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Controlled radical polymerization and in-depth mass-spectrometric characterization of poly(ionic liquid)s and their photopatterning on surfaces†

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The preparation and characterization of poly(ionic liquid)s (PILs) bearing a polystyrene backbone via reversible addition fragmentation chain transfer (RAFT) polymerization and their photolithographic patterning on silicon wafers is reported. The controlled radical polymerization of the styrenic ionic liquid (IL) monomers ([BVBIM]X, X = Cl⁻ or Tf₂N⁻) by RAFT polymerization is investigated in detail. We provide a general synthetic tool to access this class of PILs with controlled molecular weight and relatively narrow molecular weight distribution (2000 g mol⁻¹ ≤ M_n ≤ 10 000 g mol⁻¹ with dispersities between 1.4 and 1.3 for p([BVBIM]Cl); 2100 g mol⁻¹ ≤ M_p ≤ 14 000 g mol⁻¹ for p([BVBIM]Tf₂N)). More importantly, we provide an in-depth characterization of the PILs and demonstrate a detailed mass spectrometric analysis via matrix-assisted laser desorption ionization (MALDI) as well as – for the first time for PILs – electrospray ionization mass spectrometry (ESI-MS). Importantly, p([BVBIM]Cl) and p([DMVBIM]Tf₂N) were photochemically patterned on silicon wafers. Therefore, a RAFT agent carrying a photoactive group based on *ortho*-quinodimethane chemistry – more precisely photoenol chemistry – was photochemically linked for subsequent controlled radical polymerization of [BVBIM]Cl and [DMVBIM]Tf₂N. The successful spatially-resolved photografting is evidenced by surface-sensitive characterization methods such as X-ray photoelectron spectroscopy (XPS) and time-of-flight secondary ion mass spectrometry (ToF-SIMS). The presented method allows for the functionalization of diverse surfaces with poly(ionic liquid)s.

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Introduction

Ionic liquids (ILs) are organic salts with a melting point typically below 100 °C. They have found significant applications in various fields ranging from chemistry to physics and materials science.¹ Recently, the polymerization of ILs, which forms a new class of ionic polymers, so called poly(ionic liquid)s (PILs), has emerged as a multifunctional platform that combines the advantages of ILs with the general property profile of polymers.^{2–6} PILs are not only a natural extension of properties and function of ILs in materials science, but significantly advance the research scope of ionic polymers. For example, in spite of a high ion-pair packing density, the favorable solubility in a broad range of organic solvents and the achievable low glass transition temperature of structurally well-defined PILs offer new processing possibilities to fabricate ionic polymers for numerous functional materials.

Currently, the chemical structure of polymerizable ILs is dominated by the imidazolium type, which can be further classified into three major subgroups, *i.e.* the vinylimidazolium, (meth)acrylate, and styrenic ones. In the past several

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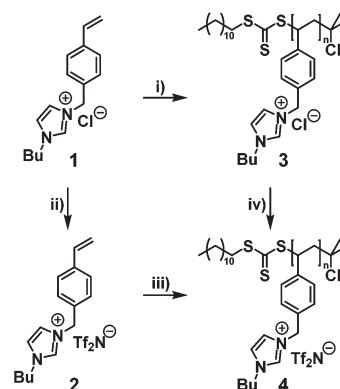
years, a wide range of polymerization techniques have been exploited to synthesize imidazolium-type PILs with defined structure characteristics. Among them, reversible deactivation radical polymerization techniques, such as the atom transfer radical polymerization (ATRP), and reversible addition–fragmentation chain-transfer (RAFT) polymerization are frequently used.^{7–12} These radical polymerization techniques allow for superior control over the polymer chain characteristics and the endgroup functionalization.

PILs bearing styrenic type IL monomer units have a rigid polystyrene backbone that can provide mechanical stability to lock the highly ionic polymer matrix. These PILs are promising candidates as support for IL networks. For example, covalent porous IL networks are often generated from styrenic type IL monomers.¹³ Additionally, PIL membrane materials for gas separation or sorption take advantage of the rigid polystyrene backbone to enhance the shape durability of membranes and the gas flux rate.¹⁴ Preparation of such PILs from styrenic IL monomers is a synthetic task that usually involves free radical polymerization, which is easy to implement and important for large scale production. Yet, to understand the intrinsic structure–property–function relationship for further optimization of their materials performance, PILs with defined chain length and narrow molecular weight distribution are absolutely necessary. In literature, ATRP of a styrenic IL monomer has been reported by the groups of Matyjaszewski and Gin.^{8,15} Both groups demonstrated that *via* ATRP, the PIL chain characteristics are tuneable and their block copolymers are accessible. An inherent problem in the polymerization of IL monomers by ATRP is, however, the inevitable ion exchange process between the required metal catalyst, copper halide and the IL monomer. Further, Cu ions may form complex anions with certain IL anions,¹⁶ which makes it even more challenging to be removed from the polymer product. Therefore, in spite of excellent control over the polymerization process *via* ATRP, the field is motivated to search for alternative polymerization techniques to overcome these challenges. One solution is to apply RAFT polymerization that generally involves no metal salt. A study by Tenhu and coworkers¹⁷ reported briefly that it is possible to polymerize a specific styrenic type chiral IL monomer, yet the RAFT polymerization of more representative and commonly used styrenic IL monomers, such as 1-butyl-3-(*p*-vinylbenzyl)-imidazolium-based IL monomers was not presented. Recently, Detrembleur and coworkers¹⁸ demonstrated the preparation of vinylimidazolium-based block PILs featuring two different counter ions *via* RAFT polymerization.

The precise characterization of PIL chain features, in general, is very challenging. Due to the strong interactions with the column materials, size exclusion chromatography results in broadened traces, or in a complete absorption on the column surface. Different teams pioneered measuring PILs with different counter ions by SEC.^{19,20} However, to date, only little efforts towards the detailed mass spectrometric analysis of these materials have successfully been carried out.^{8,9,21} Motivated by the current status, we initiated the project on

RAFT polymerization of frequently used styrenic IL monomers and provide a general synthetic protocol to access these PILs with controlled molecular weight, narrow molecular weight distribution, defined chain endgroup and no contamination from metal ion species. More precisely, we investigate the controlled radical polymerization of 1-butyl-3-(*p*-vinylbenzyl)-1*H*-imidazolium chloride ([BVBIM]Cl, (1)) and 1-butyl-3-(*p*-vinylbenzyl)-1*H*-imidazol-3-ium bis(trifluoromethanesulfonyl)imide ([BVBIM]Tf₂N, (2)) (Scheme 1) *via* RAFT polymerization and provide a comprehensive characterization of the polymers by various (mass spectrometric) analysis methods such as SEC, MALDI-ToF MS as well as ESI-MS. To the best of our knowledge, the mass spectrometric characterization *via* ESI-MS is shown for the first time in-depth.

In addition, surface design is a rapidly growing field in contemporary chemistry. PIL-decorated surfaces can serve as imaging, wetting/dewetting, and anti-biofouling interfaces.^{8,22,23} Initiated by the pioneering work of Ye *et al.* on grafted PILs *via* ROMP on TiO₂ substrates and the positive anti-fouling studies,²³ Matyjaszewski and coworkers thermally attached well-characterized, ATRP-prepared PILs on Si substrates that potentially could show anti-fouling characteristics.⁸ Furthermore, Drockenmueller and coworkers patterned azide-functionalized copolymerized PILs *via* UV irradiation and subsequent formation of highly reactive nitrenes on a silicon substrate.²⁴ Herein, we present a polymerization study of p([BVBIM]Cl) and p([BVBIM]Tf₂N) in order to investigate the mass spectrometric behavior of PILs in solution. Furthermore, we study the characterization of photochemically patterned p([BVBIM]Cl) and p[1,2-dimethyl-3-(*p*-vinylbenzyl)-1*H*-imidazol-3-ium bis(trifluoromethanesulfonyl)imide] (p([DMVIM]-Tf₂N)) *via* a surface sensitive mass spectrometric analytical method, *i.e.* time-of-flight secondary ion mass spectrometry (ToF-SIMS) as well as X-ray photoelectron spectroscopy (XPS).



Scheme 1 Synthetic approach for the preparation of p([BVBIM]Cl) (3) and p([BVBIM]Tf₂N) (4) from the monomers [BVBIM]Cl (1) and [BVBIM]Tf₂N (2), respectively. Molar ratio [M]:[CTA]:[I] for (i) 200:1:0.2 and for (iii) 200:1:0.2 (i: RAFT polymerization; ii: Tf₂N[−] vs. Cl[−] anion exchange; iii: RAFT polymerization; iv: anion exchange).



Results and discussion

Among other versatile features of the RAFT process,²⁵ the high tolerance towards functional monomers *e.g.* acrylic acids or vinyl esters, is an important characteristic. It is also expected to be a suitable candidate for the controlled radical polymerization of monomeric ILs and thus applied in the presented work. Scheme 1 summarizes the synthetic strategy for the preparation of the polymers p([BVBIM]Cl) (3) and p([BVBIM]Tf₂N) (4). It should be noted that the synthesis and properties of 1-alkyl-3-(*p*-vinylbenzyl)-imidazolium based PILs was widely investigated,^{8,15,26} yet the RAFT process of the corresponding monomers remains a challenge.

The polymerization of these ILs can either be achieved by the direct polymerization of [BVBIM]Tf₂N (2) or by the polymerization of [BVBIM]Cl (1) followed by a counter ion exchange with Tf₂N⁻. Both monomers [BVBIM]Cl (1) and [BVBIM]Tf₂N (2) were synthesized by a quaternization reaction between 4-vinylbenzyl chloride and *n*-butylimidazole with a slightly modified synthesis protocol given in the literature.²⁷ A counter ion exchange with Tf₂N⁻ resulted in 2. The polymerization of [BVBIM]Cl (1) was performed in DMSO as solvent of choice ensuring a good dissolution of both the IL monomer (1) and 2-cyano-2-propyl dodecyl trithiocarbonate (CPDT) as chain transfer agent (CTA). A molar ratio of the reagents [M]:[CTA]:[I] = 200:1:0.2 (Initiator: 1,1'-azobis(cyclohexanecarbonitrile)) was used. Interestingly, the RAFT polymerization of [BVBIM]Cl (1) is highly affected by the choice of the solvent. For instance, protic solvents, such as ethanol, methanol and water, led to a less controlled polymerization (refer to Table S1 in the ESI†). The polymerization of [BVBIM]Tf₂N (2) was performed in butyronitrile as solvent due to its favorable solubility in non-polar/aprotic solvents. In general, the common controlled polymerization techniques, *i.e.* ATRP, RAFT and NMP, are complex and challenging for IL monomers such as 1 and 2 because of their ionic character.²⁰ The RAFT polymerization of [BVBIM]Cl (1) was carried out employing CPDT as chain transfer agent in DMSO at 85 °C and VAZO-88 ($k_d = 2.01 \times 10^5 \text{ s}^{-1}$)²⁸ as radical starter. The molecular weight evolution of the SEC traces in Fig. 1(A) verifies the living character. Both the ¹H NMR (Fig. S11†) as well as the SEC results are in good agreement with the theoretically obtained molecular weights (Fig. 1(B): solid line). P([BVBIM]Cl) (3) is a charged polymer and thus interacts strongly with the column material and its polymer chains *via* hydrogen bonding of the chloride counter ion. As a consequence, halide-containing PILs prepared *via* RAFT polymerization often display a rather broad dispersity in a H₂O-SEC system.⁹

Thus, the optimum system for the SEC analysis has to be identified to suppress the intermolecular polymer and column interactions. In order to investigate the most appropriate eluent, the SEC measurements were carried out by variation of the polymer concentrations (in mg mL⁻¹) as well as sodium chloride salt concentrations (refer to Fig. S1†). The H₂O-SEC system with a water/0.3 M formic acid/0.5 g L⁻¹ NaCl mixture as eluent was identified as ideal system for the polymeric

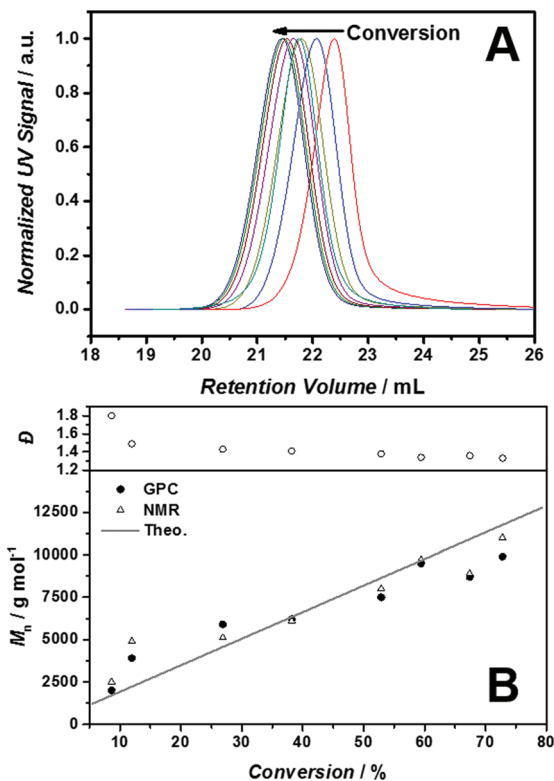


Fig. 1 (A) SEC traces of p([BVBIM]Cl) (3) *via* the RAFT mediated polymerization of [BVBIM]Cl (1) at 85 °C in DMSO. (B) Plot of molecular weight (M_n) and dispersity (D) vs. monomer conversion. Molar ratio of monomer: CTA: initiator 200:1:0.2 (CTA = 2-cyano-2-propyl dodecyl trithiocarbonate; initiator = 1,1'-azobis(cyclohexanecarbonitrile)). H₂O-SEC system conditions: water/0.3 M formic acid/0.5 g L⁻¹ NaCl, 30 °C, flow rate 1.0 mL min⁻¹.

system (3). The system was calibrated against poly(4-vinylpyridine), which forms a polycation under SEC conditions and thus allows for a more accurate estimation of the molecular weight of our PILs. Furthermore, we have demonstrated that the dispersity decreases after successful salt metathesis of the corresponding p([BVBIM]Cl) (3) ($M_n = 8900 \text{ g mol}^{-1}$; $D = 1.55$; $DP_n = 37$) against LiTf₂N resulting in p([BVBIM]Tf₂N) (4) with a dispersity of 1.2 (SEC analysis with an eluent containing 10 mM LiTf₂N and 10 mM *n*-butylimidazole). The calibration against polystyrene was in good agreement to the results obtained with the H₂O-SEC system (refer to Fig. S1†).

The polymerization of the Tf₂N⁻ containing monomeric IL 2 can be achieved in a similar synthetic approach. CPDT was used as chain transfer agent in butyronitrile at 85 °C and 1,1'-azobiscyclohexane-carbonitrile as initiator.

Fig. 2(A) shows the molecular weight evolution for the RAFT polymerization of [BVBIM]Tf₂N (4) (10 mmol LiTf₂N, 10 mmol *n*-butylimidazole, 35 °C, flow rate 1.0 mL min⁻¹). The controlled molecular weight evolution is depicted in Fig. 2(B). The determination of molecular weights of the polymers ($M_n < 10000 \text{ g mol}^{-1}$) is hampered by the presence of residual



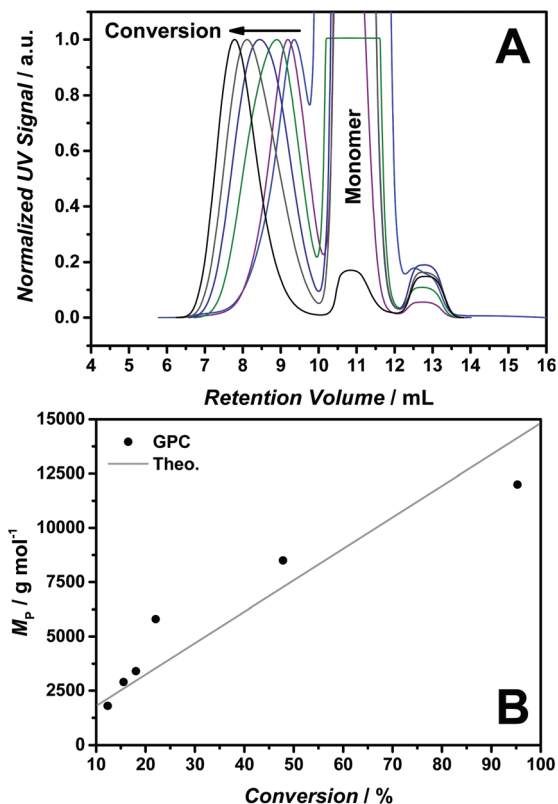


Fig. 2 (A) Molecular weight evolution of the RAFT polymerization of [BVBIM]Tf₂N (2) at 85 °C in butyronitrile. (B) Molecular weight evolution (M_p) with conversion for the RAFT mediated polymerization of 2. Molar ratio [M] : [CTA] : [I] = 200 : 1 : 0.2, 85 °C, butyronitrile (CTA: 2-cyano-2-propyl dodecyl trithiocarbonate; I = 1,1'-azobiscyclohexane carbonitrile). THF-SEC system conditions: 10 mmol LiTf₂N, 10 mmol *n*-butylimidazole, 35 °C, flow rate 1.0 mL min⁻¹.

monomer. Precipitation in methanol/water was not successful, also dialysis in water (3 d at ambient temperature as well as for 14 d at elevated temperatures of close to 40 °C) did not result

in pure PILs. The dialysis in THF or acetone according to a literature method⁸ may work for high-molecular weight PILs, however failed for our low-molecular weight polymers (due to diffusion out of the membrane). The dialysis in DMSO was successful for certain polymer samples, however, we were not able to implement the DMSO purification as a standard dialysis for Tf₂N⁻ containing PILs. Thus, the polymer signals and the monomer signal overlap in the SEC trace. The overlap is most likely due to the SEC column system in our laboratory that differs slightly from the SEC column system employed by Matyjaszewski and coworkers.^{8,20}

Thus, the peak of the molecular weight (M_p) was correlated with the conversion indicating a linear trend. However, the dispersity (D) (due to the overlap of the polymer signal and the monomer signal) was not determined and also the M_n determination *via* NMR spectroscopy was hampered by the presence of the residual monomer.

Table 1 provides an overview of the molecular weights calculated according to the monomer conversion, as well as the molecular weights obtained by NMR spectroscopy and SEC for 3 and 4. Salt exchange against bis(trifluoromethanesulfonyl)-imide (Tf₂N⁻) converts the water-soluble polymers into highly hydrophobic ones that are soluble in solvents such as THF, DMAc and acetone.²⁹ Matyjaszewski and co-workers have shown that such PILs can easily be characterized by SEC with THF as eluent containing a small concentration of the appropriate counter ion.²⁰

In contrast to SEC, mass spectrometry is often employed in order to characterize the polymer chain structure. Nevertheless, mass spectrometric analysis of PILs is a considerable challenge because of their ionic character.⁴ In particular when endgroup analysis is required, the determination of the exact mass is a crucial step for a detailed characterization of the synthesized materials, *i.e.* for subsequent post-modification reactions.³⁰ Electrospray ionization mass spectrometry (ESI-MS)³¹ and matrix assisted laser desorption ionization (MALDI)³² are the methods of choice for an in-depth characterization of the

Table 1 Characterization of p([BVBIM]Cl) (3) and p([BVBIM]Tf₂N) (4) prepared by RAFT polymerization in different solvents. Conditions for 3: 85 °C, CPDT, VAZO-88, [M] : [CTA] : [I] = 200 : 1 : 0.2, DMSO. Conditions for 4: 85 °C, CPDT, VAZO-88, [M] : [CTA] : [I] = 200 : 1 : 0.2, butyronitrile

Sample	Time [h]	$M_n^{\text{theo.}}$ [g·mol ⁻¹]	M_n^{NMR} [g·mol ⁻¹]	$M_n^{\text{SEC}*}$ [g·mol ⁻¹]	D^{SEC}	Conv. ^{NMR} [%]
3-1	1	1700	2500	2000 ^a	1.80 ^a	9
3-2	2	2300	4900	3900 ^a	1.49 ^a	12
3-3	3	4500	5100	5900 ^a	1.43 ^a	27
3-4	4	6400	6100	6200 ^a	1.41 ^a	38
3-5	5	8600	8000	7500 ^a	1.38 ^a	53
3-6	6	9600	8900	8700 ^a	1.36 ^a	59
4-1	3	2100	n.d.	1800 ^b	n.d.	12
4-2	4	2600	n.d.	2900 ^b	n.d.	16
4-3	6	3000	n.d.	3400 ^b	n.d.	18
4-4	7	3500	n.d.	5800 ^b	n.d.	22
4-5	8	7300	n.d.	8500 ^b	n.d.	48
4-6	24	14 000	n.d.	12 000 ^b	n.d.	95

*For samples 3: M_n^{SEC} [g·mol⁻¹]; for samples 4: M_p^{SEC} [g·mol⁻¹]. ^a H₂O-SEC, eluent: water/0.3 M formic acid/0.5 g L⁻¹ NaCl; temperature: 30 °C; flow rate: 1.0 mL min⁻¹. ^b THF-SEC, eluent: THF/10 mM LiTf₂N/10 mM *n*-butylimidazole; temperature: 35 °C; flow rate: 1.0 mL min⁻¹.



mixture, it is necessary to analyze the polymers with MALDI-ToF MS. The value of the repeating unit ($m/z = 240.3$ Th) between two $[3_{a,b}]^+$ peaks is in good agreement with the theoretical value ($m/z_{(\text{theo.})} = 240.2$ Th). As already discussed, the chloride degrades the alkyl chain ($[3_{b,d}]^+$ and $[3_{a,d}]^+$), however, these signals are of minor intensity (Fig. 4(A)).

Also, a RAFT endgroup conversion was studied dispersing the polymer in THF for three days. After three days, THF was evaporated, the polymer dissolved in methanol and mixed with the matrix in a ratio of 10 : 1 (matrix: sample). Clearly, THF reacted with the RAFT moiety³⁸ as indicated by the MALDI-ToF MS measurements with the identified peaks $[3 \text{ OH}]$ and $[3 \text{ SO}_2]$ (refer to the ESI Fig. S4† and Fig. 5).

Most evident are two adjacent peaks with a m/z difference of 64 Th. According to Benicewicz and coworkers,³⁷ trithiocarbonates are prone to be oxidized to the corresponding sulfonic acid in THF after storage for 48 h, and to release gaseous SO_2 (64 Th) to form the hydroxyl endgroup (as matrix the team employed dithranol doped with 0.01 M AgTFA). As observed for methanol as solvent, gaseous HCl is released during the ionization process, resulting in an uncharged polymer.

Fig. 4(B) and 6(B) illustrate the MALDI-ToF as well as the ESI-MS spectrum obtained for $p([\text{BVBIM}]\text{Tf}_2\text{N})$ **4**. In contrast to the counter ion Cl^- of (**3**) that shows a high H-bond acceptor affinity, the weakly coordinating counter ion Tf_2N^- of **4** does not interact with the protons of the imidazolium ring. The repeating unit between two $[4]^+$ peaks ($m/z = 521.1$ Th) agrees well with the theoretical value ($m/z_{(\text{theo.})} = 521.1$ Th).

As observed in the MALDI-ToF measurements as well, the quality of the ESI-MS measurement correlates with the molecular weight of the polymer. The repeating unit of $p([\text{BVBIM}]\text{Cl})$ (**3**) is present in four species (please refer to Fig. 3). The permutation of these processes together with the increasing chain length, multiple charges, and whether a proton or a sodium ion ionizes the polymer, results in high amounts of detectable ions. However, the degradation of the alkyl side chain (Fig. 3c and d) is observed with a limited probability. Thus, it is not surprising that – especially for high-molecular weight polymers – the spectrum is complex. The assignment of the dominant peaks is summarized in Table S4 in the ESI.†

Furthermore, the Coulomb interactions of Tf_2N^- appear to be stronger compared to Cl^- and thus ESI-MS measurements

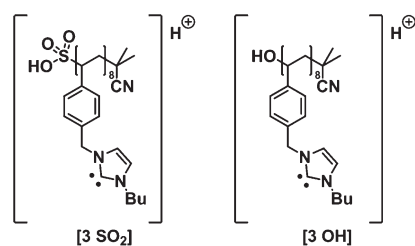


Fig. 5 Chemical structures assigned to the MALDI spectrum of $p([\text{BVBIM}]\text{Cl})$ (**3**) ($M_w = 1900 \text{ g mol}^{-1}$). PIL structure with a sulfonic acid endgroup and a hydroxyl endgroup.³⁷ The m/z difference of both peaks originates from the loss of SO_2 .

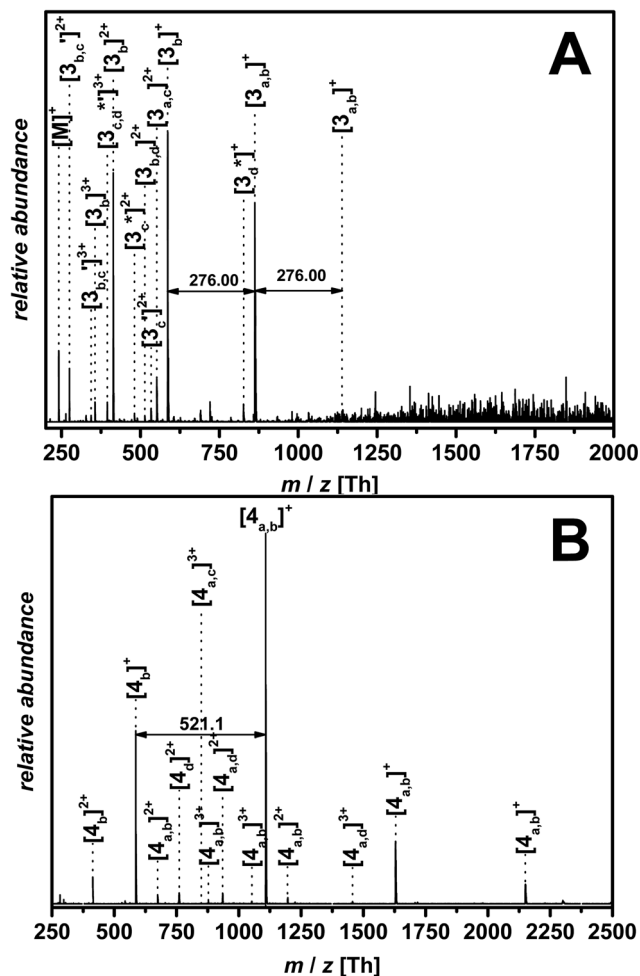


Fig. 6 Zoom of ESI-MS spectra of (A) $p([\text{BVBIM}]\text{Cl})$ (**3**) with a molecular weight of $M_w = 1900 \text{ g mol}^{-1}$ measured in THF/MeOH (3 : 2, v/v) and (B) $p([\text{BVBIM}]\text{Tf}_2\text{N})$ (**4**) measured in a mixture of acetonitrile/water (1 : 1, v/v). Please refer to the ESI† for a list of the assignments (Tables S4 and S5†).

lead to clear spectra (Fig. 6(B)). The signal difference of $m/z = 521.1$ Th is assigned to the molecular weight of the monomer unit ($m/z_{(\text{theo.})} = 521.1$ Th). The thiocarbonyl thio endgroup of **4** is clearly identified (see Table S5†). The ESI-MS analysis of the PIL bearing Tf_2N^- as counter ion is easier to evaluate since there are far less side events affecting the polymer chain during the ionization process compared to $p([\text{BVBIM}]\text{Cl})$ (**3**) polymers.

To demonstrate the versatility of the employed approach, $[\text{BVBIM}]\text{Cl}$ (**1**) and another monomer, *i.e.* 1,2-dimethyl-3-(*p*-vinylbenzyl)-1*H*-imidazol-3-ium bis(trifluoromethane-sulfonyl)-imide ($[\text{DMVBIM}]\text{Tf}_2\text{N}$ (**5**), were applied for spatially resolved light-induced surface modification. Fig. 7 details the grafting-from approach of the RAFT polymerization of **1** and **5**. The photoenol RAFT agent **7'** was photopatterned *via* photoenol chemistry onto maleimide-functionalized silicon wafers **6**. Subsequently, **1** and **5** were polymerized on the surfaces **7** by a grafting-from approach leading to highly functionalized patterned structures (**8a**, **8b**).



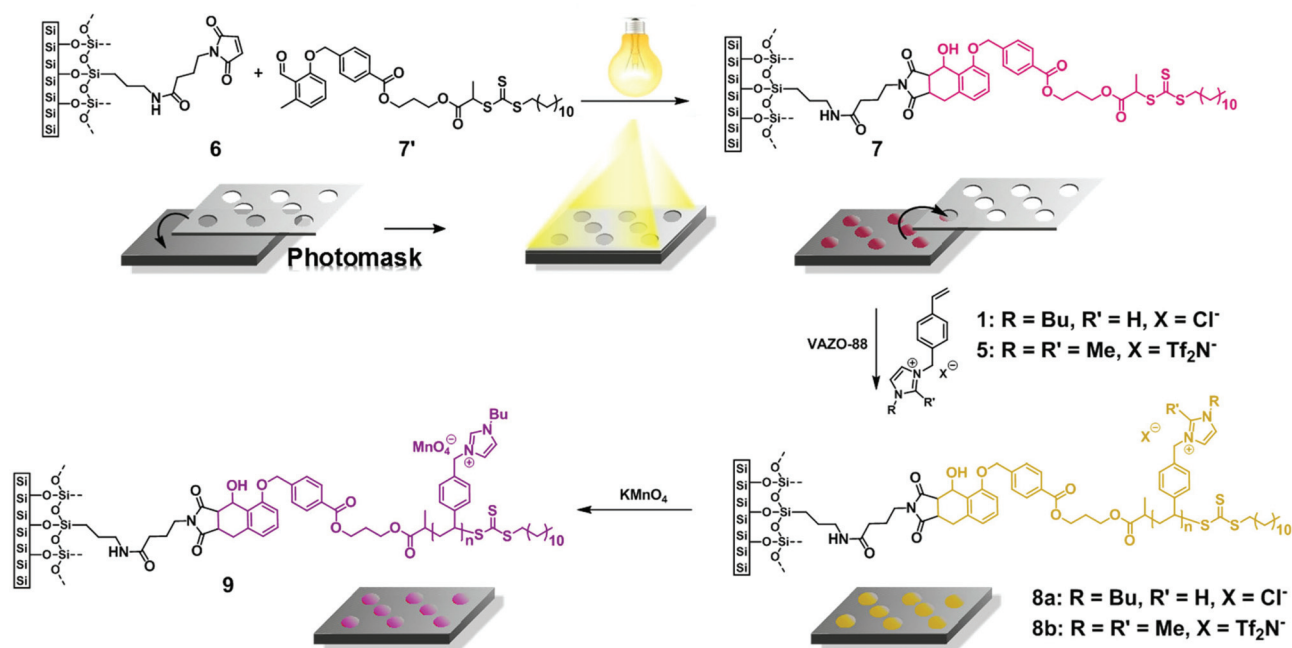


Fig. 7 Schematic illustration of the surface functionalization for the spatially resolved surface functionalization of silicon wafers with (1) and (5). (3-Aminopropyl)triethoxysilane (APTES)-coated silicon wafers were coated with 4-maleimidobutyryl chloride to achieve maleimide-functionalized surfaces (6). After irradiation, the surfaces covered with a photomask with an ARIMED B6 lamp ($\lambda_{\text{max}} = 320$ nm) in a solution of the photoenol functional CTA to result in (7) followed by RAFT polymerization of (1) from the surface. Subsequent salt metathesis of (8a) with KMnO_4 resulted in a highly violet colored meander pattern of $\text{p}([\text{BVBIM}]\text{MnO}_4)$ (9).

First, APTES-coated surfaces,³⁹ thoroughly cleaned by rinsing them with toluene, acetone, methanol and Milli Q water, were placed in a solution of freshly prepared 4-maleimidobutyryl chloride in dichloromethane to obtain maleimide-coated surface **6** acting as dienophile in the subsequent photoenol reaction with the CTA (**7'**) in acetonitrile (due to its low absorption in the UV/Vis area) for 2.5 hours.

Therefore, a photomask is placed on the maleimide-coated surfaces **6**, and irradiated with an ARIMED B6 lamp ($\lambda_{\text{max}} = 320$ nm) for spatially resolved photopatterning of the samples with a RAFT agent for subsequent RAFT polymerization of **1** and **5**. The surface was rinsed thoroughly with THF, acetone, methanol and Milli Q water, dried under a flow of nitrogen, and analyzed *via* XPS and ToF-SIMS, respectively.

The polymerization of $[\text{BVBIM}]\text{Cl}$ (**1**) was subsequently carried out in a DMSO/DMF mixture (1:1, v/v) with a monomer concentration of approx. 0.6 mol L^{-1} . After addition of 2-(dodecylthiocarbonothioyl-thio)propionic acid (DoPAT) as sacrificial RAFT agent and VAZO-88 as radical starter, the mixture was degassed *via* three consecutive freeze-pump-thaw cycles. As demonstrated by Zamfir *et al.*,⁴⁰ the grafting-from approach is best carried out in a 1:1 ratio of CTA: initiator. The high initiator concentration used is necessitated by the hopping migration mechanism explaining the steady termination of adjacent radicals on the surface. The polymerization was allowed to proceed for 7 h (50% conversion), ensuring the formation of high-molecular weight polymer on the surface.

The residual monomer as well as potential polymer attached *via* physisorption was removed by thorough rinsing of the functionalized surface **8a** with Milli Q water, methanol, acetone, and Milli Q water followed by immersing the surface in methanol for 12 h. Subsequently, the surface was ultrasonically treated in a solution containing 0.5 g L^{-1} ammonium chloride and rinsed thoroughly with acetone.

Since $\text{p}([\text{DMVBIM}]\text{Tf}_2\text{N})$ (55% conversion after 7 h) is highly hydrophobic, the residual monomer as well as potential polymer attached *via* physisorption was removed by thorough rinsing of the functionalized surface **8b** with acetone, dichloromethane, methanol and Milli Q water followed by treating the surface in a solution containing 1.0 g L^{-1} LiTf_2N . Residual salt was removed *via* ultrasonification in acetone.

Fig. 8 depicts the corresponding XP spectra. The C 1s spectrum shows the (C-C, C-H) component at 285.0 eV, (C-O, C-N) at 286.4 eV and (COO, NCO) at 288.5 eV, which are present in the maleimide (surface **6**) and the photoenol (surface **7**).⁴¹ The photo-ligation of **7** was evidenced by the sulfur signal at 163.6 eV indicating the presence of the trithiocarbonate group (refer to Fig. S13 in the ESI[†]).⁴²

After the surface-initiated polymerization to obtain **8a**, the formation of the signal at 401.8 eV, which is characteristic for the imidazole group (N^+), is detected in the N 1s spectrum. The (protonated) amine signal at 401.8 eV in the XPS increases from 0.3 at% (surface **7**) obviously stemming from a contamination (*e.g.* unreacted APTES) to 8.9 at% (surface **8a**), which



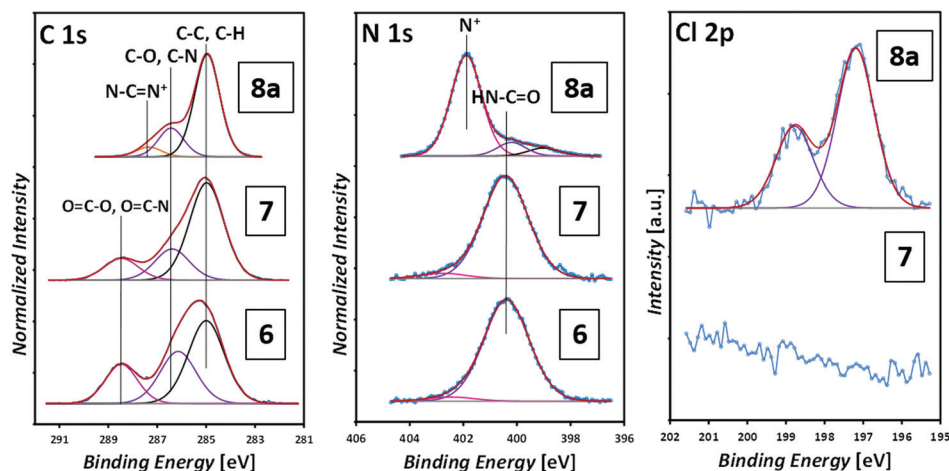


Fig. 8 XPS spectra of the maleimide functionalization (6), photoenol functionalization (7) and the grafting-from polymerization (8a) with [BVBIM]Cl (1).

can be assigned to the formation of the imidazole group. Most characteristically, the presence of a signal originating from Cl 2p indicates the counter ion, which is absent on the APTES/maleimide-coated surface 6 as well as on the RAFT agent-functionalized surface 7. ToF-SIMS images of the $C_{12}H_{25}S^-$, S^- , $C_{16}H_{21}N_2^+$ (surface 8a) and $C_{14}H_{17}N_2^+$ (surface 8b) fragments are shown in Fig. 9. ToF-SIMS images clearly confirm the covalent attachment of 7' precisely following the dotted shape of the photomask. As expected, in the areas covered by the photomask, no photoenol reaction proceeded and thus no characteristic signals stemming from 7 were observed. After successful grafting-from of [BVBIM]Cl (1), ToF-SIMS measurements in the positive mode confirm the presence of the repeating unit ($C_{16}H_{21}N_2^+$; $m/z = 241.20$ Th) on the surface 8a. Furthermore, successful grafting of [DMVBIM]Tf₂N (5) was also confirmed by its characteristic repeating unit ($C_{14}H_{17}N_2^+$; $m/z = 213.13$ Th) on the surface 8a. Since this surface was prepared as proof-of-principle for the ToF-SIMS measurement, no XPS measurements were carried out on surface 8b. In addition, in order to confirm the presence of the ionic nature of the PIL attached to the surface, the substrate was immersed in $KMnO_4$ (refer to Fig. 7). After dipping the substrate into a 50 mM $KMnO_4$ solution for only two to three seconds, the surface was thoroughly rinsed with methanol and Milli Q water. Clearly, the meander structure – assigned to the successful photopatterning – became visible immediately. The stability of PILs with a redox-active counter ion was shown by Mecerreyes and coworkers.⁴³

Thus, it is believed that the ion exchange of Cl^- against MnO_4^- proceeded on the surface 9.⁴⁴ It has to be noted that the brownish color of the pattern can be assigned to MnO_2 that was formed due to the rapid degradation of MnO_4^- (Fig. 9). The meander structure was used for this experiment to show the versatility of the presented photochemically induced approach.

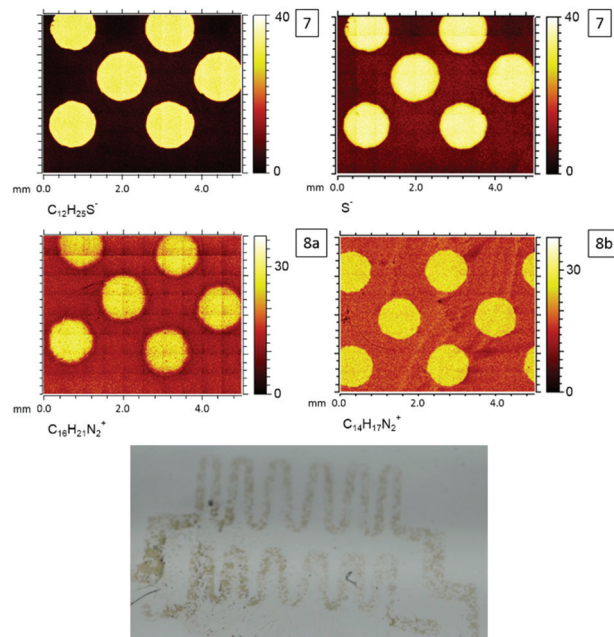


Fig. 9 ToF-SIMS images of photoenol functionalization (7), grafting-from polymerization (8a) with [BVBIM]Cl (1), grafting-from polymerization (8b) with [DMVBIM]Tf₂N (5) and detail of a photograph of the violet colored meander pattern of p([BVBIM]MnO₄) (9) resulting from the salt metathesis of 8a with $KMnO_4$. Please refer to Fig. S16[†] for the complete photograph.

Experimental

Synthesis of 1-butyl-3-(4'-vinylbenzyl)-1H-imidazol-3-ium chloride ([BVBIM]Cl, 1)

The IL was prepared by a slightly modified procedure by Bara *et al.*²⁷ 9.00 g *n*-butylimidazole (72.5 mmol, 1.00 eq.) was dissolved in 100 mL chloroform and cooled to 0 °C. Sub-



sequently, 15.4 mL vinylbenzyl chloride (16.6 g, 108.7 mmol, 1.50 eq.) was added. The mixture was stirred at 50 °C for 24 h. The mixture was extracted with water (3 × 25 mL), evaporated at reduced pressure to get rid of the residue organic solvents, and freeze-dried. 17.5 g of product **1** (63.1 mmol, 87%) was received as a colorless, highly viscous liquid.

¹H NMR (400 MHz, DMSO-*d*₆, 298 K): δ = 9.72 (s, 1 H, N=CH-N), 7.91 (d, ³J = 13.8 Hz, 2 H, Im-H), 7.50 (d, ³J = 8.2 Hz, 2 H, Ph-H), 7.46 (d, ³J = 8.2 Hz, 2 H, Ph-H), 6.72 (dd, ³J(Z) = 11.0 Hz, ³J(E) = 17.7 Hz, 1 H, H₂C=CH), 5.85 (d, ³J(E) = 17.7 Hz, 1 H, HC(H)=CH), 5.49 (s, 2 H, Bn-H), 5.27 (d, ³J(Z) = 11.0 Hz, 1 H, HC(H)=CH), 4.20 (t, ³J = 7.2 Hz, 2 H, N-CH₂-CH₂), 1.92–1.60 (m, 2 H, CH₂-CH₂-CH₂), 1.43–1.02 (m, 2 H, CH₂-CH₂-CH₃), 0.87 (t, ³J(Z) = 7.5 Hz, 3 H, CH₂-CH₃) ppm.

¹³C-NMR (100 MHz, DMSO-*d*₆, 298 K): δ = 137.56 (C), 136.17 (C), 135.93 (CH), 134.30 (CH), 128.67 (2 CH), 126.67 (2 CH), 122.79 (CH), 122.54 (CH), 115.29 (CH₂), 51.65 (CH₂), 48.67 (CH₂), 31.24 (CH₂), 18.79 (CH₂), 13.26 (CH₃) ppm.

Analytical data is in accordance to the results obtained by He *et al.*²⁰

Synthesis of 1-butyl-3-(4'-vinylbenzyl)-1H-imidazol-3-ium bis(trifluoromethanesulfonyl)imide ([BVBIM]Tf₂N, 2). 4.50 g *n*-butylimidazole (36.2 mmol, 1.00 eq.) was dissolved in 100 mL chloroform and cooled to 0 °C. Subsequently, 8.3 g vinylbenzyl chloride (54.4 mmol, 1.50 eq.) was added. The mixture was stirred at 50 °C for 24 h. The mixture was extracted with water (2 × 100 mL), and evaporated at reduced pressure to get rid of the residual organic solvents. Afterwards, 10.4 g LiTf₂N (36.2 mmol, 1.00 eq.) was added. The reaction was allowed to proceed overnight at ambient temperature. The mixture was extracted with ethyl acetate (3 × 75 mL). The combined organic solvents were washed with water (1 × 100 mL), and dried over sodium sulfate. 15.97 g of product **2** (30.7 mmol, 84%) was received as a colorless liquid.

¹H NMR (400 MHz, DMSO-*d*₆, 298 K): δ = 9.29 (s, 1 H, N=CH-N), 7.77 (s, 2 H, Im-H), 7.52 (d, ³J = 8.1 Hz, 2 H, Ph-H), 7.43 (d, ³J = 8.2 Hz, 2 H, Ph-H), 6.74 (dd, ³J(Z) = 11.0 Hz, ³J(E) = 17.7 Hz, 1 H, H₂C=CH), 5.86 (d, ³J(E) = 17.7 Hz, 1 H, HC(H)=CH), 5.42 (s, 2 H, Bn-H), 5.29 (d, ³J(Z) = 11.0 Hz, 1 H, HC(H)=CH), 4.19 (t, ³J = 7.2 Hz, 2 H, N-CH₂-CH₂), 1.79 (quin, ³J = 7.5 Hz, 2 H, CH₂-CH₂-CH₂), 1.28 (m, 2 H, CH₂-CH₂-CH₃), 0.90 (t, ³J = 7.5 Hz, 3 H, CH₂-CH₃) ppm.

¹³C{¹H} NMR (100 MHz, DMSO-*d*₆, 298 K): δ = 139.32 (C), 135.90 (C), 135.83 (CH), 131.52 (CH), 129.36 (2 CH), 127.49 (2 CH), 122.30 (CH), 122.08 (CH), 119.95 (q, ¹J = 318.0 Hz, CF₃), 115.88 (CH₂), 53.59 (CH₂), 50.31 (CH₂), 32.07 (CH₂), 19.49 (CH₂), 13.39 (CH₃) ppm.

Analytical data is in accordance to the results obtained by He *et al.*²⁰

Synthesis of 1,2-dimethyl-3-(4'-vinylbenzyl)-1H-imidazol-3-ium bis(trifluoromethanesulfonyl)imide ([DMVBIM]Tf₂N, 5). A Schlenk tube was charged with 2.00 g 1,2-dimethylimidazole (20.8 mmol, 1.00 eq.), dissolved in 10 mL CHCl₃ and cooled to 0 °C. Subsequently, 3.81 g vinylbenzyl chloride (25.0 mmol, 1.20 eq.) was added and the tube was

heated from 0 °C to 50 °C. The mixture was stirred at 50 °C for 24 h. The mixture was extracted with water (slow phase separation). Afterwards, 7.17 g LiTf₂N (25.0 mmol, 1.2 eq.) was added and the mixture stirred for 3 d at ambient temperature. The mixture was extracted with ethyl acetate (3 × 40 mL) and the organic solvent washed with water (1 × 50 mL). The organic layers were dried over sodium sulfate and the solvent was evaporated at reduced pressure. 8.67 g of product **5** (17.6 mmol, 85%) was received as a slightly orange colored liquid.

¹H NMR (400 MHz, DMSO-*d*₆, 298 K): δ = 7.67 (d, ³J = 20.2 Hz, 2 H, Im-H), 7.51 (d, ³J = 8.2 Hz, 2 H, Ph-H), 7.32 (d, ³J = 8.2 Hz, 2 H, Ph-H), 6.74 (dd, ³J(Z) = 11.0 Hz, ³J(E) = 17.7 Hz, 1 H, H₂C=CH), 5.86 (d, ³J(E) = 17.7 Hz, 1 H, HC(H)=CH), 5.41 (s, 2 H, Bn-H), 5.29 (d, ³J(Z) = 11.0 Hz, 1 H, HC(H)=CH), 3.77 (s, 3 H, N-CH₃), 2.60 (s, 3 H, N-C(CH₃)-N) ppm.

¹³C{¹H} NMR (100 MHz, DMSO-*d*₆, 298 K): δ = 144.66 (C), 137.42 (C), 135.94 (C), 134.04 (CH), 128.12 (2 CH), 124.36 (2 CH), 122.69 (CH), 121.24 (CH), 119.57 (q, ¹J = 321.8 Hz, CF₃), 115.07 (CH₂), 50.46 (CH₂), 34.78 (CH₃), 9.40 (CH₃) ppm.

ESI-MS for C₁₄H₁₇N₂⁺: *m/z* = 213.14.

Synthesis of poly[1-butyl-3-(4'-vinylbenzyl)-1H-imidazol-3-ium chloride] (3). A Schlenk flask was charged with 2.03 g 1-butyl-3-(4'-vinylbenzyl)-1H-imidazol-3-ium chloride (**1**) (7.41 mmol, 200.00 eq.), 1.8 mg 1,1'-azobis(cyclohexanecarbonitrile) (0.00741 mmol, 0.20 eq.) and 12.8 mg 2-cyano-2-propyl dodecyl trithiocarbonate (0.037 mmol, 1.00 eq.). The mixture was dissolved in DMSO, degassed by three consecutive freeze-pump-thaw cycles, filled with nitrogen, and subsequently immersed in a preheated oil bath at 85 °C. The polymerization was stopped by sudden freezing in liquid nitrogen, diluting in water and purging with air. Next, the polymer was purified by dialysis in pure water (3 d, a.t., 1 kDa MWCO).

Synthesis of poly[1-butyl-3-(4'-vinylbenzyl)-1H-imidazol-3-ium bis(trifluoromethanesulfonyl)imide] (4). A Schlenk flask was charged with 8.00 g 1-butyl-3-(4'-vinylbenzyl)-1H-imidazol-3-ium bis(trifluoromethane) sulfonimide (**2**) (15.34 mmol, 200.00 eq.), 3.8 mg 1,1'-azobis(cyclohexanecarbonitrile) (0.0153 mmol, 0.20 eq.) and 26.5 mg 2-cyano-2-propyl dodecyl trithiocarbonate (0.0767 mmol, 1.00 eq.). The mixture was dissolved in BuCN, degassed by three consecutive freeze-pump-thaw cycles, filled with nitrogen and subsequently immersed in a preheated oil bath at 85 °C. The polymerization was stopped by sudden freezing in liquid nitrogen, diluted in acetone and purging with air. After dialysis (pure water or DMSO for 3 d, a.t., 1 kDa MWCO), the product **4** was obtained as yellow substance depending on the residual monomer as a highly viscous liquid or as a solid.

Synthesis of poly[1-butyl-3-(4'-vinylbenzyl)-1H-imidazol-3-ium bis(trifluoromethanesulfonyl)imide] (4) by salt metathesis. 21.0 mg poly(1-butyl-3-(4'-vinylbenzyl)-1H-imidazol-3-ium chloride) (8900 g mol⁻¹, 0.0755 mmol, 1.00 eq.) was dissolved in 1 mL water. Subsequently, 26.0 mg LiTf₂N (0.0906 mmol, 1.20 eq.) was added, and the reaction was allowed to proceed for 24 h at ambient temperature. Next, the product was extracted with ethyl acetate (2 × 5 mL), dried over



sodium acetate, and evaporated at reduced pressure. Product 4 was received as a yellowish solid.

Synthesis of the photoenol-functionalized RAFTagent (7). The synthesis was followed the protocol of Kaupp *et al.*⁴²

Synthesis of 4-maleimidobutyryl chloride. The synthesis was followed the protocol of Herrmann *et al.*⁴⁵

APTES-functionalization of Si wafer. The Si surfaces (approx. $1 \times 1 \text{ cm}^2$) were treated with piranha solution ($\text{H}_2\text{SO}_4:\text{H}_2\text{O}_2$, 4:1 v/v) for 3 h. Subsequently, the Si wafers were rinsed thoroughly with Milli Q water and dried under a stream of nitrogen. In a further step, the activated surfaces were placed in a solution of 5.0 mL APTES, 70.0 mL toluene and 0.1 vol% triethylamine and allowed to react overnight at 80 °C. The substrates were rinsed with toluene, acetone and methanol, before being dried under a stream of nitrogen.

Maleimide-functionalization of (APTES) Si wafer. The APTES-functionalized Si substrate was immersed in 55 mM DCM solution of TEA. Equal volume of a 50 mM solution of 4-maleimidobutyryl chloride in dry dichloromethane was added subsequently under inert atmosphere. The reaction was allowed to proceed for 4 h. The surfaces were rinsed with dichloromethane, methanol and Milli Q water and left in a petri dish with methanol overnight. Next, the surfaces were rinsed with THF, methanol and Milli Q water.

Photo-patterning of DoPAT-PE (7') on the maleimide-functionalized Si wafer (6). The substrate was immersed in a 11 mM solution of DoPAT-PE (7') in acetonitrile, and subsequently degassed by purging with nitrogen for 20 minutes. Next, the wafer was irradiated for 2.5 h (UV lamp: ARIMED B6, $\lambda_{\text{max}} = 320 \text{ nm}$). Subsequently, the wafer was rinsed thoroughly with THF, acetone, methanol and Milli Q water and dried under a stream of nitrogen.

Grafting-from polymerization of butylvinylbenzylimidazolium chloride (1). The DoPAT-PE-functionalized substrate 7 was placed in a vial and flushed with argon for 5 minutes. A solution – degassed *via* three consecutive freeze–pump–thaw cycles – of 5.55 g butylvinylbenzylimidazolium chloride (1) (20.1 mmol, 1500 eq.), 4.7 mg DoPAT (0.0134 mmol, 1.00 eq.) and 3.3 mg VAZO-88 (0.0134 mmol, 1.00 eq.) was added to the wafer and the vial placed in a preheated custom-built heating block at 85 °C. After 7 h, the polymerization was stopped by sudden cooling in liquid nitrogen and bubbled with air. The solution was dialyzed in deionized water for 3 d and freeze-dried afterwards. The wafer was rinsed thoroughly with Milli Q water, methanol, acetone and Milli Q water again and left in a petri dish with methanol overnight. The wafer was again rinsed with Milli Q water, methanol and Milli Q water and left for another 24 h in a petri dish with methanol. Subsequently, the wafer was rinsed with Milli Q water, methanol and Milli Q water and dried under a stream of nitrogen.

Reaction of PIL-grafted surface (8) with KMnO_4 . The PIL-grafted surface was dipped into a 50 mM solution of KMnO_4 in water for 2 to 3 seconds. Subsequently, the surface was rinsed thoroughly with Milli Q water and dried under a stream of nitrogen.

Conclusions

In summary, we provide a useful and general synthetic tool to access imidazolium poly(ionic liquids) with styrenic IL monomer units *i.e.* p([BVBIM]Cl) and p([BVBIM]Tf₂N) *via* the RAFT polymerization technique. The mass spectrometric in-depth characterization of poly(ionic liquid)s is described giving a detailed account of the molecular structure of these polyelectrolytes. In addition, we present a light-induced photoenol driven avenue to provide spatially resolved functional interfaces. The results of our contribution will facilitate access to the discussed class of poly(ionic liquid)s to further understand their molecular weight dependent physical properties and functions. The surface patterning method can be applied to develop task-specific interfaces.

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Notes and references

- N. V. Plechkova and K. R. Seddon, *Chem. Soc. Rev.*, 2008, **37**, 123–150.
- M. D. Green and T. E. Long, *Polym. Rev.*, 2009, **49**, 291–314.
- O. Green, S. Grubjesic, S. Lee and M. A. Firestone, *Polym. Rev.*, 2009, **49**, 339–360.
- J. Lu, F. Yan and J. Texter, *Prog. Polym. Sci.*, 2009, **34**, 431–448.
- D. Mecerreyes, *Prog. Polym. Sci.*, 2011, **36**, 1629–1648.
- J. Yuan, D. Mecerreyes and M. Antonietti, *Prog. Polym. Sci.*, 2013, **38**, 1009–1036.
- Y. Gu and T. P. Lodge, *Macromolecules*, 2011, **44**, 1732–1736.
- H. He, S. Averick, E. Roth, D. Luebke, H. Nulwala and K. Matyjaszewski, *Polymer*, 2014, **55**, 3330–3338.
- H. Mori, M. Yahagi and T. Endo, *Macromolecules*, 2009, **42**, 8082–8092.
- J. Texter, V. A. Vasantha, R. Crombez, R. Maniglia, L. Slater and T. Mourey, *Macromol. Rapid Commun.*, 2012, **33**, 69–74.
- K. Vijayakrishna, S. K. Jewrajka, A. Ruiz, R. Marcilla, J. A. Pomposo, D. Mecerreyes, D. Taton and Y. Gnanou, *Macromolecules*, 2008, **41**, 6299–6308.
- J. Yuan, H. Schlaad, C. Giordano and M. Antonietti, *Eur. Polym. J.*, 2011, **47**, 772–781.
- A. Wilke, J. Yuan, M. Antonietti and J. Weber, *ACS Macro Lett.*, 2012, **1**, 1028–1031.



- 14 J. E. Bara, C. J. Gabriel, E. S. Hatakeyama, T. K. Carlisle, S. Lessmann, R. D. Noble and D. L. Gin, *J. Membrane Sci.*, 2008, **321**, 3–7.
- 15 Z. Shi, B. S. Newell, T. S. Bailey and D. L. Gin, *Polymer*, 2014, **55**, 6664–6671.
- 16 F. Jiang, C. Li, H. Fu, Y. Wang, X. Guo and G. Wu, *Sci. China Chem.*, 2015, **58**, 716–721.
- 17 E. Karjalainen, D. F. Izquierdo, V. Marti-Centelles, S. V. Luis, H. Tenhu and E. Garcia-Verdugo, *Polym. Chem.*, 2014, **5**, 1437–1446.
- 18 D. Cordella, A. Kermagoret, A. Debuigne, C. Jérôme, D. Mecerreyes, M. Isik, D. Taton and C. Detrembleur, *Macromolecules*, 2015, **48**, 5230–5243.
- 19 P. Coupillaud, J. Pinaud, N. Guidolin, J. Vignolle, M. Fèvre, E. Veaudecenne, D. Mecerreyes and D. Taton, *J. Polym. Sci., Part A: Polym. Chem.*, 2013, **51**, 4530–4540.
- 20 H. He, M. Zhong, B. Adzima, D. Luebke, H. Nulwala and K. Matyjaszewski, *J. Am. Chem. Soc.*, 2013, **135**, 4227–4230.
- 21 D. Cordella, A. Kermagoret, A. Debuigne, R. Riva, I. German, M. Isik, C. Jérôme, D. Mecerreyes, D. Taton and C. Detrembleur, *ACS Macro Lett.*, 2014, **3**, 1276–1280.
- 22 D. England, N. Tambe and J. Texter, *ACS Macro Lett.*, 2012, **1**, 310–314.
- 23 Q. Ye, T. Gao, F. Wan, B. Yu, X. Pei, F. Zhou and Q. Xue, *J. Mater. Chem.*, 2012, **22**, 13123–13131.
- 24 I. Abdelhedi-Miladi, D. Montarnal, M. M. Obadia, H. Ben Romdhane and E. Drockenmuller, *ACS Macro Lett.*, 2014, **3**, 1187–1190.
- 25 C. Barner-Kowollik, *Handbook of RAFT Polymerization*, Wiley-VCH, Weinheim, Germany, 2008.
- 26 Y. Li, G. Li, X. Wang, Z. Zhu, H. Ma, T. Zhang and J. Jin, *Chem. Commun.*, 2012, **48**, 8222–8224.
- 27 J. E. Bara, S. Lessmann, C. J. Gabriel, E. S. Hatakeyama, R. D. Noble and D. L. Gin, *Ind. Eng. Chem. Res.*, 2007, **46**, 5397–5404.
- 28 E. H. I. J. Brandrup and E. A. Grulke, *Polymer Handbook*, John Wiley and Sons, Weinheim, Germany, 1999.
- 29 R. Marcilla, J. A. Blazquez, R. Fernandez, H. Grande, J. A. Pomposo and D. Mecerreyes, *Macromol. Chem. Phys.*, 2005, **206**, 299–304.
- 30 A. S. Goldmann, M. Glassner, A. J. Inglis and C. Barner-Kowollik, *Macromol. Rapid Commun.*, 2013, **34**, 810–849.
- 31 J. Fenn, M. Mann, C. Meng, S. Wong and C. Whitehouse, *Science*, 1989, **246**, 64–71.
- 32 A. Calderaro, M.-C. Arcangeletti, I. Rodighiero, M. Buttrini, C. Gorrini, F. Motta, D. Germini, M.-C. Medici, C. Chezzi and F. De Conto, *Sci. Rep.*, 2014, **4**, 6803–6813.
- 33 P. A. Hunt, C. R. Ashworth and R. P. Matthews, *Chem. Soc. Rev.*, 2015, **44**, 1257–1288.
- 34 J. Pinaud, J. Vignolle, Y. Gnanou and D. Taton, *Macromolecules*, 2011, **44**, 1900–1908.
- 35 M. C. Kroon, W. Buijs, C. J. Peters and G.-J. Witkamp, *Thermochim. Acta*, 2007, **465**, 40–47.
- 36 H. C. M. Byrd and C. N. McEwen, *Anal. Chem.*, 2000, **72**, 4568–4576.
- 37 D. V. Zagorevskii, M. J. Nasrullah, V. Raghunadh and B. C. Benicewicz, *Rapid Commun. Mass Spectrom.*, 2006, **20**, 178–180.
- 38 M. Dietrich, M. Glassner, T. Gruending, C. Schmid, J. Falkenhagen and C. Barner-Kowollik, *Polym. Chem.*, 2010, **1**, 634–644.
- 39 B. Yameen, C. Rodriguez-Emmenegger, C. M. Preuss, O. Pop-Georgievski, E. Verveniotis, V. Trouillet, B. Rezek and C. Barner-Kowollik, *Chem. Commun.*, 2013, **49**, 8623–8625.
- 40 M. Zamfir, C. Rodriguez-Emmenegger, S. Bauer, L. Barner, A. Rosenhahn and C. Barner-Kowollik, *J. Mater. Chem. B*, 2013, **1**, 6027–6034.
- 41 C. M. Preuss, A. S. Goldmann, V. Trouillet, A. Walther and C. Barner-Kowollik, *Macromol. Rapid Commun.*, 2013, **34**, 640–644.
- 42 M. Kaupp, T. Tischer, A. F. Hirschbiel, A. P. Vogt, U. Geckle, V. Trouillet, T. Hofe, M. H. Stenzel and C. Barner-Kowollik, *Macromolecules*, 2013, **46**, 6858–6872.
- 43 R. Gracia, K. Vijayakrishna and D. Mecerreyes, *React. Funct. Polym.*, 2014, **79**, 54–58.
- 44 Y.-J. Lu, W.-L. Wong and C.-F. Chow, *Catal. Commun.*, 2015, **69**, 25–28.
- 45 A. Herrmann, G. Mihov, G. W. M. Vandermeulen, H.-A. Klok and K. Müllen, *Tetrahedron*, 2003, **59**, 3925–3935.

