 . Symmetrical stainless steel/PIL/stainless steel assembly was clamped into the 2032 coin cell and afterwards was taken out from the glovebox. EIS experiments were carried out by applying a 10 mV perturbation in the frequency range from 10^{-2} to 2×10^5 Hz at the open circuit potential (OCV). The ohmic resistance (R_{Ω}) of the sample, obtained from the Nyquist plot at the low frequency end of the semicircle, was used to calculate the ionic conductivity using the following equation:

$$\sigma = \frac{l}{A R_{\Omega}}$$

where l is the sample thickness and A is the electrode area. The measurements were carried out in the temperature range from 10 to $110\text{ }^{\circ}\text{C}$. The temperature was controlled using a programmed M-53 oven (Binder, Germany), where cells were allowed to reach thermal equilibrium for at least 30 minutes before each test.

Cyclic voltammetry (CV) and VSP potentiostat/galvanostat (Bio-Logic Science Instruments, France) were used to determine the electrochemical stability window of solid polymer electrolytes at $70\text{ }^{\circ}\text{C}$. Prior to measurement PILs (0.2 g) were dissolved in anhydrous CH_3CN (2 mL) inside the glovebox and 10 wt% of LiTFSI (0.02 g) were added under stirring until the formation of a clear solution. The solvent was evaporated under reduced pressure in a B-585 oven (Buchi Glass Drying Oven, Switzerland) and gummy samples were further dried there at $60\text{ }^{\circ}\text{C}/1\text{ mmHg}$ for 12 h. To avoid moisture or oxygen contamination samples were afterwards transferred under vacuum back to an argon-filled glovebox, where they were used for the two-electrode 2032 coin cell assembly. The two-electrode cells were compiled by sandwiching samples between the working electrode and a lithium metal foil, which served as the reference and the counter electrode, simultaneously. Stainless steel and copper disks were used as working electrodes during anodic and cathodic stability measurements, respectively. To evaluate anodic limits, potential sweeps were carried out between OCV and $5.5\text{ V vs. Li}^+/\text{Li}$ at a constant rate of 0.5 mV s^{-1} . To determine cathodic limits, potential sweeps were performed between OCV and $-0.5\text{ V vs. Li}^+/\text{Li}$ at the same constant rate.

The charge density of PILs as the number of charges per gram of material was calculated using the following equation:

$$\text{Charge density} = \frac{1000 \cdot n}{M}$$

where n is the number of charges per repeat unit and M is the molar mass of the repeat unit.

Conclusions

The aim of this paper was to obtain polyelectrolytes with enhanced ionic conductivity and electrochemical stability as well as to fill the gap in fundamental knowledge concerning the influence of repeat unit charge density on the structure/property relationships of PILs. Within this scope a series of TFSI-based cationic PILs was obtained from (1) CuAAC polyaddition using cationic ILMs and (2) subsequent *N*-alkylation of poly(1,2,3-triazole) intermediates. The synthesized PILs differed by the number of charge carriers per monomer unit (from 1 to 4 ion pairs) and by the nature of cation combinations (1,2,3-triazolium, imidazolium or ammonium as well as 1,2,3-triazolium/imidazolium or 1,2,3-triazolium/ammonium associations).

Within the studied PILs thermal stability, heat resistance (glass transition temperature), ionic conductivity and electrochemical stability were all found to be significantly affected by repeat unit charge density. Generally, the higher the repeat unit charge density of PILs the higher their T_g and the lower their ionic conductivity and thermal stability. The observed dependences passed through a maximum when PILs had two cations per monomer unit that were separated by alkyl or oxyethylene chains. For almost all combinations, 1,2,3-triazolium-containing PILs showed the highest ionic conductivity. A good cathodic stability on the Li metal electrode was seen when PILs with a mixed type of cation were tested. It is worth noting that the presence of ammonium and imidazolium cations in heterocationic PILs was able to hamper the reduction of 1,2,3-triazolium cations and to increase the cathodic limit down to $-0.4\text{ V vs. Li}^+/\text{Li}$ at $70\text{ }^{\circ}\text{C}$. The highest ionic bulk conductivity ($1.8 \times 10^{-5}\text{ S cm}^{-1}$) and anodic stability ($4.8\text{ V vs. Li}^+/\text{Li}$ at $70\text{ }^{\circ}\text{C}$) were demonstrated by PIL 15 containing solely 1,2,3-triazolium cations. Such PILs can be potential candidates for the next generation of safe and flexible polymer-based electrochemical devices requiring high conductivity in the solid state, for example supercapacitors, smart windows or actuators. However, when an application demands high electrochemical stability, it is preferred to include more stable cations such as ammonium or imidazolium in the polymer backbone. Therefore, the future application area of PILs with mixed type cations can include Li batteries.

Conflicts of interest

There are no conflicts to declare.



Acknowledgements

Merlin Cotessat is grateful to the Luxembourg Institute of Science and Technology (LIST) for the sponsorship of his internship and work in the Materials and Research Technology Department (MRT). This work was supported by the Luxembourg National Research Fund (FNR) and Agency Nationale de la Recherche (ANR) through ANR-FNR project DISAFECAP (Agreement number INTER/ANR/19/13358226). Elemental analysis and IR spectroscopy were performed with the financial support from the Ministry of Science and Higher Education of the Russian Federation using the equipment of Center for molecular composition studies of INEOS RAS. This manuscript is in honor of the 50 year anniversary of the French Polymer Group (Groupe Français des Polymères – GFP).

Notes and references

- X. Lu, M. Yu, G. Wang, Y. Tong and Y. Li, *Energy Environ. Sci.*, 2014, **7**, 2160.
- W. K. Chee, H. N. Lim, Z. Zainal, N. M. Huang, I. Harrison and Y. Andou, *J. Phys. Chem. C*, 2016, **120**, 4153–4172.
- H. Zhang, C. Li, M. Piszcz, E. Coxa, T. Rojo, L. M. Rodriguez-Martinez, M. Armand and Z. Zhou, *Chem. Soc. Rev.*, 2017, **46**, 797–815.
- L. Yue, J. Ma, J. Zhang, J. Zhao, S. Dong, Z. Liu, G. Cui and L. Chen, *Energy Storage Mater.*, 2016, **5**, 139–164.
- M. J. Park, I. Choi, J. Hong and O. Kim, *J. Appl. Polym. Sci.*, 2013, **129**, 2363–2376.
- A.-L. Pont, R. Marcilla, I. De Meatza, H. Grande and D. Mecerreyes, *J. Power Sources*, 2009, **188**, 558–563.
- A. Balducci, *J. Power Sources*, 2016, **326**, 534–540.
- G. Ayalneh Tiruye, D. Munoz-Torrero, J. Palma, M. Anderson and R. Marcilla, *J. Power Sources*, 2015, **279**, 472–480.
- I. Osada, H. de Vries, B. Scrosati and S. Passerini, *Angew. Chem., Int. Ed.*, 2016, **55**, 500–513.
- A. S. Shaplov, R. Marcilla and D. Mecerreyes, *Electrochim. Acta*, 2015, **175**, 18–34.
- J. Yuan and M. Antonietti, *Polymer*, 2011, **52**, 1469–1482.
- D. Mecerreyes, *Prog. Polym. Sci.*, 2011, **36**, 1629–1648.
- J. Yuan, D. Mecerreyes and M. Antonietti, *Prog. Polym. Sci.*, 2013, **38**, 1009–1036.
- M. M. Obadia and E. Drockenmuller, *Chem. Commun.*, 2016, **52**, 2433–2450.
- M. D. Green and T. E. Long, *Polym. Rev.*, 2009, **49**, 291–314.
- E. B. Anderson and T. E. Long, *Polymer*, 2010, **51**, 2447–2454.
- W. Qian, J. Texter and F. Yan, *Chem. Soc. Rev.*, 2017, **46**, 1124–1159.
- G. B. Appetecchi, G.-T. Kim, M. Montanino, M. Carewska, R. Marcilla, D. Mecerreyes and I. D. Meatza, *J. Power Sources*, 2010, **195**, 3668–3675.
- A. S. Shaplov, D. O. Ponkratov and Y. S. Vygodskii, *Polym. Sci., Ser. B*, 2016, **58**, 73–142.
- W. Zhang, Z. Kochovski, B. V. K. J. Schmidt, M. Antonietti and J. Yuan, *Polymer*, 2016, **107**, 509–516.
- W. Zhang and J. Yuan, *Macromol. Rapid Commun.*, 2016, **37**, 1124–1129.
- M. Yahia, S. Mei, A. P. Mathew and J. Yuan, *ACS Macro Lett.*, 2019, **8**, 1372–1377.
- H. Ohno, *Macromol. Symp.*, 2007, **249–250**, 551–556.
- B. Qiu, B. Lin and F. Yan, *Polym. Int.*, 2013, **62**, 335–337.
- A. S. Shaplov, D. O. Ponkratov, P.-H. Aubert, E. I. Lozinskaya, C. Plesse, F. Vidal and Y. S. Vygodskii, *Chem. Commun.*, 2014, **50**, 3191–3193.
- G. Colliat-Dangus, M. M. Obadia, Y. S. Vygodskii, A. Serghei, A. S. Shaplov and E. Drockenmuller, *Polym. Chem.*, 2015, **6**, 4299–4308.
- E. Drockenmuller, J. Bernard, A. Serghei, B. Zhang and R. Sood, *Polym. Chem.*, 2015, **6**, 3521–3528.
- M. Lee, U. H. Choi, R. H. Colby and H. W. Gibson, *Chem. Mater.*, 2010, **22**, 5814–5822.
- M. Lee, U. H. Choi, D. Salas-de la Cruz, A. Mittal, K. I. Winey, R. H. Colby and H. W. Gibson, *Adv. Funct. Mater.*, 2011, **21**, 708–717.
- A. S. Shaplov, E. I. Lozinskaya, R. Losada, C. Wandrey, A. T. Zdvizhkov, A. A. Korlyukov, K. A. Lyssenko, I. A. Malyskhina and Y. S. Vygodskii, *Polym. Adv. Technol.*, 2011, **22**, 448–457.
- A. Jourdain, A. Serghei and E. Drockenmuller, *ACS Macro Lett.*, 2016, **5**, 1283–1286.
- S. Yi, W. Leon, D. Vezenov and S. L. Regen, *ACS Macro Lett.*, 2016, **5**, 915–918.
- B. Qiu, B. Lin, L. Qiu and F. Yan, *J. Mater. Chem.*, 2012, **22**, 1040–1045.
- B. Qiu, B. Lin, Z. Si, L. Qiu, F. Chu, J. Zhao and F. Yan, *J. Power Sources*, 2012, **217**, 329–335.
- X. Chen, J. Zhao, J. Zhang, L. Qiu, D. Xu, H. Zhang, X. Han, B. Sun, G. Fu, Y. Zhang and F. Yan, *J. Mater. Chem.*, 2012, **22**, 18018–18024.
- M. M. Obadia, A. Jourdain, A. Serghei, T. Ikeda and E. Drockenmuller, *Polym. Chem.*, 2017, **8**, 910–917.
- Y. S. Vygodskii, O. A. Mel'nik, E. I. Lozinskaya, A. S. Shaplov, I. A. Malyskhina, N. D. Gavrilova, K. A. Lyssenko, M. Y. Antipin, D. G. Golovanov, A. A. Korlyukov, N. Ignat'ev and U. Welz-Biermann, *Polym. Adv. Technol.*, 2007, **18**, 50–63.
- R. Marcilla, F. Alcaide, H. Sardon, J. A. Pomposo, C. Pozo-Gonzalo and D. Mecerreyes, *Electrochem. Commun.*, 2006, **8**, 482–488.
- K. Yin, Z. Zhang, X. Li, L. Yang, K. Tachibana and S. Hirano, *J. Mater. Chem. A*, 2014, **3**, 170–178.
- J. M. C. Puguán and H. Kim, *J. Ind. Eng. Chem.*, 2019, **74**, 1–6.
- D. Flachard, A. Serghei, M. Fumagalli and E. Drockenmuller, *Polym. Int.*, 2019, **68**, 1591–1598.
- Y. S. Vygodskii, A. S. Shaplov, E. I. Lozinskaya, P. S. Vlasov, I. A. Malyskhina, N. D. Gavrilova, P. Santhosh Kumar and M. R. Buchmeiser, *Macromolecules*, 2008, **41**, 1919–1928.
- B. P. Mudraboyina, M. M. Obadia, I. Allaoua, R. Sood, A. Serghei and E. Drockenmuller, *Chem. Mater.*, 2014, **26**, 1720–1726.
- P. Dimitrov-Raytchev, S. Beghdadi, A. Serghei and E. Drockenmuller, *J. Polym. Sci., Part A: Polym. Chem.*, 2013, **51**, 34–38.



- 45 S. Binauld, E. Fleury and E. Drockenmuller, *J. Polym. Sci., Part A: Polym. Chem.*, 2010, **48**, 2470–2476.
- 46 S. Zhang, N. Sun, X. He, X. Lu and X. Zhang, *J. Phys. Chem. Ref. Data*, 2006, **35**, 1475.
- 47 A. S. Shaplov, E. I. Lozinskaya, D. O. Ponkratov, I. A. Malyshkina, F. Vidal, P.-H. Aubert, O. V. Okatova, G. M. Pavlov, L. I. Komarova, C. Wandrey and Y. S. Vygodskii, *Electrochim. Acta*, 2011, **57**, 74–90.
- 48 W. Ogihara, S. Washiro, H. Nakajima and H. Ohno, *Electrochim. Acta*, 2006, **51**, 2614–2619.
- 49 Y. Cao and T. Mu, *Ind. Eng. Chem. Res.*, 2014, **53**, 8651–8664.
- 50 M. M. Obadia, A. Jourdain, P. Cassagnau, D. Montarnal and E. Drockenmuller, *Adv. Funct. Mater.*, 2017, **27**, 1703258.
- 51 H. Matsumoto, H. Sakaebe and K. Tatsumi, *J. Power Sources*, 2005, **146**, 45–50.
- 52 M. Watanabe, M. L. Thomas, S. Zhang, K. Ueno, T. Yasuda and K. Dokko, *Chem. Rev.*, 2017, **117**, 7190–7239.
- 53 G. G. Eshetu, D. Mecerreyes, M. Forsyth, H. Zhang and M. Armand, *Mol. Syst. Des. Eng.*, 2019, **4**, 294–309.
- 54 G. M. A. Girard, M. Hilder, D. Nucciarone, K. Whitbread, S. Zavorine, M. Moser, M. Forsyth, D. R. MacFarlane and P. C. Howlett, *J. Phys. Chem. C*, 2017, **121**, 21087–21095.
- 55 D. Flachard, J. Rolland, M. M. Obadia, A. Serghei, R. Bouchet and E. Drockenmuller, *Chem. Commun.*, 2018, **54**, 9035–9038.
- 56 J.-H. Shin, W. A. Henderson and S. Passerini, *Electrochem. Solid-State Lett.*, 2005, **8**, A125.
- 57 S. Sanghi, E. Willett, C. Versek, M. Tuominen and E. B. Coughlin, *RSC Adv.*, 2012, **2**, 848–853.
- 58 E. S. H. El Ashry, S. Nadeem, M. R. Shah and Y. E. Kilany, *Advances in Heterocyclic Chemistry*, Elsevier, 2010, vol. 101, pp. 161–228.
- 59 X. Chen, W. Xu, M. H. Engelhard, J. Zheng, Y. Zhang, F. Ding, J. Qian and J.-G. Zhang, *J. Mater. Chem. A*, 2014, **2**, 2346.
- 60 S. Li, Y.-M. Chen, W. Liang, Y. Shao, K. Liu, Z. Nikolov and Y. Zhu, *Joule*, 2018, **2**, 1838–1856.
- 61 Y. V. Oza, D. R. MacFarlane, M. Forsyth and L. A. O'Dell, *Electrochim. Acta*, 2015, **175**, 80–86.
- 62 M. Martínez-Ibañez, E. Sánchez-Diez, L. Qiao, Y. Zhang, X. Judez, A. Santiago, I. Aldalur, J. Carrasco, H. Zhu, M. Forsyth, M. Armand and H. Zhang, *Adv. Funct. Mater.*, 2020, 2000455.
- 63 M. N. Soltani Rad, S. Behrouz, J. Mohammadtaghi-Nezhad, E. Zarenezhad and M. Agholi, *Appl. Organomet. Chem.*, 2019, **33**, e4799.
- 64 F. Menges, *Spectragryph – optical spectroscopy software, Version 1.2.9*, 2019, <http://www.effemm2.de/spectragryph/>.
- 65 E. Gómez, N. Calvar and Á. Domínguez, in *Ionic Liquids - Current State of the Art*, ed. S. Handy, InTech, London, 2015, pp. 199–228.

