Reversible crosslinking and fast stress relaxation in dynamic polymer networks via transalkylation using 1,4-diazabicyclo[2.2.2]octane†

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Dynamically crosslinked polymers offer the possibility to design materials that have the robustness of thermosetting materials and the reshapability of thermoplastic materials. A reversible crosslinker based on transalkylation chemistry using 1,4-diazabicyclo[2.2.2]octane (DABCO) was developed. The kinetics of the exchange reaction of benzyl bromide with DABCO was studied in detail by varying molar ratios and temperature. The system showed efficient exchange without addition of a catalyst. Reversibility in a polymer was studied by the crosslinking of a linear, benzyllic bromide functionalized polyacrylate with DABCO. The resulting network was insoluble in common solvents, but it readily dissolved in the presence of DABCO. Rheological measurements showed that the crosslinks are dynamic, with a strong temperature dependent relaxation that at 140 °C is faster than for other transalkylation based dynamic networks reported in the literature.

Introduction

Thermosetting polymers are covalently crosslinked networks with excellent properties such as solvent resistance and high thermal and dimensional stability.1,2 Although these properties are highly advantageous for a wide range of applications such as in the coatings,3 adhesives,4 construction,5 and automotive6 industries, recycling or reprocessing these products is very challenging because they are unable to flow upon heating. By introducing exchangeable crosslinks, enhanced properties such as reshapability,7 recyclability,7,8 (self-)healing,9,10 or stress relaxation can be obtained while retaining the advantageous properties like enhanced thermal and chemical stability.11,12 Networks containing specially designed reversible covalent crosslinks are also known as covalent adaptable networks (CANs).8,11,13–15 Depending on the nature of the exchangeable bond, a stimulus such as light,20 temperature,21 or pH triggers the reversibility of the bond.15 Many dynamic motifs for bond exchange have been developed over the past years, some examples are transesterification reactions,7,22–26 boronic ester exchange,27–29 disulfide exchange,30 controllable reversible addition–fragmentation reactions,31–33 transcarbonylation exchange,34 and (vinyllogous) urethane exchange.35–38 Because nucleophilic substitutions are usually considered to be non-reversible, they have often been overlooked for use in CANs.15 However, the reversibility of specific nucleophilic substitutions has been known for years,39 and has recently been employed in CANs.40 Kulchat and Lehn investigated the transcarbonylation of quaternary ammonium salts,41 and showed their potential for the development of dynamic covalent chemistry. Droge and coworkers were the first to report a dynamic polymeric network based on transalkylation exchange of C–N bonds.42 Network polymers containing 1,2,3-triazolium salts as reversible crosslinks were synthesized and could be recycled by compression molding. After two recycling cycles, samples with an extension at break similar to the starting material were obtained, indicating efficient network reformation. Furthermore, efficient stress relaxation was observed in rheological experiments. At a temperature of 200 °C, the characteristic relaxation time τ was only a few seconds. Following this work, Konkolewicz and coworkers designed acrylic polymeric materials which were dynamically crosslinked with anilinium salts.43 These dynamic polymeric materials show excellent malleability at 60 °C and moreover could be quantitively healed by mild heating to 60 °C or 80 °C. A more detailed kinetic study elucidated the exchange mechanism and it was found that although the exchange follows a dissociative pathway, they have low degrees...
of de-crosslinking and therefore the networks have properties resembling those of associative CANs.\textsuperscript{44} Du Prez and coworkers developed a reversible network via C–S transalkylation of trialkylsulfonium salts.\textsuperscript{45} With kinetic studies they showed that alkylbrosylates are efficient transalkylation agents to convert thioesters into trialkylsulfonium salts. These salts showed complete relaxation at elevated temperatures in rheological stress relaxation experiments, and remolding by compression molding yielded defect free samples. Recycled samples showed no significant change in mechanical properties when measured in DMTA and tensile tests. Zhang and coworkers utilized similar C–S transalkylation reactions to recycle sulfur-cured rubber.\textsuperscript{46}

Inspired by these systems, we aimed at developing a transalkylation system that can be implemented in (meth)acrylate polymers. Multifunctional (meth)acrylate polymers are a typical example of thermosetting polymers and due to their fast curing behavior and compatibility with many different functionalities,\textsuperscript{47} they have many industrial applications e.g. in dentistry,\textsuperscript{48} coatings,\textsuperscript{3,49} liquid-crystal displays,\textsuperscript{50} inks and adhesives.\textsuperscript{4,51} The use of a two-step approach of linear polymerization followed by reversible crosslinking offers opportunities for the introduction of dynamic behavior into commercially available methacrylic polymers.\textsuperscript{38} A number of different tertiary amines were considered as reversible crosslinking agent. 1,4-Diazabicyclo[2.2.2]octane (DABCO) was selected as the most promising candidate for fast exchange reactions (Scheme 1). Compared to the popular nucleophilic catalyst 4-dimethylaminopyridine (DMAP), a pyridine derivate, the bicyclic aliphatic amine DABCO is a significantly better nucleophile, yet also a much better leaving group (by a factor of $10^3$ and $10^6$ respectively)\textsuperscript{52} will transalkylate much faster than DMAP derivatives. An additional advantage is that one DABCO molecule can form a crosslink between two polymer chains, because it contains two tertiary amine functional groups. Finally, DABCO can easily be alkylated at low temperatures, making the alkylation process compatible with (meth)acrylates.\textsuperscript{53}

The work reported here investigates the potential of DABCO in transalkylation reactions to create dynamically crosslinked acrylate networks. The kinetics of the exchange reaction of benzyl bromide with DABCO was studied in detail using a small molecule model system. Additionally, we demonstrate the formation of a dynamic covalent network by crosslinking a linear benzyl bromide functionalized methacrylic prepolymer in a single step with DABCO. The cured network can be reprocessed and shows efficient stress relaxation in rheological measurements.

### Experimental

#### Materials

All commercial chemicals and solvents were used as received, unless stated otherwise. 4-(Bromomethyl)benzoic acid and 2-hydroxyethyl methacrylate were obtained from TCI EUROPE. All other chemicals for the synthesis were purchased from Sigma Aldrich. Solvents were obtained from Biosolve.

#### Characterization methods

**Differential scanning calorimetry (DSC).** $T_g$ of the polymers was measured using a TA Instruments Q2000 differential scanning calorimeter equipped with an RCS90 cooling accessory using aluminum hermetic pans. 5–10 mg of sample was used per measurement. The general method consisted of four cycles in which a sample was measured from $-50$ to $200 \, ^\circ\text{C}$ with heating and cooling rates of 10 $^\circ\text{C} \, \text{min}^{-1}$ (cycles 1 and 2), 20 $^\circ\text{C} \, \text{min}^{-1}$ and 40 $^\circ\text{C} \, \text{min}^{-1}$. The first heating run was discarded. Universal Analysis TRIOS software was used for data acquisition, midpoint at half height was used to determine $T_g$.

**Gel fraction.** The mass fraction of the crosslinked part of the material is defined as the gel fraction. Approximately 250 mg of crosslinked material was weighed and immersed in 10 mL THF for 75 hours at room temperature. The samples were filtered and dried in a vacuum oven at 60 $^\circ\text{C}$ in a vacuum oven. The gel fraction was determined using the following equation:\textsuperscript{54}

$$\text{Gel fraction} = \frac{m_{\text{dry}}}{m_{\text{initial}}}$$

where $m_{\text{initial}}$ is the initial weight of the sample, and $m_{\text{dry}}$ is the mass of the sample after extraction and subsequent drying.

**Matrix assisted laser absorption/ionization-time of flight mass spectra (MALDI-TOF).** Molar masses of the synthesized small molecules were determined using a Bruker Autoflex Speed mass spectrometer using α-cyano-4-hydroxycinnamic acid as matrix. The lactones were measured in positive mode, and the alkylation systems were measured in negative mode using 2-[4-(dimethylamino)phenyl]benzoic acid (DAPB) as matrix. The peak of interest was extracted from the spectra using MASCOT software and compared against the spectra of known compounds.

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**Scheme 1** Schematic representation of the topology rearrangements due to the transalkylation of DABCO by pendant benzyl bromide groups.
acid or trans-2-[3-(4-tert-butylphenyl)-2-methyl-2-propenylidine]-malononitrile as a matrix.

**Nuclear magnetic resonance spectroscopy (NMR).** $^1$H and $^{13}$C spectra were recorded on a 400 MHz Bruker Advance III HD (400 MHz for $^1$H NMR) spectrometer. $^1$H In situ NMR spectra were recorded using a Varian Inova 500 MHz spectrometer. Chemical shifts (δ) are expressed in ppm with respect to tetramethylsilane (TMS, 0 ppm) as an internal standard. Coupling constants are reported as J-values in Hz.

**Rheology.** The mechanical properties and stress relaxation behavior of disk-shaped samples were determined using a stress-controlled AR-G2 rheometer (TA instruments) using 8 mm parallel plate geometry. Time sweeps at constant temperature were performed before and after relaxation experiments using a strain of 1% with a frequency of 1 Hz. Stress relaxation experiments were performed at a temperature range between 110 °C and 140 °C with a strain of 1% using a constant normal force of 10 N to ensure contact. A strain of 1% is within the linear viscoelastic regime.

**Size exclusion chromatography (SEC).** Molar mass distributions of the linear polymers were measured on a Shimadzu Prominence-LC-2030C 3D equipped with two Agilent columns (mixed C and mixed D) with a combined range of 200-2 000 000 g mol$^{-1}$. The used eluent was THF with a flow rate of 1 mL min$^{-1}$ at 40 °C. The molar mass was determined relative to narrow dispersity polystyrene standards purchased from Agilent.

**Thermogravimetric analysis.** Thermal stability studies were performed in a TA instruments TGA Q500 machine. Samples were heated under nitrogen flow (60 mL min$^{-1}$) from 26 °C to 600 °C with a heating rate of 10 °C min$^{-1}$.

**Synthesis of small molecules**

The synthesis and characterization of 1,4-dibenzyl-1,4-diazabicyclo[2.2.2]octane-1,4-dimethacrylate (DABCO-polyacrylate) were characterized by reactivity ratios close to 1, no significant composition drift is expected for this copolymerization. The overall composition of the copolymer was determined by peak integration of the $^1$H NMR spectra, and was found to contain 10% BrEMA. Since copolymerization of two methacrylates are characterized by reactivity ratios close to 1, no significant composition drift is expected for this copolymerization.

**SEC analysis** showed a single peak with a $M_n = 33 \times 10^3$ g mol$^{-1}$ and a PDI $\approx 1.7$ with respect to polystyrene standards.

**Network formation**

The linear poly(BMA-co-BrEMA) (7 g) was dissolved in 70 mL tetrahydrofuran using mild sonication in an ultrasonic bath. The desired amount of DABCO (0.3 mg for a ratio of 0.6 mol DABCO per benzyl bromide group) was added and sonicated for 2 minutes, to ensure homogeneous mixing. The liquid mixture was heated for 1 hour at 60 °C resulting in the formation of an organogel. The gel was transferred to aluminum molds, and solvent was removed using a vacuum oven at 70 °C overnight.

**Sample preparation**

The crosslinked polymer was compression molded into disk or bar shaped samples using a Collin Press 300 G with an operating temperature of 140 °C, pressure of 100 bar for 20 minutes and subsequently cooled with water. To remove any thermal history prior to rheology experiments, the samples were ther-
mally treated by placing them into an oven overnight at 120 °C.

Results and discussion

Model experiments

To evaluate the efficiency of the transalkylation reaction of DABCO a kinetic study on low molecular weight model compounds was performed. First model compound $S$, (Scheme 2) was prepared by the alkylation of DABCO with two molecules of benzyl bromide (see ESI† for synthetic details). For the exchange experiments, DABCO ($D$), was added to compound $S$ in DMSO.

The formation of the monobenzyl-DABCO product ($P$) was monitored using $^1$H NMR. In Fig. 1a, the changes in the relevant NMR signals for the determination of the concentrations of $S$ and $P$ are shown (see also Fig. S1, ESI†).

The concentrations of the product, monobenzyl-DABCO ($P$) are plotted as a function of reaction time for a range of initial concentrations at a temperature of 60 °C in Fig. 1b and a range of temperatures for equimolar initial concentrations of 40 mM in Fig. 1c.

Quantitative analysis of the results shown in Fig. 1 are consistent with a reaction rate that is first-order in both reactant concentrations, and second-order rate coefficients were determined at a range of temperatures (see ESI† for more details). An Arrhenius analysis leads to an activation energy of 93 kJ mol$^{-1}$ and frequency factor $A \approx 7 \times 10^{11}$ L mol$^{-1}$ s$^{-1}$ for the second-order forward reaction rate coefficient $k$.

In a dynamic polymer network, the nucleophile in the exchange reaction is not DABCO, but a dangling DABCO group that is quaternized on one of the nitrogen atoms. Since the positive charge may be expected to reduce the nucleophilicity of the second nitrogen atom through electrostatic repulsion, it was relevant to investigate the reactivity of a monosubstituted DABCO with bisbenzyl-DABCO. It was found that monosubstituted DABCO is 3 to 5 times less reactive due to electrostatic effects, but still shows efficient exchange (for details, see ESI Fig. S3†).

To investigate whether exchange reactions take place in the absence of free amine, the reaction of $S$ with excess alkylating agent 4-methylbenzyl bromide was performed (Scheme 3). Surprisingly, exchange reactions occurred and a mixed bisbenzylated product was formed, albeit at a significantly lower rate than for the reaction of $S$ with DABCO under the same conditions (Fig. S4, ESI†).

![Scheme 2](image)

Scheme 2 Exchange reaction of bisbenzyl-DABCO ($S$) and free DABCO ($D$) for kinetic study in DMSO.

![Fig. 1](image)

Fig. 1 (a) NMR signals for the determination of the concentrations of monobenzyl-DABCO ($P$) and bisbenzyl-DABCO ($S$), $T = 60$ °C and $[S]_0 = [D]_0 = 40$ mM. (b) Formation of $P$ from $S$ and DABCO ($D$) in DMSO for a range of different initial concentrations at 60 °C, data labels in the plot denote concentrations $[S]_0 = [D]_0$ in mM. (c) Temperature dependence of the formation of $P$ using initial concentrations $[S]_0 = [D]_0 = 40$ mM.

![Scheme 3](image)

Scheme 3 Exchange reaction of bisbenzyl-DABCO excess alkylating agent.

The occurrence of trans-alkylation in the absence of free amino groups is in line with the work of Lehn and coworkers, who propose two pathways for transalkylation of
quaternary amines, a mechanism with direct attack of the amine on the benzylic carbon atom, and an indirect mechanism in which the bromide counterion acts as a nucleophile to produce benzyl bromide and an amine (Scheme S1, ESI†). Our results strongly suggest that both the indirect and direct pathway occur with the direct one being dominant. Other dynamic networks based on trans-N-alkylation reactions like those reported by Konkolewicz44 and Drockenmuller42,56 seem to follow a dissociative mechanism but nevertheless have low degrees of de-crosslinking resulting in properties resembling those of associative CANs.57

Polymer synthesis

After the successful transalkylation reaction in the model study we aimed at using this chemistry to reversibly crosslink acrylic polymers. To this end, a monomer with a benzyl bromide functional group was synthesized via an esterification reaction of 4-(bromomethyl) benzoic acid and 2-hydroxyethylmethacrylate. This monomer (BrEMA) was copolymerized with n-butyl methacrylate (BMA) in a free radical polymerization at 50 °C, using AIBN as an initiator and decanethiol (DT) as a chain transfer agent (Scheme 4) to yield a linear polymer with benzyl bromine side groups (SEC: \(M_n = 3.3 \times 10^3\) g mol\(^{-1}\) and \(D \approx 1.7\), Fig. 2a). Butyl methacrylate was selected as the comonomer to obtain a polymer with a low \(\mathcal{T}_g\) to facilitate chain mobility necessary for the exchange.58

The copolymer composition was determined from \(^1\)H NMR in THF-\(d_8\) (Fig. S5, ESI†) and it was found that the synthesized polymer contained 10 mol% of BrEMA units. The \(\mathcal{T}_g\) of the copolymer was 24 °C, as determined by DSC, Fig. 2b. The TGA thermogram, Fig. 2c, showed the typical thermal stability of a methacrylate polymer with significant thermal degradation from 260 °C onwards.

Network formation

After having established that DABCO and benzylbromide also efficiently react in THF (see ESI, Fig. S6†), crosslinking experiments with polymer were carried out in THF. The linear polymer readily dissolved in THF (700 mg in 7 mL) resulting in a clear transparent solution, Fig. 3. Upon addition of DABCO (29 mg), the mixture became slightly opaque. When left to react at room temperature the viscosity of the mixture slowly increased until after 1.5 hours a gel was obtained that was able to support its own weight upon tilting the vial. When reacted at 60 °C a gel was obtained after only 10 minutes (Fig. 3). An attempt was made to show the presence of DABCO adducts in the polymer by \(^1\)H and \(^{13}\)C NMR. A small amount of DABCO crosslinker was added to a solution of linear copolymer in THF-\(d_8\) (70 mg polymer, 0.49 mg DABCO; this corresponds to 0.0043 mmol DABCO and 0.043 mmol benzyl bromide).

After 1 hour of reaction at 60 °C an increase in viscosity was observed and the sample was analyzed by NMR.
Due to the low concentration of crosslinks, a signal of the adduct could not be observed in NMR. The DOSY NMR spectra in Fig. 4a show a clear decrease in diffusion coefficients when the copolymer was reacted with DABCO, indicating that chains are linked together by DABCO.\textsuperscript{59,60}

In order to obtain a network for mechanical testing, a batch of 7 g of linear polymer was reacted with 0.3 mg of DABCO.

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**Fig. 3** After addition of DABCO to a solution of linear polymer in THF a gel is formed. When reacted at room temperature, top pictures, the viscosity slowly increases over time until a gel is formed. When reacted at 60 °C a gel is obtained after 10 min.

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**Fig. 4** Overlay of DOSY spectra of the linear copolymer and copolymer reacted with DABCO.
(approximately 0.6 DABCO per benzyl bromide group) as a crosslinking agent by heating for 1 hour at 60 °C in THF. The resulting organogel was dried, yielding a solid polymer. The $T_g$ after crosslinking was 32 °C which is slightly higher than that of the linear polymer (Fig. S7a, ESI†). The thermal stability decreased slightly as the crosslinked polymer starts degrading from 180 °C (Fig. S7b, ESI†). To quantitatively check the formation of a network, the gel fraction was determined by extraction with THF at room temperature. The linear polymer readily dissolved in THF but after crosslinking, the gel fraction was ≥98%. This high gel fraction confirms the presence of a highly branched and/or network polymer.

The temperature dependence of the moduli was studied using oscillatory rheometry (Fig. 5).

For the linear copolymer the change from rubbery to viscous behavior is clearly seen around 95 °C. For the crosslinked system the $G'$ is larger than $G''$ for the whole temperature range again confirming the presence of a network.

**Dynamic behaviour**

After crosslinking, the polymer was shaped into disks and bars by compression molding at elevated temperatures. This is already indicative of the presence of dynamic crosslinks as this type of processing is not possible for classical thermosets. When compression molded samples were heated to 100 °C, a clear difference between the linear polymer and the crosslinked polymer was observed. The non-crosslinked sample flowed at 100 °C even when no stress was applied (Fig. 6a), while the crosslinked sample maintained its shape even when heated up to 250 °C (Fig. 6b). In addition to the processability of the crosslinked sample by compression molding, the reversibility of the network was further confirmed by treating a sample with excess DABCO in THF. When this mixture was kept overnight at room temperature, the polymer bar could be taken out in one piece, but heating to 60 °C resulted in full dissolution of the bar in 120 minutes (Fig. 6c).

For a quantitative study on the reversibility of these systems, stress relaxation experiments were performed in a rheometer. Networks with 0.6 and 0.4 mol of DABCO per mol of benzyl bromide group were tested, corresponding to excess amine and excess benzyl bromide, respectively. The experiments were performed in the linear viscoelastic regime at 1% strain. The time dependent stress relaxation moduli $G(t)$ are shown at four different temperatures for both systems in Fig. 7. Both systems show efficient stress relaxation with a rate that strongly increases with temperature, and is faster in the system with excess amine, in accordance with the stoichiometry dependence in the small molecule kinetic studies. In contrast to stress relaxation in a number of other CANs,38,45,61 relaxation could not be adequately described by a simple exponential function (characteristic of a Maxwell model). By fitting the data to a stretched exponential, (eqn (2)) starting at a time of 1s, good fits were obtained with $R^2 \geq 0.99$ (see Fig. S9 ESI†). From the deviation from a stretched exponential below 1s, that can be seen in the plots of Fig. 7, it is evident that a
fast component contributes to relaxation at short times. The best fit parameters are shown in Table 1.

\[ G(t) = G_0 e^{-\left(\frac{t}{\tau}\right)^{\beta}} \]  

(2)

Relaxation times are strongly temperature dependent, and the stretching parameters changed relatively little with temperature. Therefore, we used the fitted relaxation times, \( \tau_{\text{fit}} \), to calculate activation energies using the Arrhenius rate law (eqn (3)).

\[ \ln(\tau_{\text{fit}}) = \frac{E_a}{RT} + \ln(\tau_0) \]  

(3)

Values of 216 kJ mol\(^{-1}\) for the system with excess amine and 266 kJ mol\(^{-1}\) for the system with excess benzyl bromide, were obtained (Fig. S10, ESI†).

<table>
<thead>
<tr>
<th>Temperature [°C]</th>
<th>0.6 eq. DABCO</th>
<th>0.4 eq. DABCO</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>( G_0 ) [MPa]</td>
<td>( \tau_{\text{fit}} ) [s]</td>
</tr>
<tr>
<td>110</td>
<td>7.2</td>
<td>484</td>
</tr>
<tr>
<td>120</td>
<td>6.6</td>
<td>42</td>
</tr>
<tr>
<td>130</td>
<td>6.4</td>
<td>9</td>
</tr>
<tr>
<td>140</td>
<td>4.1</td>
<td>3</td>
</tr>
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</table>

These activation energies for mechanical relaxation are significantly higher than the activation energy for the chemical reactions obtained from the small molecule kinetics. The difference is due to the influence of polymer dynamics on the overall rate of transalkylation, but also contains a contribution from differences in polarity between the polymer matrix and highly polar DMSO in the model reaction, which strongly affects the activation energy of the chemical reaction.

Mechanical relaxation in this material also has a higher activation energy and a lower characteristic relaxation time, \( \tau \) at 140 °C than observed in other transalkylation networks (Table S2, ESI†). The triazolium based network reported by Drockenmüller and coworkers has a much lower activation energy (140 kJ mol\(^{-1}\)) and much higher relaxation times \( \tau \) at 140 °C.\(^{42}\) For other systems the activation energy is even lower and relaxation times are longer. It is worth noting here that in a very recent publication\(^{63}\) on the triazolium system the use of benzylic N-substituents resulted in relaxation times that were about two orders of magnitude lower than those in the systems with only aliphatic N-substituents, but all of the investigated systems had activation energies between 126 and 155 kJ mol\(^{-1}\). These results may suggest that part of the fast relaxations that are observed in the current studies may be caused by the fact we used a benzyl bromide substituent and not an alkyl bromide substituent.

In Fig. 8, we compare the temperature dependence of our current system with a selection of previously published systems. It is clear that, at least within this selection, the current DABCO system is quite unique with its pronounced temperature dependence. This is highly beneficial for
processing purposes as exchange is fast at high temperatures but the material is stable at the use temperature.

**Conclusions**

In summary, we have demonstrated that the alkylation of benzyl bromide with DABCO creates a dynamic bond with efficient exchange at elevated temperatures. A linear polymer with benzyl bromide side groups was crosslinked using DABCO as a divalent crosslinker without addition of a catalyst, resulting in a highly crosslinked polymer network. The polymer networks with excess of amine and excess of bromine both showed efficient stress relaxation in rheological step-stress relaxation experiments. At 140 °C, relaxation times for both stoichiometries are faster than observed in a range of systems based on transesterification reactions.65 Several orders of magnitude faster than observed in a range of systems based on transesterification reactions.65–68 Since the networks are made from easily accessible chemicals and the exchange reaction is efficient at relatively low temperatures, this system offers a simple route towards moldable, recyclable network polymers. We expect that changing size and nucleophilicity of the counterion will give access to networks with a wide range of relaxation temperatures.

**Conflicts of interest**

There are no conflicts to declare.

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**References**

