Dynamic Polyamide Networks via Amide–Imide Exchange

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ABSTRACT: The diamide–imide equilibrium was successfully exploited for the synthesis of dynamic covalent polymer networks in which a dissociative bond exchange mechanism leads to high processibility at temperatures above \( \approx 110 \, ^\circ C \). Dynamic covalent networks bridge the gap between thermostsets and thermoplastic polymers. At the operating temperature, when the network is fixed, dynamic covalent networks are elastic solids, while at high temperatures, chemical exchange reactions turn the network into a processible viscoelastic material. Upon heating a dissociative network, the viscosity may also decrease due to a shift of the chemical equilibrium; in such materials, the balance between processibility and excessive flow is important. In this study, a network is prepared that upon heating to above \( \approx 110 \, ^\circ C \) dissociates to a significant extent due to a shift in the amide–imide equilibrium of a bisimide, pyromellitic diimide, in combination with poly(tetrahydrofuran) diamines. At room temperature, the resulting materials are elastic rubbers with a tensile modulus of 2–10 MPa, and they become predominantly viscous above a temperature of approximately 110 \( ^\circ C \), which is dependent on the stoichiometry of the components. The diamide–imide equilibrium was studied in model reactions with NMR, and variable temperature infrared (IR) spectroscopy was used to investigate the temperature dependence of the equilibrium in the network. The temperature-dependent mechanical properties of the networks were found to be fully reversible, and the material could be reprocessed several times without loss of properties such as modulus or strain at break. The high processibility of these networks at elevated temperatures creates opportunities in additive manufacturing applications such as selective laser sintering.

INTRODUCTION

Dynamic covalent networks (DCNs) are polymer networks that combine the advantages of thermosts and thermoplastic polymers due to the presence of chemically reversible cross-links. While DCNs possess typical features of thermosts, such as chemical resistance and dimensional stability, they may obtain thermoplastic characteristics under an external stimulus, such as heat or light, which allows them to be recycled and reprocessed. In the past decade, a large variety of reversible chemical reactions have been explored for the synthesis of DCNs, including cycloadditions, trans-esterification, transamination, olefin metathesis, disulfide exchange, thiol–thioester exchange, siloxane–silanol exchange, imine–amine exchange, and other reactions.

Many reversible reactions used in the synthesis of DCNs are associative in nature, and the resulting materials have been coined “vitrimers” because their temperature-dependent viscosity follows Arrhenius behavior, reminiscent of that of SiO\(_2\) glass. Because the cross-link density is constant, vitrimers generally are solvent resistant and retain structural integrity even at high temperatures.

In dissociative DCNs, the cross-link density decreases with temperature due to a shift of the equilibrium. Nevertheless, an Arrhenius relationship between viscosity and temperature exists when the fraction of dissociated functional groups remains small. If a significant fraction of exchangeable groups dissociates upon heating, the viscosity drops more strongly with temperature, following Williams–Landell–Ferry (WLF) behavior, similar to what is found for linear polymers around and above their glass transition temperature. Such dissociative networks offer the prospect of having a low-viscosity regime that facilitates melt processing, but excessive dissociation will give rise to undesirable flow. The challenge in these materials is to strike a balance between processibility and material integrity to provide a sufficiently wide processing window.

Recently, neighboring group participation has been introduced as a strategy to accelerate exchange reactions in DCNs. Involvement of a functional group close to the exchanging moiety increases the reaction rate by providing a low free energy pathway to exchange via formation of a cyclic...
intermediate, a dissociative process. In their seminal work, Du Prez and co-workers showed a neighboring carboxylic acid group-participated transesterification of a phthalic acid derivative via a reactive anhydride intermediate.\textsuperscript{28} Faster transesterification in networks with neighboring group participation resulted in higher stress relaxation rates as compared to analogous networks without the ortho carboxylic acid group. Subsequently, our group incorporated an ortho sulfonic acid group as the neighboring group in aromatic polyesters,\textsuperscript{29} resulting in even faster relaxation and improved recyclability. Temperature-dependent relaxation showed Arrhenius behavior for these networks, which are therefore vitrimer-like. Arrhenius-like relaxation behavior was also reported in dissociative DCNs based on transamidation with neighboring group participation in an acetoacetyl-based amide.\textsuperscript{30}

Although the introduction of neighboring groups in these networks has led to faster dynamics, accessing the low-viscosity regime of dissociative networks also requires tuning thermodynamics. The equilibrium must be shifted to the cyclic intermediate to lower the cross-link density at high temperatures.

In the current work, we introduce DCNs based on transamidation via a relatively stable imide intermediate and we show that these networks combine substantial exchange rates at intermediate temperatures with a pronounced shift in the diamide−imide equilibrium that leads to network dissociation at high temperatures (Scheme 1). Facile reactions
of phthalimides with alkylamines in the aminolysis of phthalimide and the de-cross-linking of polyimides cross-linked with diamines at a high temperature under vacuum have been reported in the literature. These precedents encouraged us to explore the diamide-imide equilibrium, first in a small-molecule model reaction and then used in dynamic covalent networks formed from dianhydrides and diamines. The dynamic nature and flow properties of the networks are studied as a function of temperature.

RESULTS AND DISCUSSION

Model Reaction. To study the temperature dependence of the diamide-imide equilibrium, model reactions were performed using N-octylphthalimide (I) and octylamine (II) in toluene-d8 (Figure 1a) between 25 and 100 °C. The progression of this reaction was monitored by 1H NMR and the concentrations of I, II, and the product N1,N2-dioctylphthalimide (III) were determined based on the integrals of the NMR peaks (Figure 1b). The reaction was considered to reach the equilibrium state once the concentrations of compounds remained constant over time. As shown in Figure 1c for a reaction temperature of 60 °C, the concentration of III gradually decreased over time, while the concentration of III increased; after around 250 h, the concentrations no longer changed. Similar experiments as those shown in Figure 1 were carried out at several different temperatures between 25 and 100 °C and after achieving equilibrium, the equilibrium constants, K, were determined as

$$K = \frac{[III]}{[I][II]}$$

and the results are summarized in Table 1.

| temp. (°C) | [I]eq (mol·L⁻¹) | [II]eq (mol·L⁻¹) | [III]eq (mol·L⁻¹) | K (L⁻¹·mol⁻¹) |
|------------|----------------|----------------|----------------|-------------|
| 25         | 0.0456         | 0.135          | 0.0122         | 1.98        |
| 60         | 0.0419         | 0.119          | 0.0084         | 1.68        |
| 80         | 0.0424         | 0.120          | 0.0079         | 1.55        |
| 90         | 0.0426         | 0.120          | 0.0077         | 1.51        |
| 100        | 0.0429         | 0.120          | 0.0074         | 1.43        |

It is clear from these data that with increasing temperature, the equilibrium shifts toward the imide. The temperature dependence of this diamide-imide equilibrium was then further quantified by plotting the Van’t Hoff curve (Figure 2) of the above model reaction. In toluene, the equilibrium constant K decreases with increasing temperature from 1.99 at 25 °C to 1.43 at 100 °C and shifts toward the imide, which is in line with the literature. The standard reaction enthalpy (ΔH°) and entropy (ΔS°) were determined from the slope and intercept of the plot and resulted in ΔH° = −3.7 kJ·mol⁻¹ and ΔS° = −6.6 J·mol⁻¹·K⁻¹.

Synthesis of Dynamic Polyamide Networks. Based on the diamide-imide exchange reaction, a catalyst-free amide-imide dynamic covalent network is designed and synthesized in three steps as shown in Scheme 2. In the first step, stoichiometric amounts of poly(tetrahydrofuran)-diamine (PTHF-diamine) and pyromellitic dianhydride (PDMA) were reacted in anhydrous tetrahydrofuran (THF) for 30 min at ambient temperature to form poly(amic acid). The FTIR spectrum of the product after removal of THF (see Supporting Information (SI) Figure S7) confirmed full conversion of anhydride to amic acid groups based on the disappearance of the anhydride bands at 1854 and 1775 cm⁻¹. Subsequently, poly(amic acid) was dissolved in THF and refluxed at 66 °C overnight for imidization. After removal of the solvent, imidization was then completed by heating at 80 °C in vacuo for 15 h. After drying, Fourier transform infrared (FTIR) spectroscopy was performed to verify formation of imide from the appearance of bands at 1780 and 1725 cm⁻¹ (see SI Figure S7) and further confirmed by 1H NMR spectra of poly(amic acid) and polyimide (see SI Figure S2). The number-average molar mass, Mₐ, of the linear polyimide was determined with size-exclusion chromatography in THF (see SI Figure S8) to be 20 kg·mol⁻¹ relative to polystyrene standards. To this linear polyamide, various amounts of PTHF-diamine were added as cross-linker to form polyamide-imide networks, denoted as PTHF-PMDA-X, where X represents the combined feeding ratio of diamine to dianhydride, with X = 1 being the linear polyamide and X = 2 being the network cross-linked with one additional equivalent of PTHF-diamine with respect to PMDA. Gel fractions of 84, 82, and 69% were determined with a hydrogen bond-breaking solvent mixture MeOH/THF for PTHF-PMDA-2, PTHF-PMDA-1.75, and PTHF-PMDA-1.5, respectively (see the SI for details).

Amide–Imide Exchange in the Network. The diamide-imide equilibrium in the PTHF-PMDA-2 network was investigated with variable-temperature infrared (VT-IR) spectroscopy by monitoring the intensities of the diamide and imide bands over a range of temperatures from 30 to 155 °C at a heating rate of 1 °C·min⁻¹. The IR spectra in Figure 3a show the carbonyl bands of the imide at 1780 and 1725 cm⁻¹, and of the diamide at 1630 and 1550 cm⁻¹. The position of the diamide bands, as well as that of the N–H stretch band at 3310 cm⁻¹ (see SI Figure S9), indicates that the diamides are hydrogen bonded, probably leading to aggregation of the tetraamides in columnar stacks similar in structure to those found in tetraamide columnar liquid crystals or benzene-triamide thermoplastic elastomers. As shown in Figure 3b, the intensities of the diamide bands decrease, while for those of imide the bands increase upon heating. Above 140 °C, the bands corresponding to the diamide groups almost disappear, indicating the dissociation of the network. Similarly, the...
reformation of the diamide bonds was studied by cooling the sample (see SI Figure S9). Analysis of the band intensities as a function of temperature gave a $K$ value that decreased from $\sim 5$ M$^{-1}$ at 115 °C to 0.8 at 155 °C (see SITable S1).

(Thermo)Mechanical and Rheological Properties. Thermal properties of the networks were first studied by thermogravimetric analysis (TGA) and differential scanning calorimetry (DSC). Thermograms are shown in Figure 4.

From the TGA results (Figure 4a), it can be concluded that the networks are stable up to 200 °C and have a degradation temperature $T_{d,5\%} \geq 300$ °C. Furthermore, the networks contain (probably PTHF) crystalline regions below 0 °C as can be concluded from the cold crystallization and melting peaks in the DSC thermograms (Figure 4b).

To further characterize the (thermo)mechanical properties of the networks, dog-bone-shaped samples were prepared using a hot press at 125 °C for 30 min under a pressure of 100 bar and studied with tensile testing and dynamic mechanical thermal analysis (DMTA). The results of these studies are shown in Figure 5.

Tensile tests were carried out in triplicate and representative stress–strain curves at room temperature are shown in Figure 5a (see the SI for more details). From these measurements, it
is clear that the Young’s modulus increases with increasing amount of cross-linking diamine in PTHF-PMDA-X: $E \approx 1$ MPa for $X = 1.5$, $E \approx 2$ MPa for $X = 1.75$, and $E \approx 3$ MPa for $X = 2$. Furthermore, the material with the lowest amount of cross-linker, PTHF-PMDA-1.5, has a significantly higher elongation at break than the other two materials. Qualitatively, these results are expected as a higher cross-link density leads to stiffer materials. This is further confirmed by the DMTA results shown in Figure 5b, where the increasing storage modulus of the rubbery plateau with increasing $X$ is directly related to an increasing cross-link density. It is interesting to note that only the least cross-linked network showed a melting transition between $-60$ and $10^\circ$C, while PTHF-PMDA-1.75 and PTHF-PMDA-2 remain amorphous (see Figure 4b). The dog-bone-shaped samples after compression molding were all transparent; hence, the crystallinity of the PTHF-PMDA-1.5 network might stem from the cold crystallization of pendant (and/or free) PTHF chains during the conditioning step prior to DMTA measurements as shown in Figure 5b. At temperatures above $120^\circ$C, the storage modulus drops precipitously, indicative of the transition to viscous flow of the materials. Furthermore, there is a shift to higher temperatures for this transition with increasing cross-link density. It is clear that this viscous flow is related to the dynamic character of the dynamic covalent networks. We hypothesize that the abrupt transition is the combined (and possibly cooperative) effect of a shift in the diamide/imide equilibrium and dissociation of tetraamide stacks. Since the nature of the exchange reaction is dissociative, the observed flow may be accompanied by a greater (or even complete) disintegration of the network.

The dynamic nature of the polyamide-imide networks was further investigated using shear rheology with disk-shaped samples of 8 mm diameter and 1 mm thickness. The frequency-dependent shear moduli at varying temperature of the PTHF-PMDA-2 network are presented in Figure 6a. The significantly decreasing values for $G'$ at higher frequencies for $T > 120^\circ$C are consistent with an increased, but not a complete, disintegration of the network at higher temperatures; if disintegration of the network were complete, a stronger frequency dependence of the moduli would be expected, as well as $G'' > G'$ at high frequencies. The oscillatory frequency sweep measurements further reveal the transient nature of the PTHF-PMDA network, evident by the significant drop of $G'$ at lower frequencies ($<10^{-2}$ rad s$^{-1}$) for temperatures above and including $120^\circ$C. At $130^\circ$C, PTHF-PMDA-2 shows a crossover of $G'$ and $G''$ at a frequency of $0.02$ rad s$^{-1}$. Similar results were obtained for PTHF-PMDA-1.5 and PTHF-PMDA-1.75 (see the SI for more details).

Stress relaxation measurements of the PTHF-PMDA-2 network at temperatures between 80 and $130^\circ$C are shown in Figure 6b. As the temperature decreases, the relaxation curves shift to longer time scales as the exchange reaction slows down and the equilibrium shifts toward diamide from imide, resulting in a higher cross-link density and less network mobility. The relaxation behavior cannot be adequately described by a Maxwell model (i.e., a single exponential decay), which describes relaxation in many of the reported dynamic covalent networks quite well.$^{37-39}$ In a number of studies,$^{40-43}$ a stretched exponential decay (eq 2) was used to describe the relaxation of the stress $\sigma(t)$, where the exponent $\beta$ is often considered as a factor taking into account the
dispersity in a particular relaxation mode, with characteristic relaxation time $\tau_{fi}$

$$
\sigma(t) = \sigma_0 \exp\left\{ -\left(\frac{t}{\tau}\right)^{\beta} \right\}
$$

(2)

Although it is not possible to fit the relaxation data using this model over the entire range of time, we nevertheless decided to use this relatively simple model to numerically characterize the stress relaxation behavior. Since our primary interest lies in network relaxation caused by transamidation characterized by longer relaxation times, data were fitted for $t \geq 10$ s (see SI for more details). The results of curve fitting using eq 2 are shown in Figure 6b and the best-fit values of each parameter are summarized in Table 2. Adequate fits are obtained at longer times, but at most temperatures relaxation below $t = 10$ s is strongly underestimated, indicating that separate, faster relaxation modes are important at short time scales. These modes are more important at lower temperatures and are responsible for the aberrant fit results at 80 °C, with an unrealistically low value of 0.39 s for $\tau$ in PTHF-PMDA-2. At this temperature, a large contribution of the faster modes around $t \approx 10$ s overlaps strongly with relaxation through transamidation and obscures the dynamic covalent relaxation time. Above 80 °C, the data follow the trends expected for a thermally activated chemical exchange in combination with a shift in the diamide--imide equilibrium to lower the cross-link density. Fitted relaxation times and the values of $\sigma_0$ continuously decrease with increasing temperatures, and the values of fitted $\sigma_0$ are close to the stresses observed in oscillatory shear rheology at a strain of 1% and $\omega = 6.28$ rad·s$^{-1}$ (see SI Figure S23). The very low values for $\beta$, however, indicate a very high dispersity of relaxation times and suggests that relaxation at longer time scales is also a complex process that may involve dissociation of hydrogen-bonded aggregates in addition to the contribution from transamidation. A more detailed calculation of the retardation spectra is nontrivial, and is beyond the scope of the current study.

The creep response of the materials was investigated by applying a constant shear stress of 10 kPa to the material for 1200 s at varying temperatures and monitoring the strain as a function of time. The results for PTHF-PMDA-2 are shown in Figure 7a (for the other two systems, see the SI). In all systems, below 110 °C, less than 3% strain is observed after the initial response, more than 90% of which is recovered after the release of stress. Above 110 °C, the strain linearly increases with time after the initial response and the recovery drops to 29% of the deformation (see SI Figure S18 for creep recovery experiments). To compare the three materials better, the strain rates were determined from the strain—time curves for 200 s onward and are shown as a function of temperature in Figure 7b. The strain rates for all three networks increase drastically above 110 °C with a higher temperature dependence, evident from the steeper slope in Figure 7b. The temperature at which these changes occur corresponds to the transition to viscous flows as observed in the DMTA thermograms (Figure 5b).

Reprocessibility and Reversibility of Dynamic Poly(amide–imide) Networks. To confirm the reprocessibility of the dynamic poly(amide–imide) networks, the PTHF-PMDA-2 samples were shredded and compression molded for three cycles. The stress—strain curves of the samples after each reprocessing cycle presented in Figure 8a show there is neither a significant loss in the Young’s modulus nor in the elongation at break.

Furthermore, the reversibility of the network was studied using oscillatory shear rheology to heat and cool the sample for six cycles at a rate of 1 °C·min$^{-1}$ and a strain of 1% and $\omega = 6.28$ rad·s$^{-1}$. As can be seen from Figure 8b, the temperature dependences of the moduli remain the same, and despite some hysteresis in the reformation of the network upon cooling, the network fully restores if given sufficient time, allowing processing or reprocessing of the network without significant loss of mechanical properties.

| network | $X = 2$ | $X = 1.75$ | $X = 1.5$ |
|---------|---------|------------|------------|
| temp. (°C) | $\sigma_0$ (kPa) | $\beta$ | $\tau$ (s) | $\sigma_0$ (kPa) | $\beta$ | $\tau$ (s) | $\sigma_0$ (kPa) | $\beta$ | $\tau$ (s) |
| 80      | 29      | 0.06      | 0.39      | 10        | 0.25      | 1.3 × 10^4 | 9.5        | 0.37      | 1.5 × 10^4 |
| 90      | 13      | 0.13      | 2.1 × 10^4 | 9.6      | 0.27      | 9.3 × 10^1 | 9.4        | 0.39      | 9.7 × 10^3 |
| 100     | 12      | 0.13      | 1.8 × 10^4 | 8.5      | 0.29      | 4.5 × 10^1 | 8.3        | 0.43      | 5.0 × 10^3 |
| 110     | 7.7     | 0.27      | 1.3 × 10^4 | 6.6      | 0.34      | 2.0 × 10^1 | 6.1        | 0.48      | 2.1 × 10^3 |
| 120     | 4.8     | 0.36      | 1.9 × 10^3 | 4.0      | 0.44      | 5.2 × 10^1 | 3.3        | 0.57      | 4.4 × 10^2 |
| 130     | 4.4     | 0.31      | 98        | 1.8      | 0.44      | 1.3 × 10^1 | 1.8        | 0.58      | 1.1 × 10^2 |

Figure 7. Shear rheology of PTHF-PMDA-X networks at 80, 90, 100, 110, 120, and 130 °C. (a) Creep results for PTHF-PMDA-2 at a constant stress of 10^3 Pa. (b) Comparison of strain rates (derived from the creep data acquired in (a) using $t > 200$ s) for different PTHF-PMDA-X networks (where $X = 2, 1.75, 1.5$).
Figure 8. (a) Stress–strain curves of the PTHF-PMDA-2 network with three cycles of reprocessing by shredding and compression molding the samples acquired by tensile testing at a strain rate of 30 mm·min⁻¹ at room temperature. (b) Reversibility of the network dissociation/reforption was studied using shear rheology for six cycles of heating and cooling from 80 to 140 °C at a rate of 1 °C·min⁻¹ (strain of 1% and ω = 6.28 rad·s⁻¹).

■ CONCLUSIONS
A novel type of dissociative dynamic covalent network has been synthesized by making use of the dynamic equilibrium between diamides and imides derived from pyromellitic dianhydride. Above 80 °C, the networks show significant stress relaxation due to chemical exchange of amides via imide intermediates. The temperature dependence of the diamide–imide equilibrium leads to a significant, but fully reversible reduction of cross-link density in the network above 110 °C. These features result in a moldable material with a strong drop in viscosity at high temperatures that aids its (re)processing. Probably as a result of extensive hydrogen bonding and aggregation of the tetraamide, viscosity loss is limited and material integrity is retained at intermediate temperatures.

We expect that the amide–imide chemistry demonstrated in this paper will find use in applications where temporarily reaching low viscosities is important, e.g., improving interfacial adhesion in additive manufacturing techniques such as fused filament fabrication.

■ ASSOCIATED CONTENT

Supporting Information
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.macromol.1c01389.

Experimental details, characterization methods, and additional results of NMR studies, DSC, TGA, and rheology supporting data (PDF)

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