

Article

Polymerizable Ionic Liquids for Solid-State Polymer Electrolytes

Robert Löwe^{1,2}, Thomas Hanemann^{1,2} and Andreas Hofmann^{1,*} 

¹ Institute for Applied Materials, Karlsruhe Institute of Technology, Hermann-von-Helmholtz-Platz 1, D-76344 Eggenstein-Leopoldshafen, Germany; robert.loewe@kit.edu (R.L.); thomas.hanemann@kit.edu (T.H.)

² Department of Microsystems Engineering, University of Freiburg, Georges-Köhler-Allee 102, D-79110 Freiburg, Germany

* Correspondence: andreas.hofmann2@kit.edu; Tel.: +49-721-608-25920

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Abstract: Eight new polymerizable ammonium-TFSI ionic liquids were synthesized and characterized with respect to an application in energy storage devices. The ionic liquids feature methacrylate or acrylate termination as polymerizable groups. The preparation was optimized to obtain the precursors and ionic liquids in high yield. All products were characterized by NMR and IR spectroscopy. Phase transition temperatures were obtained by DSC analysis. Density, viscosity and ionic conductivity of the ionic liquids were compared and discussed. The results reveal that the length of attached alkyl groups as well as the methyl group at the polymerizable function have significant influences on the ionic liquids physicochemical properties. Ionic conductivity values vary between 0.264 mS cm^{-1} for $[\text{C}_2\text{N}_{\text{A},22}]\text{TFSI}$ and 0.080 mS cm^{-1} for $[\text{C}_8\text{N}_{\text{MA},22}]\text{TFSI}$ at $25 \text{ }^\circ\text{C}$. Viscosity values are within a range of 0.762 Pa s for $[\text{C}_2\text{N}_{\text{A},22}]\text{TFSI}$ and 1.522 Pa s for $[\text{C}_6\text{N}_{\text{MA},22}]\text{TFSI}$ at $25 \text{ }^\circ\text{C}$.

Keywords: polymerizable ionic liquids; property-structure relationship; conductivity; viscosity; density

1. Introduction

Ionic liquids (ILs) possess a wide range of applications, being employed in CO_2 adsorption [1], catalysis [2] or as solvents [3]. They are promising candidates in drug design [4] and drug delivery systems [5] due to their high biological activity [6] and the possibility to combine bioactive cations with bioactive anions, e.g., the salicylate (Sal) ionic liquid (IL) $[\text{EMIM-OSal}][\text{Sal}]$ [7]. Within the field of energy storage systems, such as lithium ion batteries, ionic liquids are discussed as alternatives to conventional electrolytes [8]. Remarkable properties for electrochemical applications are low flammability, low vapor pressure, a broad electrochemical window (ca. 4–6 V) [8] and a high tunability of properties by structural variations and usage of IL mixtures.

For particular applications, immobilization of the ionic liquids is beneficial. In terms of drug design, a solid dosage form can lead to a higher convenience of products. In lithium ion batteries the electrolyte can move freely within the cell and thus lead to unwanted side reactions. A common example is the reactions of the aluminum current collector with bis(trifluoromethanesulfonyl)imide (TFSI) containing ionic liquids [9]. This type of degradation mechanism can be avoided if the electrolyte is fixed between the electrodes. Different approaches to immobilize ionic liquids are known. It is relatively easy to distinguish between materials in which the ILs are absorbed within a matrix [10] and materials in which the ionic liquids are fixed by covalent bonding [11,12]. The latter is an interesting approach for transferring the outstanding properties of ionic liquids to solid materials and covalent networks. In our research, we focus on ILs that feature acrylate or methacrylate functional groups for this particular purpose. Polymerization and reactions such as the alkene hydrothiolation are candidates to link the ILs to a covalent network.

A common method for the preparation of ammonium ionic liquids is a reaction pathway with two steps. In a first step, the tertiary amine is converted in a quaternary ammonium salt by Menshutkin reaction [13,14]. Within the second step, the halogen anion is exchanged for an organic anion, e.g., PF₆, BF₄ or TFSI [15–17]. Preparation of ionic liquids starting from the polymerizable amines 2-(diethylamino)ethyl methacrylate (DEAEMA) and 2-(diethylamino)ethyl acrylate (DEAEA) is rarely described in the literature [18]. However, there are few reports on the synthesis of the halogen precursors by Menshutkin reaction [13,14]. It is known that the turnover of DEAEA to the desired precursor lowers drastically when the quaternization reaction is performed with alkyl bromides containing more than two carbon atoms. Regarding to the literature, yields decrease from 62% (DEAEA + C₂H₅Br) [19] to 13% (DEAEA + C₄H₉Br) [19] and 2.1% (DEAEA + C₆H₉Br) [18]. This is consistent with the general trend that is found for the reaction rate constant of S_N2 reactions which decreases with increasing size of the alkylating agent [20]. In terms of reaction mechanism, the Menshutkin reaction can be seen as a special type of S_N2 reaction [21]. In comparison to other S_N2 type reactions, the reactions rate is promoted by increasing solvent polarity [21]. Synthesis of similar ionic liquids starting from the methyl substituted amines 2-(dimethylamino)ethyl methacrylate (DMAEMA) or 2-(dimethylamino)ethyl acrylate (DMAEA) were described by several authors [18,22]. In the literature, alkyl bromides and acetone or chloroform were used as reagent and solvent, respectively, thus resulting in high yields for the precursors, e.g., 89% for [C₄N_{MA,11}]Br [22] and 82.6% for [C₆N_{MA,11}]Br [18]. In contrast to the earlier mentioned precursors, the yields for the quaternization of DMAEMA and DMAEA are still reasonable for alkyl substituents with more than two carbon atoms.

For application in energy storage systems the relationship between structure variations and fundamental properties such as glass transition temperature, viscosity and ionic conductivity are of great interest. Two common models for data interpretation are the Arrhenius equation (1) and the Vogel–Fulcher–Tammann (VFT) Equation (2)

$$\eta = A_{\eta}e^{E_{\eta}/RT}; \quad \sigma = A_{\sigma}e^{-E_{\sigma}/RT} \quad (1)$$

$$\eta = A_{\eta}e^{k_{\eta}/(T-T_0)}; \quad \sigma = A_{\sigma}e^{-k_{\sigma}/(T-T_0)} \quad (2)$$

with A_{η} and A_{σ} as the viscosity and conductivity at infinite temperature, E_{η} and E_{σ} as flow and conductivity activation energies, the ideal gas constant R , the temperature T , k_{η} and k_{σ} as constants and T_0 as ideal glass transition temperature [23]. In the literature the ideal parameter T_0 is described as temperature where free volume disappears so that cooperative movement of chain segments, as well as motion related phenomena such as diffusion or migration of molecules and ions becomes impossible [23]. Due to kinetic reasons, this state cannot be reached within a limited time scale and T_0 is found to be lower than experimentally measured values for T_g [23]. Correlation of the different fundamental properties offers further possibilities for analysis and interpretation of the experimental results. Scaling the Arrhenius plot to experimentally measured T_g values leads to the Arrhenius plot of fragilities or so-called Angell plot. Thereby, glass builders are classified as *strong* or *fragile*. The liquid structure of *fragile* materials responses very sensitive to temperature changes, which is associated with a relative narrow glass transition temperature range. *Strong* glass builders response in an opposite way [24,25]. Another interpretation is made by the Walden plot. In 1906 Paul Walden released his underlying empirically found relationship, that the product of viscosity of a solvent and its molar conductivity is constant [26]. Within the plot, experimental data are related to a line of an ideal electrolyte solution, which commonly is a 1-molar KCl solution. In the field of ionic liquids, the position of the ILs line is compared to the ideal line to classify the ILs in *superionic* liquids, *good* ILs and *poor* ILs concerning the decoupling of one charge carrier from the viscosity determined structure [27]. In the case of *superionic* the ILs line is located above the ideal line, which leads to the interpretation that one charge carrier species is partially decoupled from the viscous bulk. Slightly below and directly on the ideal line, ILs are classified as *good* ionic liquids featured by a quasi-lattice state and a low ionicity. The more the ILs line is below the ideal line, the more this state is shifted to direct ion pairing and higher ionicities [27].

In this study, we focus on the systematic preparation and characterization of polymerizable ionic liquids starting from DEAEMA and DEAEA. First, we describe the optimization of the reaction conditions to achieve almost quantitative precursor yields even for steric demanding alkyl chains and the adapted anion exchange. Subsequently, the characterization of the IL products is presented with respect to an application of the ionic liquids in energy storage systems. Structure-property relationships of the monomeric ILs are analyzed and their adaptability on polymerized ionic liquids are discussed.

2. Results and Discussion

2.1. Synthesis

Eight new ionic liquids were prepared in a two-step synthesis, shown in Figure 1. In the first step the amines DEAEMA and DEAEA were converted into quaternary ammonium bromide or iodide salts using the Menshutkin reaction [13,14]. In a following metathesis reaction, the halogen anions were exchanged by the organic anion bis(trifluoromethanesulfonyl)imide (TFSI).

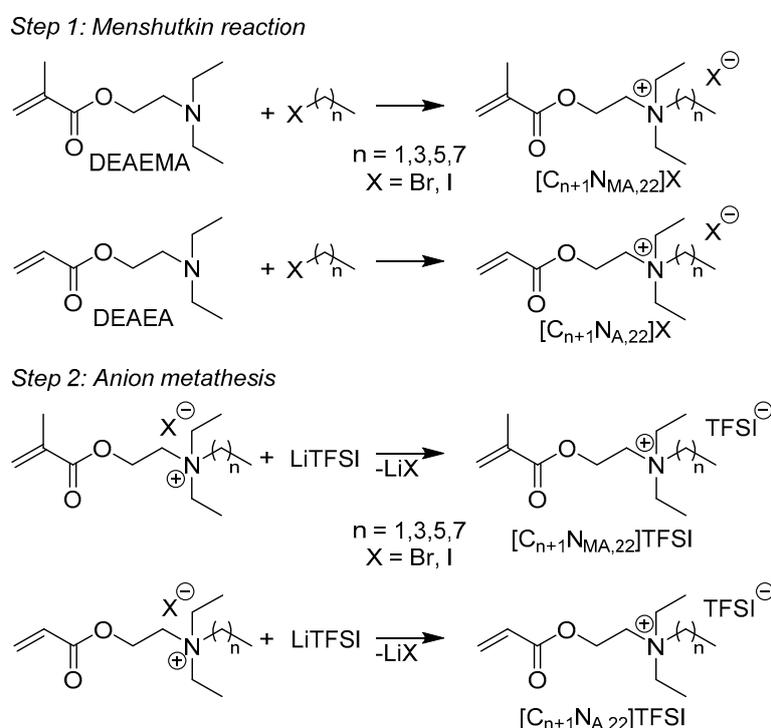


Figure 1. Reaction scheme for the preparation of the IL and notation.

Compared to the dimethyl substituted ammonium ionic liquids known from literature [11,22], two ethyl substituents are attached to the nitrogen atom in DEAEMA and DEAEA. This variation is associated with a higher steric hindrance at the amine function. We found that the yields of the quaternization reaction to bromide precursors are much lower when comparable reaction conditions from the literature were used with DEAEMA and DEAEA as starting materials. With increasing lengths of alkyl chains, the yields dropped drastically and fell below a reasonable level for C₄–C₈ rests. In order to achieve the quaternary ammonium salts in high yields, different reaction conditions were applied. Table 1 gives an overview of selected reactions. To avoid declining yields with increasing chain lengths, the reaction time and temperature were increased, thus leading to a partial degradation of the double bonds. Referring to Daoudi et al., the reaction of DEAEMA with *n*BuBr in dichloromethane was supposed to proceed almost quantitatively when the mixture was heated 8 h under reflux [28]. Applying similar conditions (16 h of reflux), the reaction failed and no product was obtained. Yields were significantly enhanced by the combination of alkyl iodides and acetone as reactants and solvent, respectively, as well as

by the usage of acetonitrile as solvent when alkyl bromides were used. In S_N2 type reactions iodide is the better leaving group compared to bromide, due to its lower basicity. Solvents with higher polarity affect the Menshutkin reaction positively [21]. Accordingly, the combination of acetonitrile as solvent and alkyl iodides as reactants led to almost quantitative yields, while the excess of alkyl halides was reduced.

Table 1. Selected reaction conditions and yields of the intermediate products.

Product	Solvent	Reaction Time	Temperature	Alkyl Halide Excess	Yield
[C ₂ N _{MA,22}]Br	Acetone	6 d	47.0 °C	1.4	60.7%
[C ₄ N _{MA,22}]Br	Acetone	6 d	47.0 °C	1.4	7.0%
[C ₆ N _{MA,22}]Br	Acetone	11 d	56.0 °C	1.4	51.5% ^a
[C ₄ N _{MA,22}]Br	CH ₂ Cl ₂	16 h	39.6 °C	1.2	0.0%
[C ₆ N _{MA,22}]I	Acetone	5 d	56.0 °C	1.4	66.3%
[C ₆ N _{MA,22}]Br	Acetonitrile	6 d	56.0 °C	1.4	54.0%
[C ₂ N _{MA,22}]I	Acetonitrile	1 d	45.0 °C	1.1	96.4%
[C ₄ N _{MA,22}]I	Acetonitrile	3 d	60.0 °C	1.1	97.5%
[C ₆ N _{MA,22}]I	Acetonitrile	3 d	60.0 °C	1.1	95.9%

^a ¹H NMR spectroscopy revealed that approx. 25% of the double bonds were degraded during the reaction.

All synthesized ammonium bromide compounds are strongly hygroscopic and feature a high solubility in water. Hence, using bromide intermediates, the anions exchange was executed in an aqueous solution from which the desired product precipitates. The synthesized iodide intermediates show a significantly lower hygroscopy and are therefore easier to handle under ambient conditions. A drawback compared to bromides is the accompanied decrease in water solubility, which makes a metathesis in pure water difficult for most of the precursor iodides. Instead the meth(acrylate) iodides were dissolved in acetone and added dropwise to an aqueous LiTFSI solution from which the IL phase precipitated immediately. The attempt to remove undesirable residues by extraction of the ionic liquids with chloroform and washing the organic phase with water points out to be impractical. The solubility of the ILs in chloroform is strongly limited and two organic phases appeared. The high viscosity and adhesion on the glass surface of the ionic liquid enriched phases, makes an effective separation from the aqueous phase almost impossible, when a separation funnel is used. In contrast, washing the IL monomers directly several times with purified water and decanting the aqueous phase, removed the remaining acetone and lithium halide salts from the products in an easy and effective manner. Table 2 gives an overview of the achieved yields concerning the two-step synthesis with ammonium iodide salts as intermediate products.

Table 2. Particular yields of all reaction steps and resulting overall yields for the ILs.

Precursor	Yield [%]	Ionic Liquid	Yield [%]	Overall Yield [%]
[C ₂ N _{MA,22}]I	96.4	[C ₂ N _{MA,22}]TFSI	90.7	87.4
[C ₄ N _{MA,22}]I	97.5	[C ₄ N _{MA,22}]TFSI	88.8	86.6
[C ₆ N _{MA,22}]I	95.9	[C ₆ N _{MA,22}]TFSI	91.7	87.9
[C ₈ N _{MA,22}]I	93.2	[C ₈ N _{MA,22}]TFSI	90.7	84.5
[C ₂ N _{A,22}]I	98.5	[C ₂ N _{A,22}]TFSI	85.0	83.7
[C ₄ N _{A,22}]I	96.6	[C ₄ N _{A,22}]TFSI	80.5	77.8
[C ₆ N _{A,22}]I	96.1	[C ₆ N _{A,22}]TFSI	89.5	86.0
[C ₈ N _{A,22}]I	89.4	[C ₈ N _{A,22}]TFSI	89.3	79.8

To accelerate the preparation of the ILs, a one-pot synthesis for [C₂N_{MA,22}]TFSI was carried out, by way of example. Without optimization of reaction conditions, a total yield of 66% was achieved, making the one-pot synthesis easily accessible to industrial scales.

It was found in the study that for long-term-storage the ionic liquids must be handled with an inhibitor to avoid any polymerization. An appropriate inhibitor which can be used when oxygen is present in the sample is MEHQ (Hydroquinone monomethyl ether) [29].

2.2. Density

Density values of all prepared TFSI-ionic liquids were measured between 5 °C and 60 °C. The resulting data are plotted in Figure 2 and fitted to a linear function $y = m \times x + n$. The calculated parameters are presented in Table 3. Density values of the acrylates are higher than the values of the methacrylate ILs with respect to the same alkyl chain lengths. With an increasing length of the attached alkyl group, density values decrease. Longer chains and the variation from acrylate to methacrylate groups lead to increased bulkiness and steric demand of the molecules, which is expressed in lower density values.

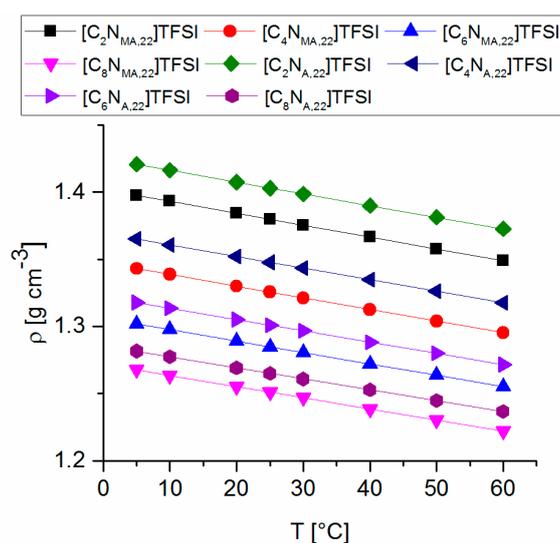


Figure 2. Density values of the synthesized TFSI-Ionic Liquids.

Table 3. Parameters from linear fit of density values ($y = m \times x + n$).

IONIC Liquid	m	$n/\text{g}\cdot\text{cm}^{-3}$
$[\text{C}_2\text{N}_{\text{MA},22}]\text{TFSI}$	-8.86882×10^{-4}	1.40202
$[\text{C}_4\text{N}_{\text{MA},22}]\text{TFSI}$	-8.70826×10^{-4}	1.34746
$[\text{C}_6\text{N}_{\text{MA},22}]\text{TFSI}$	-8.44608×10^{-4}	1.30603
$[\text{C}_8\text{N}_{\text{MA},22}]\text{TFSI}$	-8.29652×10^{-4}	1.27181
$[\text{C}_2\text{N}_{\text{A},22}]\text{TFSI}$	-8.76545×10^{-4}	1.42503
$[\text{C}_4\text{N}_{\text{A},22}]\text{TFSI}$	-8.64216×10^{-4}	1.36936
$[\text{C}_6\text{N}_{\text{A},22}]\text{TFSI}$	-8.40463×10^{-4}	1.32206
$[\text{C}_8\text{N}_{\text{A},22}]\text{TFSI}$	-8.18980×10^{-4}	1.28568

As the linear fitting shows, the lines are almost parallel. The slopes indicate a tendency to flatten with increasing alkyl chain length. The parameter n shows the differences between the calculated densities at 0 °C, which decrease with increasing chain length towards smaller density values. Density values of new ionic liquids of the same homologous series can be estimated on basis of the presented interpretation.

2.3. Thermal Analysis

DSC analysis was applied to the halogen salts and TFSI-ionic liquids. Data of the melting points result from a single measurement. For glass transition temperatures the probes were heated twice from -150 °C to 180 °C with 10 K min^{-1} . The given T_g values are the values of detected inflection points of the first occurring glass transition ($T_g(L)$) for each substance at the first heating curve.

As shown in Figure 3, two second order phase transitions, $T_g(L)$ and $T_g(U)$, appear in DSC analysis. $T_g(L)$ refers to the conventional glass transition and is therefore labeled as T_g in the further text. The appearance of $T_g(U)$ might be associated with semi-crystalline, smectic or para-crystalline phases as discussed in literature for semi-crystalline polymers [30].

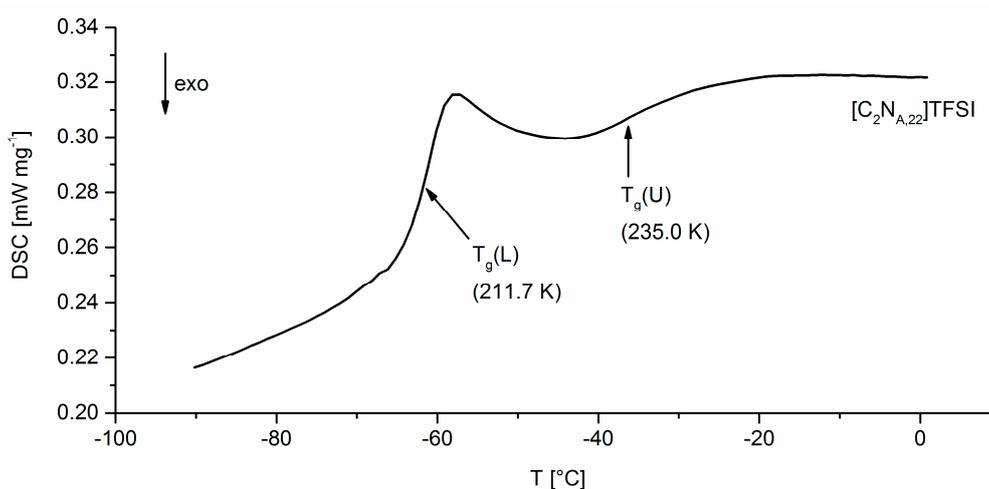


Figure 3. DSC heating curve of $[C_2N_{A,22}]TFSI$ which stands exemplary for all synthesized ILs.

Table 4 shows the resulting data of DSC analysis and gives two different values for T_g ($T_g(L)$). The $T_{g,1}$ value was measured two weeks after synthesis in a single probe analysis. The second value, $T_{g,2}$, is the average glass transition temperature of three probes, which were of the same batch, but stored eight weeks before DSC analysis was applied. All freshly synthesized ionic liquids were diluted in acetone (50 wt%) and stored at -40 °C. Before any analysis, the acetone was completely removed under vacuum at room temperature within an argon glove box.

The data indicate that the intermediates melting points decrease with increasing length of the introduced alkyl chains, except $[C_4N_{MA,22}]I$. The acrylate iodides show lower melting points than the methacrylate iodides. The studied bromid salt intermediates feature higher melting points than the analogous iodides.

The glass transition temperatures of the ionic liquids tend to decrease with increasing alkyl length. It is remarkable that the $T_{g,2}$ values are in average 4.3 K higher than $T_{g,1}$ values. There is evidence that this observation is due to partial polymerization processes, although the ionic liquids are diluted in acetone (50 wt%) and stored at -40 °C. The process of double bond degradation is also seen on the morphology of the ionic liquids after the acetone is removed. In contrast to freshly prepared ionic liquids, the probes start to thread after storing them several weeks. For analysis in Arrhenius plot of fragilities $T_{g,1}$ values are used due to the evidence that these values are less influenced by autopolymerization.

Table 4. Melting points (T_m) and glass transition temperatures (T_g) of the products. Standard uncertainties u for values from this work are $u(T_m) = 2$ K; $u(T_g) = 2$ K.

Halogen Intermediate Salt			Ionic Liquid				
Entry	Product	Onset [K]	Entry	Product	$T_{g,1}$ [K] ¹	$T_{g,2}$ [K] ²	Standard Deviation ² s
1	$[C_2N_{MA,22}]I$	339.9	12	$[C_2N_{MA,22}]TFSI$	219.2	223.0	0.3
2	$[C_4N_{MA,22}]I$	353.3	13	$[C_4N_{MA,22}]TFSI$	216.8	223.2	1.2
3	$[C_6N_{MA,22}]I$	332.9	14	$[C_6N_{MA,22}]TFSI$	215.2	220.4	1.0
4	$[C_8N_{MA,22}]I$	317.7	15	$[C_8N_{MA,22}]TFSI$	213.2	218.4	0.2
5	$[C_2N_{A,22}]I$	330.3	16	$[C_2N_{A,22}]TFSI$	211.7	215.5	0.9
6	$[C_4N_{A,22}]I$	321.2	17	$[C_4N_{A,22}]TFSI$	214.5	218.2	1.1
7	$[C_6N_{A,22}]I$	318.3	18	$[C_6N_{A,22}]TFSI$	211.6	216.1	0.4
8	$[C_8N_{A,22}]I$	291.6	19	$[C_8N_{A,22}]TFSI$	212.6	214.4	0.6
9	$[C_2N_{MA,22}]Br$	348.4					
10	$[C_4N_{MA,22}]Br$	345.1					
11	$[C_6N_{MA,22}]Br$	339.9					

¹ Measurement two weeks after synthesis and storage at -40 °C diluted in acetone (50 wt%). Acetone was completely removed under vacuum before measurement. ² Measurements after eight weeks of storage at same conditions as before.

2.4. Rheological Behavior

The viscosity of the ionic liquid monomers was studied in two manners. First, a constant temperature of 20 °C and varying shear rates from 1 s⁻¹ to 250 s⁻¹ were applied, resulting in shear rate independent viscosity for all the prepared ILs (within 1 s⁻¹ to 250 s⁻¹). Second, a temperature program from 20 °C to 100 °C at a constant shear rate of 100 s⁻¹ was applied. The dependence of the viscosity values on the temperature is given in Figure 4.

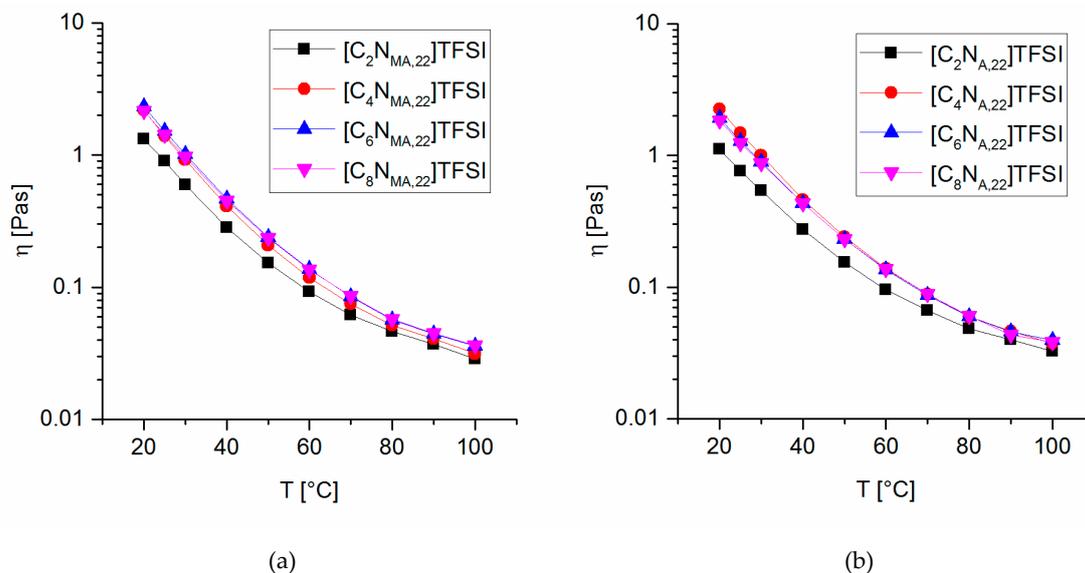


Figure 4. Viscosity values at various temperatures for methacrylate ILs (a) and acrylate ILs (b).

The results reveal that the ethyl-substituted ILs have the lowest viscosity values within the studied temperature range. Butyl- to octyl-substituted derivatives show low divergence in the viscosity values and temperature dependence and thus, no correlation between the chain lengths and the viscosity values was found for these derivatives. In general, the methacrylate ionic liquids tend to higher viscosities than the analogous acrylates at same temperatures. The data indicate that bulky groups within the ILs promote higher viscosity values. Phenomena such as entanglements increase the internal friction and are most likely to be causal for this observation. Tokuda et al. found the same property-structure relationship for imidazolium TFSI ionic liquids and correlated it to the influence on the ionic self-diffusivity [31].

2.5. Electrochemical Characterization

The ionic conductivity of the TFSI ionic liquids was calculated from the corresponding impedance spectra. The closed, airtight measuring cells for the liquid monomers were assembled inside an argon-purged glovebox (O_2 and $H_2O < 0.5$ ppm). Electrochemical impedance spectroscopy experiments were done at temperatures between -40 °C and 100 °C. During the measurements, the cells were located inside an oven. After the adjusted temperatures reached the desired values, further 20 min were given to the system before measurement to ensure a steady state. Voltage for potentiostatic impedance spectroscopy was set to 0.000 V. Amplitudes of 10 mV were applied with varying frequencies within the range of 100 Hz to 1 MHz. Figure 5 shows the temperature dependency of the ionic conductivity values. From 100 °C to 10 °C, the conductivity values decrease logarithmically. Below 5 °C there is an abrupt flatten of the conductivity curves, which indicates a phase transition within the given temperature range. This is consistent with the appearance of the upper glass transition temperature, $T_g(U)$, in DSC. Due to the limitation of ionic mobility, the conductivity values strongly approach $\sigma(T_g)$ with onset of the upper phase transition.

There is a significant impact of the structural modifications on the ionic conductivity, showing a straight structure-property relationship. It was found, that increasing chain lengths of the introduced alkyl group correlate with lower ionic conductivities. Furthermore, the acrylates show higher conductivity values than the methacrylate derivatives at the same alkyl chain lengths and temperatures. The conductivities at 25 °C ranging from 0.264 mS cm⁻¹ ([C₂N_{A,22}]TFSI) down to 0.080 mS cm⁻¹ ([C₈N_{MA,22}]TFSI), as given in Table 5.

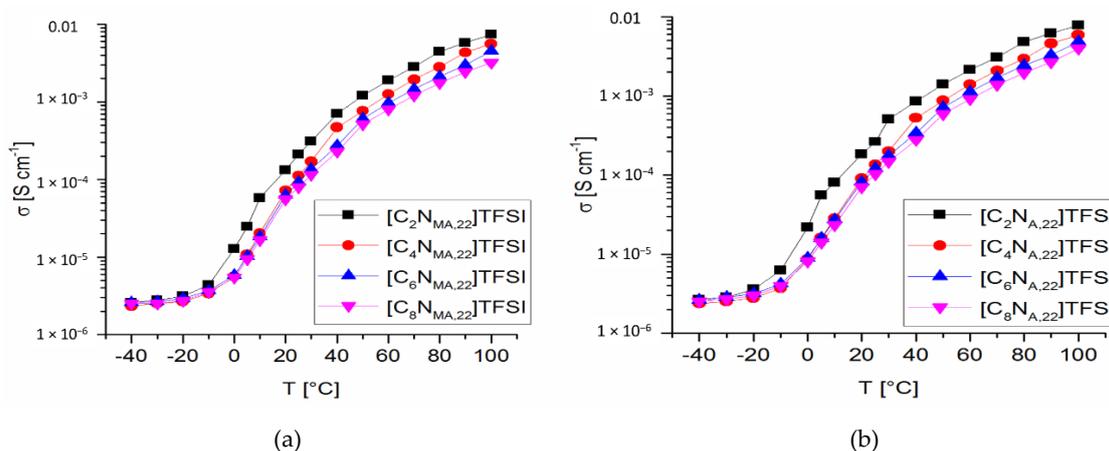


Figure 5. Ionic conductivity values at various temperatures for methacrylate ILs (a) and acrylate ILs (b).

Table 5. Experimentally measured values of ionic conductivity (κ), density (d), viscosity (η) and calculated molar conductivity (Λ_m) at 25 °C and at pressure $p = 0.1$ MPa. Standard uncertainties u for values from this work are $u(d) = 0.001$ g cm⁻³; $u(\eta) = 0.05 \cdot \eta$; $u(\kappa) = 0.03 \cdot \kappa$; $u(p) = 5$ kPa; $u(\Lambda_m) = 0.06 \cdot \Lambda_m$; $u(T_\kappa) = 0.1$ K; $u(T_\eta) = 0.1$ K; $u(T_d) = 0.01$ K (specified temperature during conductivity (T_κ), density (T_d) and viscosity (T_η) measurement).

Ionic Liquid	Ionic Conductivity [mS cm ⁻¹]	Molar Conductivity [mS cm ² mol ⁻¹]	Density [g cm ⁻³]	Viscosity [Pa s]
[C ₂ N _{MA,22}]TFSI	0.211	75.7	2.790	0.905
[C ₄ N _{MA,22}]TFSI	0.112	44.2	2.537	1.395
[C ₆ N _{MA,22}]TFSI	0.093	39.8	2.334	1.522
[C ₈ N _{MA,22}]TFSI	0.080	36.8	2.162	1.425
[C ₂ N _{A,22}]TFSI	0.264	90.3	2.920	0.762
[C ₄ N _{A,22}]TFSI	0.136	51.1	2.650	1.477
[C ₆ N _{A,22}]TFSI	0.122	50.3	2.425	1.284
[C ₈ N _{A,22}]TFSI	0.103	46.1	2.241	1.246

2.6. Analysis of Structure-Property Relationships

The variation between acrylate and methacrylate terminal function as well as the length of the introduced alkyl chain significantly affect the ionic liquids physicochemical properties. The measured densities decrease with increasing chain lengths and steric demand of the side chains, to remain the same trends found for the ionic conductivity on the molar conductivity. Ionic conductivity values decrease and viscosity values tend to increase with increasing steric demand of the attached side chains. Similar influences of structural variations on density, rheological behavior and conductivity were reported in the literature for imidazolium TFSI ILs, indicating a general trend for structure-property relationship of ionic liquids [31].

Considering the influence of the attached alkyl chains on the conductivity and viscosity values in Table 5, it turns out that the step from C₂ to C₄ chains is accompanied with a more pronounced property change than any other step in chain prolongation. For both homologous series, acrylates and methacrylates, the ionic conductivity drops almost 50% from C₂ to C₄ chains, whereas the step from C₄ to C₈ rests leads to a decrease in the order of 25% to 30%. The trend found for the viscosity values shows a similar behavior.

There is a significant increase in viscosity when going from C₂ to C₄ groups. From C₄ to C₈ side chains there is no straight property-structure relationship identifiable and the occurring variations of viscosity values are much less pronounced than caused by the step from C₂ to C₄.

Since all studied ionic liquids are featured by the same anion, the variance in the chemico-physical values is primarily driven by the structural differences of the cations. On one side, longer and bulkier sidechains lead to a higher steric demand of the molecules on the molecular level, thus resulting in lower densities. On the other side, higher densities cause stronger intermolecular forces, concerning anion-cation interactions as well as attractive forces between the organic cations, such as aliphatic π -interactions. As expected from this consideration, it was found that the glass transition temperatures tend to decrease with increasing length of the substituted alkyl groups due to higher free volumes, but contradicts the influence of the methyl group at the double bond. Table 6 shows that the studied acrylate ILs feature lower glass transition temperatures than the respective methacrylates. Therefore, it is assumed that the higher steric freedom of the acrylate function without attached methyl group has a more dominant influence on T_g than stronger electrostatic interactions between the cations.

The introduced alkyl chains are accompanied with an electron donating inductive effect. Longer alkyl chains will lead to a reduction of charge density at the cationic center. Thus, induce a more independent mobility of cations and anions and result in a higher ionic conductivity. Experimentally the opposite observation was made on the ionic conductivity. Therefore, it is stated that the electrostatic effect of alkyl chains has a negligible impact on the ionic conductivity. The adverse influence of longer chains on the cations mobility is most likely to limit the conductivity in impedance spectroscopy experiments.

The influence of the substituents steric demand on the cations mobility can be derived by the comparison of viscosity values of the different ILs. Figure 6 shows the empirical analysis of the viscosity values in terms of an Arrhenius plot (equation 1). The data between 20 °C and 60 °C were linear fitted and the flow activation energy was calculated within this temperature range. The results are given in Table 7 and show the same behavior as discussed for the viscosity values.

Table 6. Comparison of experimentally determined T_g and calculated T_0 from VFT fit. Standard uncertainties u for values from this work are $u(T_g) = 2$ K.

Ionic Liquid	T_g [K] ^{1,2}	T_0 [K] ³	T_g/T_0
[C ₂ N _{MA,22}]TFSI	219.2	201.8	1.09
[C ₄ N _{MA,22}]TFSI	216.8	208.7	1.04
[C ₆ N _{MA,22}]TFSI	215.2	197.7	1.09
[C ₈ N _{MA,22}]TFSI	213.2	200.0	1.07
[C ₂ N _{A,22}]TFSI	211.7	186.8	1.13
[C ₄ N _{A,22}]TFSI	214.5	202.7	1.06
[C ₆ N _{A,22}]TFSI	211.6	193.7	1.09
[C ₈ N _{A,22}]TFSI	212.6	190.6	1.11

¹ Experimentally measured values are provided. ² The measurement was performed two weeks after synthesis and storage at -40 °C diluted in acetone (50 wt%). Acetone was completely removed under vacuum before measurement. ³ Values are calculated from VFT fits of ionic conductivity values (Figure 7b).

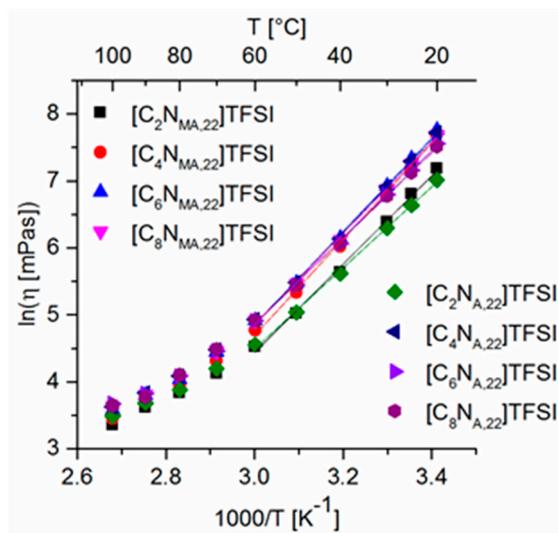


Figure 6. Arrhenius fit of viscosity values.

Table 7. Calculated Flow activation energies from Arrhenius fit are shown (20 °C–60 °C).

Ionic Liquid	E_{η} [kJ mol ⁻¹]
[C ₂ N _{MA,22}]TFSI	54.9 ± 1.5
[C ₄ N _{MA,22}]TFSI	59.7 ± 1.3
[C ₆ N _{MA,22}]TFSI	58.1 ± 1.1
[C ₈ N _{MA,22}]TFSI	56.7 ± 1.0
[C ₂ N _{A,22}]TFSI	50.3 ± 1.0
[C ₄ N _{A,22}]TFSI	57.1 ± 1.1
[C ₆ N _{A,22}]TFSI	54.0 ± 0.9
[C ₈ N _{A,22}]TFSI	53.1 ± 0.8

With respect to the experimentally measured glass transition temperatures and the viscosities, Figure 7a represents the Arrhenius plot of fragility. Due to the analysis, all studied ionic liquids are classified as *fragile*. They react sensitive to thermal excitation and thus show a narrow temperature range of glass transition.

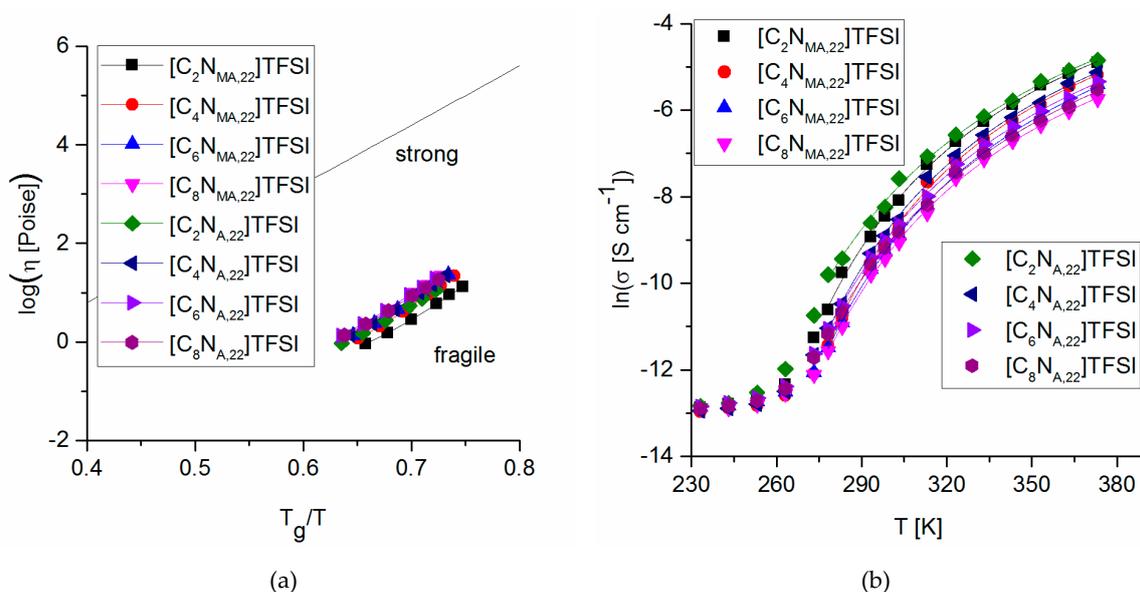


Figure 7. (a) Arrhenius plot of fragilities; (b) VFT fit of ionic conductivity values.

Empirical description of ionic conductivity values is done by VFT fit shown in Figure 7b (Equation (2)). The analysis was initialized by iterative non-linear least squares to give values for T_0 (Table 6). Due to this procedure, the ionic conductivity data fit well to the VFT equation in a temperature range between 5 °C and 100 °C. Below this range, conductivity values approach the conductivity at T_g . As expected, T_0 values are below the measured T_g values and T_g/T_0 is within the same range as found for other TFSI-ionic liquids in the literature [23].

The correlation of molar conductivity and viscosity of the studied ionic liquids is plotted in a Walden plot. Figure 8 shows the resulting linear relationship, which indicates a completely dissociation of all ions in the system. The data for all ionic liquids are located slightly below the ideal Walden line, which implies that the charge carriers are coupled to the viscosity determining structure. In terms of the Walden plot, all products are classified as *good* ionic liquids. The lines of the studied ILs are very close to each other, which shows that there are only small differences in ionicities between the ILs. This supports the already discussed assumption that the strength of cation-anion interactions is not the causal driving force for the found structure-property relationships.

Regarding the application of the studied ionic liquids in electrolyte systems, acrylates with short alkyl chains seem to outperform methacrylates and more bulky substituents due to the lower viscosity and higher conductivity. Although they found similar behaviors for the ionic liquids, Zhang et al. showed that the polymer of undoped $[C_7N_{MA,11}]TFSI$ reveals a higher ionic conductivity than the $[C_4N_{MA,11}]TFSI$ polymer [22]. Therefore, the property-relationships of the pure ILs cannot be directly transferred on their polymers. Further, additional material properties such as strength and brittleness of the polymers and the influences of conducting salt and additives must be regarded separately for particular applications.

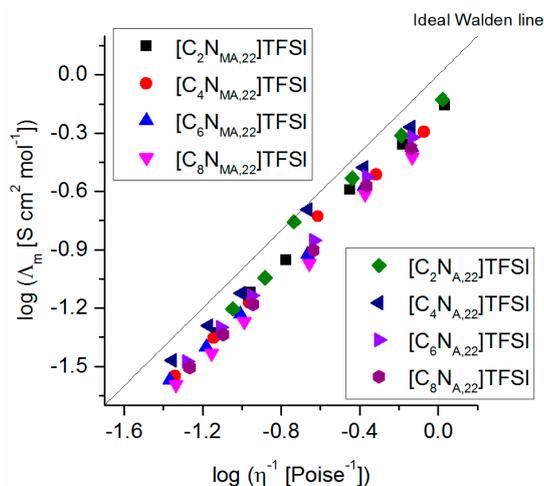


Figure 8. Walden plot for the studied TFSI ionic liquids.

In pretests, the monomers were successfully polymerized. For example, compound $[C_2N_{MA,22}]TFSI$ can be polymerized via UV induced bulk polymerization with 1 mol-% initiator (Irgacure 651) resulting in hard and brittle glasses. However, a broad range of parameters influences the polymeric matters properties. Thus, there are many parameters influencing the polymers properties starting with the general type of chosen polymerization method, type of initiator, initiator concentration and so on. Additionally, a huge toolbox of analytical methods for characterization, such as DSC, electrochemical impedance spectroscopy, cyclic voltammetry or size-exclusion chromatography, is available. With respect to an application in energy storage systems further parameters such as the type of applied conducting salt, salt concentration or additives come in addition. All these effects influence the structure and properties of the polymers. The presented results are focusing on the synthesis and characterization of the polymerizable monomers. In respect of its huge scope, the study of the polymerized ionic liquid materials will soon be presented separately.

3. Conclusions

In this study, we have synthesized and characterized eight new ionic liquids to improve the understanding of how structural changes affect the physicochemical properties of ionic liquids. Reaction conditions were optimized to obtain almost quantitative yields for the yield limiting quaternization reaction. The study reveals that an increasing side chain length at the ammonium moiety as well as and methyl group at the acrylate function accompany with increasing viscosity and decreasing ionic conductivity values. At 25 °C the ionic conductivity of the studied ionic liquids varies between 0.264 mS cm⁻¹ for [C₂N_{A,22}]TFSI and 0.080 mS cm⁻¹ and viscosity values between 0.762 Pa s [C₂N_{A,22}]TFSI and 1.522 Pa s [C₆N_{MA,22}]TFSI were found. Ionicity of the investigated ionic liquids shows low diversity in Walden plot. It is therefore assumed that the structure-property relationships of tetraalkylated ammonium ionic liquids are mainly driven by the steric properties of the cation on molecular level. It was found that the examined ionic liquids tend to autopolymerize when storing them diluted in acetone (50 wt%) at -40 °C. For the permanent usage at ambient conditions and elevated temperatures an inhibitor system will be required to prevent degradation of the double bonds. The data reveal that the polymerizable ionic liquids with short sidechains have favorable properties for an application in ionic conducting materials. Nevertheless, the literature indicate that the structure-property relationships found for the monomeric ionic liquids cannot be transferred on their polymers without further investigation.

4. Experimental Section

4.1. Materials

2-(Diethylamino)ethyl methacrylate (99%), 2-(diethylamino)ethyl acrylate (95%), bromoethane (98%), 1-bromobutane (99%), 1-bromohexane (≥99%), phenothiazine (≥98%) and bis(trifluoromethane)sulfonimide lithium salt (99%) were purchased from Sigma-Aldrich (Munich, Germany). Iodoethane (98%), 1-iodobutane (98%), 1-iodohexane (98%), 1-iodooctane (98%) and molecular sieves (3° Å, 8-12 mesh) were received from Acros Organics (Geel, Belgium). Acetone (≥98.8%) and diethyl ether (anhydrous) were purchased from Fisher Scientific (Schwerte, Germany). Water (HPLC grade) and acetonitrile (≥99.7%; HPLC grade) were purchased from Carl Roth (Karlsruhe, Germany) and Alfa Aesar (Karlsruhe, Germany), respectively. All chemicals were used as received.

4.2. Measurements

NMR spectra were recorded using an *Avance III HD 500 MHz* spectrometer (Bruker, Germany) and ATR-FTIR spectra with a *Vertex 70* (Bruker, Germany). Dynamic viscosities of the ILs were measured with a *Gemini HR Nano* rheometer (Bohlin, UK) and cone-plate system (CP4/40). For impedance spectroscopy a *Zahner Zennium* potentiostat (Zahner, Germany) and *TSC 1600 Closed* measurement cells (rhd, Germany) located inside a *SH-261* oven (Espec, USA) were applied. The water content of the ILs was measured by Karl Fischer-titration with a *KF Coulometer 831* (Metrohm, Germany) coupled with a *KF Thermoprep 860* oven (Metrohm, Germany) at 180 °C and flow rate of 150 mL min⁻¹. Melting points and glass transition temperatures were measured by DSC analysis using a *DSC 204 F1 Phoenix* (Netzsch, Germany) with temperature programs between -150 °C and 180 °C and heating rates of 10 K min⁻¹. Elemental analysis was carried out with a *vario MICRO* (Elementar, Germany) and a *SARTORIUS M2P* was used as analytical balance. Densities of ionic liquids were measured with a *DMA 4500 M* (Anton Paar, Austria) between 0.0 °C and 60.0 °C.

4.3. Synthesis

All solid products are stored in amber glass flasks inside an argon-filled glovebox. All TFSI ionic liquids are diluted to 50 wt% in acetone under argon and stored under exclusion of light at -30 °C. After removing the acetone, the ILs show less than 10 ppm water in Karl Fischer-titration. For synthesis and washing of ionic liquids deionized water (HPLC grade) was used without exceptions.

***N,N,N*-Triethyl-*N*-[2-(methacryloyloxy)ethyl]ammonium bromide ([C₂N_{MA,22}]Br)**

A mixture of 2-(diethylamino)ethyl methacrylate (DEAEMA, 10.99 g, 59.32 mmol) and a excess of bromoethane (9.24 g, 84.84 mmol) in 30 mL acetone was stirred under argon-oxygen atmosphere (80:20% *v/v*) at 47 °C for 6 days and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum for 10 h. The product was received as a white powder (10.61 g, 60.8% yield). M.p.: 111.2 °C; ¹H NMR (500 MHz, CDCl₃): δ (ppm) 1.45 (t, 9H, CH₃); 1.95 (s, 3H, CH₃); 3.64 (q, 6H, CH₂); 3.97 (t, 2H, CH₂); 4.76 (t, 2H, CH₂); 5.68 (s, 1H, CH₂); 6.12 (s, 1H, CH₂); ¹³C NMR (125 MHz, CDCl₃): δ (ppm) 8.3 (CH₃); 18.3 (CH₂); 54.4 (CH₂); 56.0 (CH₂); 57.8 (CH₂); 127.5 (CH₂); 135.1 (C); 166.4 (C=O); IR (cm⁻¹): 2979 ν(C-H); 1717 ν(C=O); 1638 ν(C=C); 1479 δ_{as}(CH₂); 1390 δ_s(CH₃); 1320 δ(CH₂); 1298 ν(C-O); 1168 ν(C-O); 1002 δ(CH₂); 947 δ(C-H); 784 δ(CH₂);

***N*-Butyl-*N,N*-diethyl-*N*-[2-(methacryloyloxy)ethyl]ammonium bromide ([C₄N_{MA,22}]Br)**

A mixture of 2-(diethylamino)ethyl methacrylate (DEAEMA, 11.12 g, 60.00 mmol) and a excess of 1-bromobutane (11.51 g, 84.00 mmol) in 30 mL acetone was stirred under argon-oxygen atmosphere (80:20% *v/v*) under reflux for 11 days and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum for 24 h. The product was received as a white powder (9.49 g, 49.1% yield). M.p.: 107.9 °C; ¹H NMR (500 MHz, CDCl₃): δ (ppm) 1.02 (t, 3H, CH₃); 1.47 (m, 8H, CH₃, CH₂); 1.76 (m, 2H, CH₂); 1.96 (s, 3H, CH₃); 3.44 (m, 2H, CH₂); 3.67 (q, 4H, CH₂); 4.01 (t, 2H, CH₂); 4.69 (t, 2H, CH₂); 5.70 (s, 1H, CH₂); 6.14 (s, 1H, CH₂); ¹³C NMR (125 MHz, CDCl₃): δ (ppm) 8.4 (CH₃); 13.7 (CH₃); 18.3 (CH₃); 19.8 (CH₂); 24.1 (CH₂); 55.4 (CH₂); 56.6 (CH₂); 57.8 (CH₂); 58.9 (CH₂); 127.5 (CH₂); 135.1 (C); 166.5 (C=O); IR (cm⁻¹): 2963 ν(C-H); 1721 ν(C=O); 1640 ν(C=C); 1457 δ_{as}(CH₂); 1399 δ_s(CH₃); 1300 ν(C-O); 1159 ν(C-O); 1013 δ(CH₂); 934 δ(C-H); 788 δ(CH₂); Elemental analysis calcd (%) for C₁₄H₂₈BrNO₂: C, 52.18; H, 8.76; N, 4.35. Found (%): C, 51.59; H, 8.49; N, 4.41.

***N,N*-Diethyl-*N*-hexyl-*N*-[2-(methacryloyloxy)ethyl]ammonium bromide ([C₆N_{MA,22}]Br)**

A mixture of 2-(diethylamino)ethyl methacrylate (DEAEMA, 5.59 g, 30.02 mmol) and a excess of 1-bromohexane (6.93 g, 42.00 mmol) in 15 mL acetonitrile was stirred under argon-oxygen atmosphere (80:20% *v/v*) at 56 °C for 6 days and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum. The product was received as a white powder (5.67 g, 53.9% yield). M.p.: 102.7 °C; ¹H NMR (500 MHz, CDCl₃): δ (ppm) 0.87 (t, 3H, CH₃); 1.33 (m, 6H, CH₂); 1.43 (t, 6H, CH₃); 1.74 (m, 2H, CH₂); 1.93 (s, 3H, CH₃); 3.40 (m, 2H, CH₂); 3.64 (q, 4H, CH₂); 3.97 (t, 2H, CH₂); 4.65 (t, 2H, CH₂); 5.66 (s, 1H, CH₂); 6.11 (s, 1H, CH₂); ¹³C NMR (125 MHz, CDCl₃): δ (ppm) 8.3 (CH₃); 16.9 (CH₃); 18.3 (CH₃); 22.2 (CH₂); 22.4 (CH₂); 26.1 (CH₂); 31.2 (CH₂); 55.0 (CH₂); 56.6 (CH₂); 57.9 (CH₂); 59.0 (CH₂); 127.4 (CH₂); 135.1 (C); 166.4 (C=O); IR (cm⁻¹): 2928 ν(C-H); 1717 ν(C=O); 1640 ν(C=C); 1446 δ_{as}(CH₂); 1399 δ_s(CH₃); 1298 ν(C-O); 1163 ν(C-O); 1009 δ(CH₂); 945 δ(C-H); 813 δ(CH₂); Elemental analysis calcd (%) for C₁₆H₃₂BrNO₂: C, 54.85; H, 9.21; N, 4.00. Found (%): C, 54.73; H, 9.17; N, 4.01.

***N,N,N*-Triethyl-*N*-[2-(methacryloyloxy)ethyl]ammonium iodide ([C₂N_{MA,22}]I)**

A mixture of 2-(diethylamino)ethyl methacrylate (DEAEMA, 27.789 g, 150.0 mmol), a slight excess of iodoethane (25.735 g, 165.0 mmol) and phenothiazine (0.6 g, 3.0 mmol) as inhibitor in 15 mL acetonitrile was stirred under argon atmosphere at 45 °C for 24 h and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum. The product was received as a white powder (49.32 g, 96.4% yield). M.p.: 102.7 °C; ¹H NMR (500 MHz, acetone-D₆): δ (ppm) 1.46 (t, 9H, CH₃); 1.96 (s, 3H, CH₃); 3.73 (q, 6H, CH₂); 4.02 (t, 2H, CH₂); 4.72 (t, 2H, CH₂); 5.74 (s, 1H, CH₂); 6.14 (s, 1H, CH₂); ¹³C NMR (125 MHz, acetone-D₆): δ (ppm) 7.6 (CH₃); 17.6 (CH₂); 53.9 (CH₂); 55.6 (CH₂); 58.0 (CH₂); 126.1

(CH₂); 135.8 (C); 166.1 (C=O); IR (cm⁻¹): 2974 ν (C-H); 1719 ν (C=O); 1635 ν (C=C); 1457 δ_{as} (CH₂); 1397 δ_s (CH₃); 1300 ν (C-O); 1165 ν (C-O); 1000 δ (CH₂); 949 δ (C-H); 782 δ (CH₂); Elemental analysis calcd (%) for C₁₂H₂₄INO₂: C, 42.24; H, 7.09; N, 4.10. Found (%): C, 42.24; H, 7.11; N, 4.10.

N-Butyl-*N,N*-diethyl-*N*-[2-(methacryloyloxy)ethyl]ammonium iodide ([C₄N_{MA,22}]I)

A mixture of 2-(diethylamino)ethyl methacrylate (DEAEMA, 27.789 g, 150.0 mmol), a slight excess of 1-iodobutane (30.363 g, 165.0 mmol) and phenothiazine (0.6 g, 3.0 mmol) as inhibitor in 15 mL acetonitrile was stirred under argon atmosphere at 60 °C for 3 days and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum. The product was received as a slightly yellowish powder (54.02 g, 97.5% yield). M.p.: 116.1 °C; ¹H NMR (500 MHz, acetone-D₆): δ (ppm) 1.02 (t, 3H, CH₃); 1.49 (m, 8H, CH₃, CH₂); 1.90 (m, 2H, CH₂); 1.96 (t, 3H, CH₃); 3.62 (m, 2H, CH₂); 3.75 (q, 4H, CH₂); 4.00 (m, 2H, CH₂); 4.72 (m, 2H, CH₂); 5.75 (t, 1H, CH₂); 6.15 (s, 1H, CH₂); ¹³C NMR (125 MHz, acetone-D₆): δ (ppm) 7.4 (CH₃); 13.0 (CH₃); 17.5 (CH₃); 19.5 (CH₂); 23.7 (CH₂); 54.4 (CH₂); 56.0 (CH₂); 57.9 (CH₂); 58.2 (CH₂); 126.0 (CH₂); 135.8 (C); 166.0 (C=O); IR (cm⁻¹): 2963 ν (C-H); 1724 ν (C=O); 1640 ν (C=C); 1458 δ_{as} (CH₂); 1401 δ_s (CH₃); 1300 ν (C-O); 1163 ν (C-O); 1013 δ (CH₂); 960 δ (C-H); 813 δ (CH₂); Elemental analysis calcd (%) for C₁₄H₂₈INO₂: C, 45.53; H, 7.64; N, 3.79. Found (%): C, 45.48; H, 7.64; N, 3.73.

N,N-Diethyl-*N*-hexyl-*N*-[2-(methacryloyloxy)ethyl]ammonium iodide ([C₆N_{MA,22}]I)

A mixture of 2-(diethylamino)ethyl methacrylate (DEAEMA, 27.789 g, 150.0 mmol), a slight excess of 1-iodohexane (34.992 g, 165.0 mmol) and phenothiazine (0.6 g, 3.0 mmol) as inhibitor in 15 mL acetonitrile was stirred under argon atmosphere at 60 °C for 3 days and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum. The product was received as a white powder (57.18 g, 95.9% yield). M.p.: 95.7 °C; ¹H NMR (500 MHz, acetone-D₆): δ (ppm) 0.91 (t, 3H, CH₃); 1.39 (m, 6H, CH₂); 1.49 (t, 6H, CH₃); 1.92 (m, 2H, CH₂); 2.00 (s, 3H, CH₃); 3.62 (m, 2H, CH₂); 3.75 (q, 4H, CH₂); 4.01 (m, 2H, CH₂); 4.72 (s, 2H, CH₂); 5.75 (s, 1H, CH₂); 6.15 (s, 1H, CH₂); ¹³C NMR (125 MHz, acetone-D₆): δ (ppm) 7.4 (CH₃); 13.3 (CH₃); 17.5 (CH₃); 21.7 (CH₂); 22.2 (CH₂); 25.8 (CH₂); 31.1 (CH₂); 54.4 (CH₂); 56.0 (CH₂); 57.9 (CH₂); 58.4 (CH₂); 126.0 (CH₂); 135.8 (C); 166.0 (C=O); IR (cm⁻¹): 2954 ν (C-H); 1721 ν (C=O); 1640 ν (C=C); 1456 δ_{as} (CH₂); 1403 δ_s (CH₃); 1296 ν (C-O); 1157 ν (C-O); 1060 δ (CH₂); 1011 δ (CH₂); 936 δ (C-H); 808 δ (CH₂); Elemental analysis calcd (%) for C₁₆H₃₂INO₂: C, 48.37; H, 8.12; N, 3.53. Found (%): C, 48.43; H, 8.13; N, 3.50.

N,N-Diethyl-*N*-[2-(methacryloyloxy)ethyl]-*N*-octylammonium iodide ([C₈N_{MA,22}]I)

A mixture of 2-(diethylamino)ethyl methacrylate (DEAEMA, 27.789 g, 150.0 mmol), a slight excess of 1-iodooctane (39.621 g, 165.0 mmol) and phenothiazine (0.6 g, 3.0 mmol) as inhibitor in 15 mL acetonitrile was stirred under argon atmosphere at 75 °C for 3 days and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum. The product was received as a white powder (59.47 g, 93.2% yield). M.p.: 80.5 °C; ¹H NMR (500 MHz, acetone-D₆): δ (ppm) 0.88 (m, 3H, CH₃); 1.32 (m, 6H, CH₂); 1.46 (m, 10H, CH₃, CH₂); 1.91 (m, 2H, CH₂); 1.96 (s, 3H, CH₃); 3.64 (m, 2H, CH₂); 3.76 (q, 4H, CH₂); 4.06 (s, 2H, CH₂); 4.73 (s, 2H, CH₂); 5.74 (s, 1H, CH₂); 6.15 (s, 1H, CH₂); ¹³C NMR (125 MHz, acetone-D₆): δ (ppm) 7.9 (CH₃); 13.6 (CH₃); 17.7 (CH₃); 22.0 (CH₂); 22.4 (CH₂); 26.2 (CH₂); 29.0 (CH₂); 31.6 (CH₂); 54.6 (CH₂); 56.3 (CH₂); 58.2 (CH₂); 58.6 (CH₂); 126.1 (CH₂); 135.9 (C); 166.1 (C=O); IR (cm⁻¹): 2928 ν (C-H); 1724 ν (C=O); 1640 ν (C=C); 1454 δ_{as} (CH₂); 1399 δ_s (CH₃); 1291 ν (C-O); 1154 ν (C-O); 1060 δ (CH₂); 1009 δ (CH₂); 936 δ (C-H); 806 δ (CH₂); Elemental analysis calcd (%) for C₁₈H₃₆INO₂: C, 50.82; H, 8.53; N, 3.29. Found (%): C, 50.51; H, 8.48; N, 3.27.

N-[2-(Acryloyloxy)ethyl]-*N,N,N*-triethylammonium iodide (**[C₂N_{A,22}II**])

A mixture of 2-(diethylamino)ethyl acrylate (DEAEA, 25.686 g, 150.0 mmol), a slight excess of iodoethane (25.735 g, 165.0 mmol) and phenothiazine (0.6 g, 3.0 mmol) as inhibitor in 15 mL acetonitrile was stirred under argon atmosphere at 45 °C for 24 h and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum. The product was received as a white powder (48.33 g, 98.5% yield). M.p.: 93.1 °C; ¹H NMR (500 MHz, acetone-D₆): δ (ppm) 1.47 (t, 9H, CH₃); 3.72 (q, 6H, CH₂); 3.97 (m, 2H, CH₂); 4.73 (s, 2H, CH₂); 6.01 (dd, 1H, CH₂); 6.24 (m, 1H, CH); 6.44 (dd, 1H, CH₂); ¹³C NMR (125 MHz, acetone-D₆): δ (ppm) 7.3 (CH₃); 53.8 (CH₂); 55.4 (CH₂); 57.5 (CH₂); 127.8 (CH); 131.6 (CH₂); 164.8 (C=O); IR (cm⁻¹): 2979 ν(C-H); 1719 ν(C=O); 1624 ν(C=C); 1461 δ_{as}(CH₂); 1401 δ_s(CH₃); 1271 ν(C-O); 1183 ν(C-O); 1086 δ(CH₂); 1064 δ(CH₂); 1020 δ(CH₂); 998 δ(C-H); 960 δ(C-H); 808 δ(CH₂); Elemental analysis calcd (%) for C₁₁H₂₂INO₂: C, 40.38; H, 6.78; N, 4.28. Found (%): C, 40.61; H, 6.78; N, 4.52.

N-[2-(Acryloyloxy)ethyl]-*N*-butyl-*N,N*-diethylammonium iodide (**[C₄N_{A,22}II**])

A mixture of 2-(diethylamino)ethyl acrylate (DEAEA, 15.945 g, 93.1 mmol), a slight excess of 1-iodobutane (18.849 g, 102.4 mmol) and phenothiazine (0.6 g, 3.0 mmol) as inhibitor in 15 mL acetonitrile was stirred under argon atmosphere at 75 °C for 2 days and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum. The product was received as a slightly yellowish powder (31.96 g, 96.6% yield). M.p.: 84.0 °C; ¹H NMR (500 MHz, acetone-D₆): δ (ppm) 1.02 (t, 3H, CH₃); 1.47 (m, 8H, CH₃, CH₂); 1.89 (m, 2H, CH₂); 3.63 (m, 2H, CH₂); 3.75 (q, 4H, CH₂); 4.01 (m, 2H, CH₂); 4.74 (s, 2H, CH₂); 6.02 (m, 1H, CH₂); 6.24 (m, 1H, CH); 6.44 (m, 1H, CH₂); ¹³C NMR (125 MHz, acetone-D₆): δ (ppm) 7.6 (CH₃); 13.1 (CH₂); 19.5 (CH₂); 23.8 (CH₂); 54.5 (CH₂); 56.1 (CH₂); 57.7 (CH₂); 58.3 (CH₂); 127.8 (CH); 131.7 (CH₂); 164.8 (C=O); IR (cm⁻¹): 2965 ν(C-H); 1724 ν(C=O); 1622 ν(C=C); 1459 δ_{as}(CH₂); 1406 δ_s(CH₃); 1260 ν(C-O); 1188 ν(C-O); 1071 δ(CH₂); 1022 δ(CH₂); 965 δ(C-H); 806 δ(CH₂); Elemental analysis calcd (%) for C₁₃H₂₆INO₂: C, 43.95; H, 7.38; N, 3.94. Found (%): C, 44.18; H, 7.38; N, 4.00.

N-[2-(Acryloyloxy)ethyl]-*N,N*-diethyl-*N*-hexylammonium iodide (**[C₆N_{A,22}II**])

A mixture of 2-(diethylamino)ethyl acrylate (DEAEA, 25.686 g, 150.0 mmol), a slight excess of 1-iodohexane (34.992 g, 165.0 mmol) and phenothiazine (0.6 g, 3.0 mmol) as inhibitor in 15 mL acetonitrile was stirred under argon atmosphere at 60 °C for 3 days and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum. The product was received as a white powder (55.26 g, 96.1% yield). M.p.: 81.1 °C; ¹H NMR (500 MHz, acetone-D₆): δ (ppm) 0.91 (t, 3H, CH₃); 1.41 (m, 6H, CH₂); 1.48 (t, 6H, CH₃); 1.92 (m, 2H, CH₂); 3.62 (q, 2H, CH₂); 3.74 (s, 4H, CH₂); 3.99 (m, 2H, CH₂); 4.74 (s, 2H, CH₂); 6.02 (m, 1H, CH₂); 6.24 (m, 1H, CH); 6.45 (m, 1H, CH₂); ¹³C NMR (125 MHz, acetone-D₆): δ (ppm) 7.4 (CH₃); 13.3 (CH₂); 21.8 (CH₂); 22.2 (CH₂); 25.8 (CH₂); 31.1 (CH₂); 54.4 (CH₂); 56.0 (CH₂); 57.6 (CH₂); 58.5 (CH₂); 127.8 (CH); 131.6 (CH₂); 164.8 (C=O); IR (cm⁻¹): 2954 ν(C-H); 1721 ν(C=O); 1624 ν(C=C); 1456 δ(CH₂); 1406 δ_s(CH₃); 1263 ν(C-O); 1188 ν(C-O); 1071 δ(CH₂); 1022 δ(CH₂); 971 δ(C-H); 936 δ(C-H); 806 δ(CH₂); Elemental analysis calcd (%) for C₁₅H₃₀INO₂: C, 47.00; H, 7.89; N, 3.65. Found (%): C, 47.05; H, 7.89; N, 3.66.

N-[2-(Acryloyloxy)ethyl]-*N,N*-diethyl-*N*-octylammonium iodide (**[C₈N_{A,22}II**])

A mixture of 2-(diethylamino)ethyl acrylate (DEAEA, 25.686 g, 150.0 mmol), a slight excess of 1-iodooctane (39.621 g, 165.0 mmol) and phenothiazine (0.6 g, 3.0 mmol) as inhibitor in 15 mL acetonitrile was stirred under argon atmosphere at 60 °C for 3 days and under exclusion of UV irradiation. All volatile components were removed under reduced pressure. The precipitate was washed with ethyl ether and dried under vacuum. The product was received as a white powder

(55.16 g, 89.4% yield). M.p.: 54.4 °C; ^1H NMR (500 MHz, acetone- D_6): δ (ppm) 0.89 (t, 3H, CH_3); 1.33 (m, 6H, CH_2); 1.47 (m, 10H, CH_3 , CH_2); 1.92 (m, 2H, CH_2); 3.62 (m, 2H, CH_2); 3.75 (q, 4H, CH_2); 4.01 (m, 2H, CH_2); 4.74 (s, 2H, CH_2); 6.01 (m, 1H, CH_2); 6.24 (m, 1H, CH); 6.45 (m, 1H, CH_2); ^{13}C NMR (125 MHz, acetone- D_6): δ (ppm) 7.5 (CH_3); 13.5 (CH_2); 21.9 (CH_2); 22.4 (CH_2); 26.2 (CH_2); 28.9 (CH_2); 31.6 (CH_2); 54.5 (CH_2); 56.1 (CH_2); 57.7 (CH_2); 58.5 (CH_2); 127.8 (CH); 131.6 (CH_2); 164.8 (C=O); IR (cm^{-1}): 2926 ν (C-H); 1728 ν (C=O); 1640 ν (C=C); 1454 δ_{as} (CH_2); 1406 δ_{s} (CH_3); 1267 ν (C-O); 1188 ν (C-O); 1075 δ (CH_2); 1009 δ (CH_2); 978 δ (C-H); 804 δ (CH_2); Elemental analysis calcd (%) for $\text{C}_{17}\text{H}_{34}\text{INO}_2$: C, 49.64; H, 8.33; N, 4.40. Found (%): C, 49.46; H, 8.22; N, 3.35.

N,N,N-Triethyl-*N*-[2-(methacryloyloxy)ethyl]ammonium bis(trifluoromethanesulfonyl)imide
 ($[\text{C}_2\text{N}_{\text{MA},22}]\text{TFSI}$)

The respective intermediate $[\text{C}_2\text{N}_{\text{MA},22}]\text{I}$ (22.180 g, 65.0 mmol) was dissolved in water (15 mL) and added to a solution of LiTFSI (19.501 g, 67.9 mmol) in water (15 mL). The ionic liquid precipitated instantly and the mixture was stirred further for 10 min at room temperature. The aqueous top layer was removed and the remaining organic layer was washed six times with water (40 mL), respectively. No iodide ions were detected in the last washing water by AgNO_3 solution. The ionic liquid was diluted in acetone (40 mL) and dried over molecular sieves (10 g, 3 Å) for 24 h. The solution was filtrated (PTFE, 0.2 μm) and the acetone removed under reduced pressure. The ionic liquid was stirred at room temperature under vacuum (6 mbar) for 2 days to give the pure product as colorless oil (29.16 g, 90.7% yield). ^1H NMR (500 MHz, CDCl_3): δ (ppm) 1.28 (t, 9H, CH_3); 1.87 (s, 3H, CH_3); 3.32 (q, 6H, CH_2); 3.53 (m, 2H, CH_2); 4.45 (m, 2H, CH_2); 5.62 (m, 1H, CH_2); 6.03 (s, 1H, CH_2); ^{13}C NMR (125 MHz, CDCl_3): δ (ppm) 7.3 (CH_3); 18.0 (CH_2); 54.0 (CH_2); 55.3 (CH_2); 57.3 (CH_2); 120.1 (q, CF_3); 127.3 (CH_2); 135.1 (C); 166.4 (C=O); IR (cm^{-1}): 2996 ν (C-H); 1726 ν (C=O); 1638 ν (C=C); 1457 δ_{as} (CH_2); 1399 δ_{s} (CH_3); 1349 ν (SO_2); 1179 ν (C-O); 1135 ν (S=O); 1053 ν (S=O); 1002 δ (CH_2); 742 δ_{s} (CF_3); 654 δ (SNS); 614 δ_{a} (SO_2); Elemental analysis calcd (%) for $\text{C}_{14}\text{H}_{24}\text{F}_6\text{N}_2\text{O}_6\text{S}_2$: C, 34.01; H, 4.89; N, 5.67. Found (%): C, 34.11; H, 4.80; N, 5.73.

One-pot synthesis of $[\text{C}_2\text{N}_{\text{MA},22}]\text{TFSI}$

DEAEMA (3.149 g, 17.0 mmol) and 1-iodoethane (2.917 g, 18.7 mmol) were added to a solution of LiTFSI (5.100 g, 17.8 mmol) in acetonitrile (5 mL). The mixture was stirred at elevated temperature (45 °C) for 16 h. All volatile components were removed under reduced pressure. The remaining oily phase was washed with water (120 mL in total) in six portions. The IL phase was dissolved in acetone (15 mL) and the solution was dried over molecular sieves (2.5 g, 3 Å) for 3 days. The solution was filtrated (PTFE, 0.2 μm) and the acetone was removed under reduced pressure. The ionic liquid was stirred at room temperature under vacuum (6 mbar) to give the pure product as colorless oil (5.56 g, 66.2% yield).

N-Butyl-*N,N*-diethyl-*N*-[2-(methacryloyloxy)ethyl]ammonium bis(trifluoromethanesulfonyl)imide
 ($[\text{C}_4\text{N}_{\text{MA},22}]\text{TFSI}$)

The respective intermediate $[\text{C}_4\text{N}_{\text{MA},22}]\text{I}$ (24.003 g, 65.0 mmol) was suspended in water (30 mL) and the supernatant water was added to a solution of LiTFSI (19.501 g, 67.9 mmol) in water (15 mL). The undissolved intermediate salt was solved in acetone (15 mL) and added dropwise to the aqueous phase. The ionic liquid precipitated instantly and the mixture was stirred for 10 min at room temperature. Further processing was realized as described above for $[\text{C}_2\text{N}_{\text{MA},22}]\text{TFSI}$ to give the product as colorless oil (30.17 g, 88.8% yield). ^1H NMR (500 MHz, CDCl_3): δ (ppm) 0.94 (t, 3H, CH_3); 1.30 (t, 6H, CH_3); 1.36 (q, 2H, CH_2); 1.62 (m, 2H, CH_2); 1.90 (s, 3H, CH_3); 3.18 (m, 2H, CH_2); 3.35 (q, 4H, CH_2); 3.57 (m, 2H, CH_2); 4.47 (m, 2H, CH_2); 5.64 (s, 1H, CH_2); 6.06 (s, 1H, CH_2); ^{13}C NMR (125 MHz, CDCl_3): δ (ppm) 7.4 (CH_3); 13.3 (CH_3); 18.0 (CH_3); 19.4 (CH_2); 23.5 (CH_2); 54.5 (CH_2); 55.8 (CH_2); 57.2 (CH_2); 58.4 (CH_2); 119.8 (q, CF_3); 127.2 (CH_2); 135.0 (C); 166.4 (C=O); IR (cm^{-1}): 2970 ν (C-H); 1724 ν (C=O); 1638 ν (C=C); 1459 δ_{as} (CH_2); 1349 ν (SO_2); 1181 ν (C-O); 1135 ν (S=O); 1053 ν (S=O); 949 δ (C-H); 788 δ (CH_2); 742 δ_{s} (CF_3); 614 δ (SNS); Elemental analysis calcd (%) for $\text{C}_{16}\text{H}_{28}\text{F}_6\text{N}_2\text{O}_6\text{S}_2$: C, 36.78; H, 5.40; N, 5.36. Found (%): C, 37.10; H, 5.26; N, 5.37.

N,N-Diethyl-*N*-hexyl-*N*-[2-(methacryloyloxy)ethyl]ammonium bis(trifluoromethanesulfonyl)imide
([C₆N_{MA,22}]TFSI)

The respective intermediate [C₆N_{MA,22}]I (26.74 g, 65.0 mmol) was dissolved in acetone (40 mL) and added dropwise to a solution of LiTFSI (19.501 g, 67.9 mmol) in water (40 mL). The ionic liquid precipitated instantly and the mixture was stirred for 10 min at room temperature. Further processing was realized as described above for [C₂N_{MA,22}]TFSI to give the product as colorless oil (32.81 g, 91.7% yield). ¹H NMR (500 MHz, CDCl₃): δ (ppm) 0.85 (t, 3H, CH₃); 1.30 (m, 12H, CH₃, CH₂); 1.63 (m, 2H, CH₂); 1.90 (s, 3H, CH₃); 3.17 (m, 2H, CH₂); 3.36 (q, 4H, CH₂); 3.57 (m, 2H, CH₂); 4.47 (s, 2H, CH₂); 5.64 (s, 1H, CH₂); 6.06 (s, 1H, CH₂); ¹³C NMR (125 MHz, CDCl₃): δ (ppm) 7.3 (CH₃); 13.7 (CH₃); 18.0 (CH₃); 21.6 (CH₂); 22.3 (CH₂); 25.7 (CH₂); 30.9 (CH₂); 54.4 (CH₂); 55.8 (CH₂); 57.2 (CH₂); 58.6 (CH₂); 119.8 (q, CF₃); 127.2 (CH₂); 135.0 (C); 166.3 (C=O); IR (cm⁻¹): 2963 ν(C-H); 2932 ν(C-H); 2864 ν(C-H); 1724 ν(C=O); 1638 ν(C=C); 1459 δ_{as}(CH₂); 1399 δ_s(CH₃); 1348 ν(SO₂); 1181 ν(C-O); 1135 ν(S=O); 1055 ν(S=O); 949 δ(C-H); 788 δ(CH₂); 742 δ_s(CF₃); 654 δ(SNS); 616 δ_a(SO₂); Elemental analysis calcd (%) for C₁₈H₃₂F₆N₂O₆S₂: C, 39.27; H, 5.86; N, 5.09. Found (%): C, 39.48; H, 5.74; N, 5.07.

N,N-Diethyl-*N*-[2-(methacryloyloxy)ethyl]-*N*-octylammonium bis(trifluoromethanesulfonyl)imide
([C₈N_{MA,22}]TFSI)

The IL was prepared accordingly to the procedure described above for [C₆N_{MA,22}]TFSI starting from [C₈N_{MA,22}]I (27.65 g, 65.0 mmol) in acetone (40 mL) and LiTFSI (19.501 g, 67.9 mmol) in water (40 mL). The product was received as colorless oil (34.11 g, 90.7% yield). ¹H NMR (500 MHz, CDCl₃): δ (ppm) 0.87 (t, 3H, CH₃); 1.27 (m, 6H, CH₃); 1.33 (m, 10H, CH₂); 1.65 (s, 2H, CH₂); 1.93 (s, 3H, CH₃); 3.20 (m, 2H, CH₂); 3.38 (q, 4H, CH₂); 3.59 (m, 2H, CH₂); 4.50 (m, 2H, CH₂); 5.67 (s, 1H, CH₂); 6.09 (s, 1H, CH₂); ¹³C NMR (125 MHz, CDCl₃): δ (ppm) 7.4 (CH₃); 14.0 (CH₃); 18.0 (CH₃); 21.7 (CH₂); 22.5 (CH₂); 26.1 (CH₂); 28.9 (CH₂); 31.5 (CH₂); 54.5 (CH₂); 55.8 (CH₂); 57.2 (CH₂); 58.6 (CH₂); 119.8 (q, CF₃); 127.2 (CH₂); 135.0 (C); 166.4 (C=O); IR (cm⁻¹): 2959 ν(C-H); 2932 ν(C-H); 2857 ν(C-H); 1726 ν(C=O); 1638 ν(C=C); 1461 δ_{as}(CH₂); 1399 δ_s(CH₃); 1351 ν(SO₂); 1183 ν(C-O); 1135 ν(S=O); 1055 ν(S=O); 947 δ(C-H); 788 δ(CH₂); 742 δ_s(CF₃); 654 δ(SNS); 616 δ_a(SO₂); Elemental analysis calcd (%) for C₂₀H₃₆F₆N₂O₆S₂: C, 41.52; H, 6.27; N, 4.84. Found (%): C, 41.64; H, 6.05; N, 4.78.

N-[2-(Acryloyloxy)ethyl]-*N,N,N*-triethylammonium bis(trifluoromethanesulfonyl)imide
([C₂N_{A,22}]TFSI)

The IL was prepared accordingly to the procedure described above for [C₂N_{MA,22}]TFSI starting from [C₂N_{A,22}]I (24.451 g, 75.0 mmol) in water (15 mL) and LiTFSI (22.501 g, 68.38 mmol) in water (15 mL). The product was received as colorless oil (30.64 g, 85.0% yield). ¹H NMR (500 MHz, acetone-D₆): δ (ppm) 1.44 (t, 9H, CH₃); 3.62 (q, 6H, CH₂); 3.83 (m, 2H, CH₂); 4.67 (s, 2H, CH₂); 6.00 (dd, 1H, CH₂); 6.22 (m, 1H, CH); 6.42 (dd, 1H, CH₂); ¹³C NMR (125 MHz, acetone-D₆): δ (ppm) 6.9 (CH₃); 53.7 (CH₂); 55.1 (CH₂); 57.3 (CH₂); 120.1 (q, CF₃); 127.6 (CH); 131.7 (CH₂); 164.8 (C=O); IR (cm⁻¹): 2996 ν(C-H); 1730 ν(C=O); 1638 ν(C=C); 1461 δ_{as}(CH₂); 1410 δ_s(CH₃); 1349 ν(SO₂); 1177 ν(C-O); 1135 ν(S=O); 1053 ν(S=O); 996 δ(CH₂); 788 δ(CH₂); 742 δ_s(CF₃); 654 δ(SNS); 614 δ_a(SO₂); Elemental analysis calcd (%) for C₁₃H₂₂F₆N₂O₆S₂: C, 32.50; H, 4.62; N, 5.83. Found (%): C, 32.45; H, 4.40; N, 5.83.

N-[2-(Acryloyloxy)ethyl]-*N*-butyl-*N,N*-diethylammonium bis(trifluoromethanesulfonyl)imide
([C₄N_{A,22}]TFSI)

The IL was prepared accordingly to the procedure described above for [C₂N_{MA,22}]TFSI starting from [C₄N_{A,22}]I (26.645 g, 75.0 mmol) in water (20 mL) and LiTFSI (22.501 g, 68.38 mmol) in water (15 mL). The product was received as colorless oil (30.49 g, 80.5% yield). ¹H NMR (500 MHz, acetone-D₆): δ (ppm) 1.00 (s, 3H, CH₃); 1.43 (s, 8H, CH₃, CH₂); 1.83 (s, 2H, CH₂); 3.48 (s, 2H, CH₂); 3.62 (s, 4H, CH₂); 3.84 (s, 2H, CH₂); 4.67 (s, 2H, CH₂); 6.00 (m, 1H, CH₂); 6.21 (m, 1H, CH); 6.42 (m, 1H, CH₂); ¹³C NMR (125 MHz, acetone-D₆): δ (ppm) 7.0 (CH₃); 12.9 (CH₂); 19.4 (CH₂); 23.4 (CH₂);

54.2 (CH₂); 55.7 (CH₂); 57.3 (CH₂); 58.1 (CH₂); 120.1 (q, CF₃); 127.6 (CH); 131.7 (CH₂); 164.8 (C=O); IR (cm⁻¹): 2961 ν(C-H); 2935 ν(C-H); 2866 ν(C-H); 1730 ν(C=O); 1638 ν(C=C); 1463 δ_{as}(CH₂); 1408 δ_s(CH₃); 1351 ν(SO₂); 1179 ν(C-O); 1135 ν(S=O); 1055 ν(S=O); 790 δ(CH₂); 742 δ_s(CF₃); 654 δ(SNS); 616 δ_a(SO₂); Elemental analysis calcd (%) for C₁₅H₂₆F₆N₂O₆S₂: C, 35.43; H, 5.15; N, 5.51. Found (%): C, 35.45; H, 5.15; N, 5.57.

N-[2-(Acryloyloxy)ethyl]-*N,N*-diethyl-*N*-hexylammonium bis(trifluoromethanesulfonyl)imide
([C₆N_{A,22}]TFSI)

The IL was prepared accordingly to the procedure described above for [C₆N_{MA,22}]TFSI starting from [C₆N_{A,22}]I (24.915 g, 65.0 mmol) in acetone (40 mL) and LiTFSI (19.501 g, 67.9 mmol) in water (30 mL). The product was received as colorless oil (31.20 g, 89.5% yield). ¹H NMR (500 MHz, CDCl₃): δ (ppm) 0.89 (s, 3H, CH₃); 1.33 (s, 12H, CH₃, CH₂); 1.66 (s, 2H, CH₂); 3.21 (m, 2H, CH₂); 3.38 (m, 4H, CH₂); 3.60 (s, 2H, CH₂); 4.52 (s, 2H, CH₂); 5.96 (m, 1H, CH₂); 6.11 (m, 1H, CH); 6.44 (m, 1H, CH₂); ¹³C NMR (125 MHz, CDCl₃): δ (ppm) 7.4 (CH₃); 13.7 (CH₂); 21.7 (CH₂); 22.3 (CH₂); 25.8 (CH₂); 31.0 (CH₂); 54.5 (CH₂); 55.8 (CH₂); 56.9 (CH₂); 58.7 (CH₂); 119.8 (q, CF₃); 126.9 (CH); 132.8 (CH₂); 165.1 (C=O); IR (cm⁻¹): 2972 ν(C-H); 2883 ν(C-H); 1732 ν(C=O); 1637 ν(C=C); 1463 δ_{as}(CH₂); 1408 δ_s(CH₃); 1349 ν(SO₂); 1179 ν(C-O); 1135 ν(S=O); 1053 ν(S=O); 790 δ(CH₂); 740 δ_s(CF₃); 653 δ(SNS); 614 δ_a(SO₂); Elemental analysis calcd (%) for C₁₇H₃₀F₆N₂O₆S₂: C, 38.06; H, 5.64; N, 5.22. Found (%): C, 37.70; H, 5.63; N, 5.13.

N-[2-(Acryloyloxy)ethyl]-*N,N*-diethyl-*N*-octylammonium bis(trifluoromethanesulfonyl)imide
([C₈N_{A,22}]TFSI)

The IL was prepared accordingly to the procedure described above for [C₆N_{MA,22}]TFSI starting from [C₈N_{A,22}]I (26.739 g, 65.0 mmol) in acetone (40 mL) and LiTFSI (19.501 g, 67.9 mmol) in water (30 mL). The product was received as colorless oil (32.77 g, 89.3% yield). ¹H NMR (500 MHz, CDCl₃): δ (ppm) 0.88 (t, 3H, CH₃); 1.28 (m, 6H, CH₂); 1.34 (t, 10H, CH₃, CH₂); 1.66 (s, 2H, CH₂); 3.20 (m, 2H, CH₂); 3.39 (q, 4H, CH₂); 3.60 (m, 2H, CH₂); 4.52 (m, 2H, CH₂); 5.96 (dd, 1H, CH₂); 6.12 (m, 1H, CH); 6.45 (dd, 1H, CH₂); ¹³C NMR (125 MHz, CDCl₃): δ (ppm) 7.4 (CH₃); 14.0 (CH₂); 21.8 (CH₂); 22.5 (CH₂); 26.1 (CH₂); 28.9 (CH₂); 31.5 (CH₂); 54.5 (CH₂); 55.8 (CH₂); 56.9 (CH₂); 58.7 (CH₂); 119.8 (q, CF₃); 126.9 (CH); 132.9 (CH₂); 165.1 (C=O); IR (cm⁻¹): 2959 ν(C-H); 2932 ν(C-H); 2860 ν(C-H); 1732 ν(C=O); 1638 ν(C=C); 1465 δ_{as}(CH₂); 1408 δ_s(CH₃); 1349 ν(SO₂); 1179 ν(C-O); 1135 ν(S=O); 1055 ν(S=O); 790 δ(CH₂); 742 δ_s(CF₃); 656 δ(SNS); 616 δ_a(SO₂); Elemental analysis calcd (%) for C₁₉H₃₄F₆N₂O₆S₂: C, 40.42; H, 6.07; N, 4.96. Found (%): C, 41.04; H, 6.01; N, 5.06.

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