Abstract

The composition of low-$T_g$ $n$-butylacrylate-$\text{block-(acetoxyaceto)ethyl}$ acrylate block polymers is investigated as a strategy to tune the properties of dynamically cross-linked vinylogous urethane vitrimers. As the proportion of the cross-linkable block is increased, the thermorheological properties, structure, and stress relaxation evolve in ways that cannot be explained by increasing cross-link density alone. Evidence is presented that network connectivity defects such as loops and dangling ends are increased by microphase separation. The thermomechanical and viscoelastic properties of block
copolymer-derived vitrimers arise from the subtle interplay of microphase separation and network defects.

**Introduction**

Block copolymers that combine rubbery and glassy segments self-assemble into nanostructured materials with emergent properties not seen in the individual homopolymers. These properties are highly tunable by adjusting the volume fraction of each block and their architecture.\(^1\) When the low-$T_g$ segment is the majority block, microphase separation produces a rubbery matrix physically cross-linked by spherical glassy domains. Thermoplastic elastomers (TPEs), the most commercially successful example of this phenomenon, combine the processability of thermoplastics and the elasticity of chemically cross-linked rubbers.\(^2\) As the volume fraction of the glassy block is increased, high-impact thermoplastics with bicontinuous morphologies are formed.\(^3\) Above the $T_g$ of the glassy block, these block copolymers behave as viscous liquids, enabling extrusion and injection molding of virgin and recycled material; however, this transition also defines the upper operating temperature of the material. Under stress, chain pull-out from the hard domains can also occur, ultimately leading to failure. To replace or reinforce these physical cross-links without sacrificing processability, researchers have explored various supramolecular interactions, including H-bonds,\(^4,5\) metal-ligand interactions,\(^6,7,8\) $\pi-\pi$ stacking,\(^9\) and ionic interactions.\(^10,11\)

Here, we explore the incorporation of dynamic covalent bonds in rubbery diblock copolymers, varying their composition to create vitrimers with highly tunable properties. Vitrimers are an emerging class of covalent adaptable networks (CANs)\(^12\) that also bridge thermoplastics and thermosetting polymers through associative exchange reactions.\(^13\) Unlike TPEs, supramolecular networks, and many dissociative CANs, vitrimers do not experience a viscosity drop at elevated temperatures because
reconfiguration of the network topology occurs without loss of covalent connectivity. Furthermore, the associative mechanism is thought to provide higher solvent resistance, since many vitrimers can swell in good solvents without dissolving. Following pioneering work by Bowman and Leibler, many researchers have adapted various associative exchange reactions for use in vitrimers. These efforts have focused on controlling vitrimer flow through applying new reactions, catalyzing bond exchange, or modifying reactivity of existing mechanisms. Macromolecular structure and assembly might also be harnessed to tune vitrimer properties. For example, cross-link density, chain mobility, and phase separation have all been investigated in conjunction with vitrimers.

While the first generation of vitrimers focused on high-$T_g$ thermoset-like materials, our group and others have become interested in the challenge of designing catalyst-free elastomeric vitrimers. With high-$T_g$ materials, even if the small-molecule exchange reaction can occur at low temperatures, it will not proceed appreciably within the network until $T_g$ is exceeded. For low-$T_g$ materials, however, the associative exchange reaction must be carefully considered to enable reprocessing at reasonable temperatures but avoid creep under service conditions. Additionally, the identity and structure of the rubbery polymer matrix can influence the mechanism of the exchange reaction and the rate of flow. The majority of elastomeric vitrimers have used telechelic or randomly grafted prepolymer, resulting in vitrimers with a relatively uniform distribution of the exchangeable groups. Segregating the exchangeable groups into one segment of a block copolymer provides new opportunities to tune the properties of the resulting vitrimer while keeping the chemical identity of the components constant. In particular, self-assembly of the vitrimer by microphase separation into cross-link-rich and cross-link-poor domains can be programmed by the structure of the block prepolymer to dramatically influence the vitrimer’s thermomechanical properties. We envisioned that minority blocks bearing exchangeable groups could serve an analogous role to the glassy domains of TPEs, while conferring the superior solvent and temperature resistance common to vitrimers.
During the course of our study, Sumerlin, Epps, and co-workers reported a glassy vitrimer derived from the symmetric block copolymer poly(butyl methacrylate)-block-poly(2-(acetoxyaceto)ethyl methacrylate), or BMA-\textit{b}-AAEMA, wherein dynamic cross-linking occurs solely in the AAEMA block using the vinylogous urethane dynamic cross-link developed first by Du Prez and co-workers. They attributed differences in the viscoelastic properties, relative to a statistical copolymer-derived vitrimer, to the presence of phase separation. This initial contribution focused on thin-film structure and suggested the potential of block copolymer-derived vitrimers by comparing their high-temperature stress relaxation and creep with those of a statistical copolymer-derived vitrimer.

To build a more complete picture of the design space surrounding block copolymers as vitrimer precursors, we vary the composition of a low-\(T_g\) diblock copolymer to study the effect of weight fraction of the cross-linking domain and study a poly(\(n\)-butylacrylate)-\textit{block}-poly(2-(acetoxyaceto)ethyl acrylate) (PnBA-\textit{b}-PAAEA) system. We provide the full details of vitrimer synthesis, thermomechanical characterization, stress relaxation, and microphase separated structure. Using dynamic light scattering (DLS) and diffusion-ordered NMR spectroscopy (DOSY NMR), we show that under the influence of dynamic cross-linking, block copolymers evolve from hyperbranched polymers to infinite networks. Dynamic mechanical thermal analysis (DMTA) and stress relaxation measurements reveal that rubbery plateau modulus and flow rates do not increase as expected when cross-linker is increased. We employ small-angle x-ray scattering (SAXS) to assess the microstructure of the prepolymer and vitrimers. By studying this series of vitrimers using this broad suite of techniques, we provide experimental comparisons to recent theories describing vitrimer stress relaxation and elucidate general strategies for tuning vitrimer properties with block copolymer composition.

**Results**
**Synthesis of prepolymer**s. We synthesized diblock polymers of \( n \)-butyl acrylate (nBA) and a ketoester-containing acrylate monomer ((2-acetoacetoxy)ethyl acrylate, AAEA)\(^{25} \) using reversible addition-fragmentation chain-transfer (RAFT) polymerization\(^{26} \). Because more common end-group removal protocols\(^{27} \) were unsuccessful for this system, the trithiocarbonate chain ends were removed by reductive photolysis\(^{28} \). We prepared five prepolymer\s with increasing mole fraction of AAEA, \textbf{Block-13}, \textbf{Block-24}, \textbf{Block-34}, \textbf{Block-48}, and \textbf{Block-66}, in which the number refers to the weight\% functional monomer present in the prepolymer, as determined by \(^1\text{H} \) NMR (see SI for details). The length of each block was systematically varied to adjust the morphologies and cross-linking densities of the prepolymers and vitrimers, including minority functional block\s (\textbf{Block-13}, \textbf{Block-24}, \textbf{Block-34}), a symmetrical diblock (\textbf{Block-48}), and majority functional block (\textbf{Block-66}). \textbf{Table 1} shows that the M\(_n\) of the copolymers fell within a narrow range, from 8 to 11 kg/mol, with low D (1.07 to 1.28). We targeted molecular weights well below the entanglement molecular weight of poly(nBA), 29 kg/mol, to avoid mechanical effects arising from backbone entanglement\(^{29} \). All of the prepolymer\s were viscous liquids. Differential scanning calorimetry (DSC) of these diblock copolymers revealed two \( T_g \)'s below room temperature, consistent with microphase separation between PnBA (\( T_g \approx 54 \, ^\circ\text{C} \)) and PAEEA-rich domains (\( T_g \approx 22 \, ^\circ\text{C} \), \textbf{Figure 1}).
Scheme 1. Synthesis of block copolymers used in this study.

Table 1. Characteristics of PnBA-b-PAAEA prepolymers

|               | Block-13 | Block-24 | Block-34 | Block-48 | Block-66 |
|---------------|----------|----------|----------|----------|----------|
| mol% AAEA     | 9        | 17       | 25       | 37       | 55       |
| Mₘ (kg/mol) ᵃ | 8.0      | 8.3      | 9.3      | 11.0     | 9.2      |
| D ᵇ           | 1.13     | 1.18     | 1.28     | 1.19     | 1.07     |
| Tₘ¹ (°C) ᵇ    | –50      | –50      | –45      | –46      | –55      |
| Tₘ² (°C) ᵇ    | –37      | –35      | –32      | –28      | –35      |

ᵃ Determined by gel permeation chromatography relative to PS standards. ᵇ Determined by differential scanning calorimetry (second heating cycle).
**Figure 1:** DSC data of all PnBA-\(b\)-PAAEA prepolymer samples and PAAEA homopolymer with the first heating cycle as a solid line and the second heating cycle as a dotted line (heating rate of 5°C/min).

**Cross-linking prepolymer to form vitrimer.** To form the vitrimer (denoted with suffix -Net), we chose tris(2-aminoethyl)amine (TREN) as the cross-linker, and the theoretical concentration of free primary amine in all networks was 33 mol% excess relative to ketoester functional groups on the polymer (Scheme 2). This stoichiometry was previously shown by Sumerlin and coworkers to enable reprocessing on reasonable time scales for poly(methacrylate) vitrimers cross-linked by TREN.\(^{26}\)

**Scheme 2.** Cross-linking of the block copolymers by vinylogous urethanes.

(a) Cross-linking by TREN

(b) Vinylogous urethane exchange

Surprisingly, after adding the cross-linker to the block polymer solutions (170 mg/mL polymer), we did not observe gelation even after standing in a sealed container overnight. As block-selective covalent cross-linking is a common strategy to stabilize block polymer micelles,\(^{30}\) we suspected that
the formation of cross-linked aggregates in solution was occurring instead of network formation. To further understand this phenomenon, we turned to dynamic light scattering (DLS) and diffusion-ordered NMR spectroscopy (DOSY NMR).

DLS of Block-34 solutions in THF show a peak centered at ~2 nm consistent with unimers and suggesting that the block prepolymer by themselves do not aggregate (Figure 2a). Upon adding TREN, cross-linked Block-34 nanoparticles form quantitatively within minutes. These 10-nm particles do not further aggregate even after 18 hours. These results suggest that ketoester-amine condensation occurs readily, but the prepolymer sequence prevents an infinite network from percolating through the sample at this concentration. Additionally, 1H NMR after addition of TREN in THF-d8 shows broadening of the signals corresponding to the ethyl linker at 4.3 ppm, and appearance of broad signals at 2.6 and 9 ppm, consistent with successful condensation. The diffusion coefficient of Block-34 in THF-d8 is 2.0 x 10⁻⁶ cm²/s; after addition of TREN, the diffusion coefficient decreases, indicating the formation of hyperbranched cross-linked polymers with larger hydrodynamic radii. After a week at 25 °C, gelation is not observed and the diffusion coefficient of the condensed species is 9.6 x 10⁻⁷ cm²/s (Figure 2b).
Figure 2. (a) DLS data for solutions of cross-linked Block-34. Adding TREN to a 170 mg/mL THF solution of Block-34 shows that the formation of nanoscale aggregates is relatively fast. Nanoparticles do not change in size after 18 hours. (b) DOSY NMR of Block-34 in THF-\textit{d}$_8$ (10 mg/mL) before and after the addition of TREN.

If a network is to form, the cross-link dynamism must allow for connectivity changes and eventual particle merging, leading toward a critical gel concentration. Indeed, we obtained vitrimers as transparent, pale yellow films after solvent removal and annealing using the following sequence. First, the block copolymer solutions were reacted with TREN, and the solutions were left capped for at least 2 hours to ensure complete conversion of all ketoester moieties.$^{31}$ After briefly degassing the solutions using vacuum and a sonicator, we poured the solutions into a mold. The vitrimers were then obtained after slow evaporation of the THF solvent and annealing at 55 °C for 18 hours. A
representative photograph of Block-66 is shown in Figure 3. All the vitrimers in this study had gel fractions (see SI) of 94% or greater (Table 2), with the exception of Block-13, which remained a tacky, viscoelastic solid at room temperature even after removal of solvent and annealing (Figures S-16 and S-17). The swelling ratios decrease as the amount of functional block increases, as expected based on the increase in sites capable of forming cross-links.

![Northwestern](image)

**Figure 3.** Photograph of Block-66. Each sample is approximately 20 x 6.5 x 0.5 mm.

| Table 2. Properties of vitrimers |
|----------------------------------|
|                                 |
| **wt% TREN** | Block-24-Net | Block-34-Net | Block-48-Net | Block-66-Net |
| Swelling in THF, 55 °C | Gel Fraction (%) | 97 | 94 | 98 | >99 |
| Swelling Ratio (%) | 259 | 181 | 128 | 60 |
| Tg1 (°C) | –26 | –27 | – | – |
| Tg2 (°C) | 96 | 79 | 95 | 99 |
| E' (MPa) | 2.2 | 1.2 | 6.0 | 30.9 |
| τ* (s) | 190 °C | 49 ± 15 | 10 ± 4 | 12 ± 3 | 11 ± 1 |
| 180 °C | 110 ± 55 | 20 ± 11 | 18 ± 2 | 18 ± 3 |
| 170 °C | 197 ± 85 | 40 ± 18 | 41 ± 9 | 33 ± 5 |
| 160 °C | 408 ± 210 | 74 ± 36 | 73 ± 7 | 62 ± 12 |
| Activation Parameters | Ea (kJ/mol) | 116 ± 5 | 112 ± 4 | 112 ± 16 | 97 ± 3 |
| ln(τ0) | –26.3 ± 1.4 | –26.8 ± 1.0 | –26.9 ± 4.2 | –22.8 ± 0.8 |

*a Determination of gel fraction described in SI. b Temperature at the peak of tan(δ). c Rubbery plateau storage modulus from DMTA at 130 °C. d Each point is average of three runs. Uncertainty is the Student’s T-test. e Uncertainties are the standard error of the slope and intercept of the Arrhenius plot.
Dynamic Mechanical Thermal Analysis. DMTA of the Block vitrimers, which we performed under an atmosphere of nitrogen, reveal a rich viscoelastic landscape (Figure 4). Block-24-Net and Block-34-Net both display a clear drop in their $E'$ traces that is accompanied by a peak in their tan($\delta$) traces at approximately –26 °C. This transition corresponds to the $T_g$ of PnBA. As the nBA fraction of the prepolymer decreases in Block-48 and Block-66, the tan($\delta$) peak corresponding to its $T_g$ disappears. Above –26 °C, $E'$ drops gradually for all samples; this behavior is reminiscent of a thermoplastic and is consistent with the presence of a PnBA domain. This gradual decrease is also similar to DMTA behavior observed by Coates, Hillmyer and co-workers for imine CANs derived from gradient copolymers. In the block copolymer vitrimers, however, the $E'$ and tan($\delta$) traces display a second, high-temperature transition, corresponding to the $T_g$ of the PAEAA domain. Unlike a thermoplastic elastomer, the $E'$ enters a rubbery plateau instead of terminal flow. The area under this second transition in the tan($\delta$) trace increases as the volume fraction of the functional block increases. Additionally, the glass transition generally broadens as the cross-linking domain increases in volume.

Figure 4. DMTA data: $E'$ (solid lines) and tan($\delta$) (dashed lines) (heating rate of 3 °C/min).
Curiously, **Block-24-Net** has a higher rubbery plateau $E'$ than **Block-34-Net**, despite the greater fraction of functional monomers in **Block-34**. A further increase of the functional block in **Block-58-Net** and **Block-66-Net**, however, increases the rubbery plateau modulus as expected. In contrast, the swelling ratios decrease monotonically as the %AAEA is increased, consistent with increasing cross-link density. These data suggest that the cross-link densities of the vitrimers are different in the dry and swollen states, and will be examined further in the Discussion.

**Stress Relaxation.** CAN recyclability is attributed to the presence of dynamic bonds, which facilitate the rearrangement of network topology to bridge the surfaces of smaller fragments and macroscopically re-form a continuous material. Stress relaxation in vitrimers is considered a fingerprint for recyclability and offers quantitative insight into the timescale required for bond swapping and subsequent topology rearrangement. We performed stress relaxation experiments under an atmosphere of nitrogen at temperatures ranging from 160–190 °C. We chose this range to avoid any complexities related to the glass transition. The inhomogeneous structure of the **Block** vitrimers suggests that they would display multiple stress relaxation modes, deviating from ideal Maxwellian behavior. In the context of vitrimers, Torkelson$^{33}$ and others have estimated that a stretched exponential (eq 1) can capture these deviations:

$$\frac{E}{E_0} = \exp \left( - \left( \frac{t}{\tau^*} \right)^B \right)$$  \hspace{1cm} (eq 1)

$E/E_0$ is the normalized relaxation modulus, $t$ is the time in seconds, $\tau^*$ is the characteristic relaxation time, and $\beta$ is a fitting factor that is a proxy for the homogeneity of the relaxation process. In the upper limiting case ($\beta = 1$), the relaxation is that of a single Maxwell element (i.e., the material possesses a single relaxation mode), while the lower limiting case ($\beta \sim 0$) represents multiple relaxation modes. In the case of the block-copolymer-derived vitrimers, $\beta$ values lie between 0.41 and 0.62,
suggesting that the materials are indeed inhomogeneous and cannot be accurately represented by a single Maxwell element.

**Figure 5.** (a) Representative stress relaxation data at 160 °C. (b) Arrhenius plots for stress relaxation in the range of 160–190 °C. (c) Full stress relaxation profile for Block-48-Net at 190°C. (d) Continuous relaxation spectrum for (c).

Flow activation energy $E_a$ may be estimated by fitting $\tau^*$ data to an Arrhenius-like equation (ESI, Eq S-2). Interestingly, we do not observe a trend in $\tau^*$ or $E_a$ as the nominal cross-link density (i.e. AEAA fraction) is increased. The vitrimer with the lowest AEAA fraction, **Block-24-Net**, exhibits significantly longer relaxation times. The relaxation times for **Block-34-Net**, **Block-48-Net**, and **Block-66-Net**, in contrast, are nearly identical at 160 °C (Figure 5a). In contrast, the $E_a$ values of
Block-24-Net, Block-34-Net, Block-48-Net are in fact quite similar (112–116 kJ/mol, within error), while that of Block-66-Net is slightly lower (97 kJ/mol) (Figure 5b, Table 2).

A “full” relaxation profile of a Block vitrimer shows that a limited amount of stress is retained at long time scales, even at 190 °C (Figure 5c). To further support the notion of inhomogeneity in the relaxation process, we obtained the continuous relaxation spectrum of Block-48-Net at 190 °C (Figure 5d; see SI for details). We found that multiple relaxation modes emerged, including one at extremely long timescale. These data are consistent with creep measurements performed by Sumerlin, Epps, and co-workers, who demonstrated that a symmetrical block copolymer-derived vitrimer was able to resist creep at high temperature, relative to a statistical copolymer-derived vitrimer with identical composition. Similarly, we found that a control vitrimer derived from a statistical copolymer displays a single relaxation mode and relaxes all stress within 200 s at 190 °C (Figure S-22).

**Small-Angle X-ray Scattering.** SAXS data confirm microphase separation in Block prepolymer and their resulting vitrimers (Figure 6, solid lines). Block-24 and Block-34 have SAXS patterns consistent with hexagonally-packed cylinders with principal domain spacing \( d = (2\pi/q^*) = 11.9 \) nm and 11.3 nm, respectively, where \( q^* \) is the first scattering peak, while Block-48 and Block-66 have SAXS patterns consistent with lamellae with principal domain spacing 14.3 and 11.1 nm (Table 3). Block-13, which did not form a network upon addition of TREN, displays a single broad peak by SAXS despite exhibiting two \( T_g \)’s by DSC. Variable-temperature SAXS data reveal that the order-disorder transition temperature \( (T_{ODT}) \) follows the trend Block-13 \(<\) Block-24 \(<\) Block-34 \(\sim\) Block-66 \(<\) Block-48 (Figure S-25). The diblock Block-48 did not exhibit an ODT up to 215 °C, consistent with its symmetrical structure and slightly higher molecular weight.
**Figure 6.** SAXS data for Block prepolymerms and vitrimers. Curves have been shifted vertically for clarity. Data for Block-24 and Block-34 and the corresponding vitrimers were acquired on a lab-based (Ganesha) instrument, while other data were acquired using synchrotron radiation; see SI for details.

**Table 3. Morphological characteristics of block prepolymerms and vitrimers**

|                  | Block-13 | Block-24 | Block-34 | Block-48 | Block-66 |
|------------------|----------|----------|----------|----------|----------|
| **Before cross-linking** | morphology | DIS | HEX | HEX | LAM | LAM |
| $T_{ODT}$ (°C) $^b$ | - | 115 | 135 | >215 | 145 |
| $d$ (nm) $^c$ | 11.6 | 11.9 | 11.3 | 14.1 | 11.0 |
| **After cross-linking** | $d$ (nm) $^c$ | 10.3 | 11.5 | 11.1 | 12.9 | 12.9 |
| $\Delta d$ (%) $^d$ | -11.6 | -3.4 | -1.8 | -8.1 | 17 |

$^a$ This sample did not form an elastomeric network upon addition of TREN. $^b$ Lower-bound estimate determined by SAXS from the transition in 1/intensity vs. 1/T (see Figure S-26). $^c$ Determined by SAXS from the first scattering peak, $d = (2\pi/q^*).$ $^d$ Change in $d$ value before and after cross-linking with TREN.

All vitrimers, in contrast, exhibit a single broad peak by SAXS, suggesting disorganized but microphase-separated morphologies (Figure 6, dashed lines). Annealing at higher temperatures or for longer times did not improve the organization of the vitrimers (Figure S-24). This disorganization is consistent with what Sumerlin, Epps, and co-workers observed in bulk films obtained by solution-casting their block copolymer-derived vitrimer. All of the samples with PAEAA as the minority or symmetrical block, Block-13 through Block-48, displayed a decrease in the $d$ value after cross-linking, while Block-66 displayed an increased principal domain spacing after cross-linking.

Clearly, introduction of TREN disrupts long-range ordering of the block copolymer microdomains. We hypothesized that this loss of order could arise from thermodynamic origins if the segregation strength of PnBA-$b$-PAEAA is lowered by formation of the vinylogous urethanes, or by the dramatic increase in dispersity that accompanies branching as cross-links form. $^{34}$ Alternatively, disorganization could be ascribed to kinetics if associative cross-linking in the disordered solution state creates barriers to formation of the ordered morphologies. To probe these possibilities, we condensed $N,N-$
dimethylethylenediamine onto the representative prepolymers Block-34 and Block-66 (Scheme 3). This condensation with a primary amine simulates the chemistry of the functional block after cross-linking but denies the polymers the ability to form a network. In the resulting linear diblock copolymers, Block-34-Amine and Block-66-Amine, we also observed the loss of order by SANS (Figure 6). These results suggest that the change in chemistry from ketoester to vinylogous urethane side chains reduces the magnitude of the Flory–Huggins interaction parameter, $\chi$, with the PnBA blocks. Attractive interactions between the carboxylate end group (derived from the RAFT chain-transfer agent) and the TREN amines may also compatibilize the blocks.

Scheme 3. Condensation to form linear vinylogous urethane diblock copolymers.

To increase the segregation strength of the cross-linked block copolymer, we next crosslinked Block-48, the prepolymer with the highest $T_{ODT}$, with a polar diamine, 4,7,10-trioxa-1,13-tridecanediamine. We rationalized that this polar linker might disfavor mixing of the crosslinked PAAEA block with the nonpolar PnBA block, and serve as a competitive H-bond acceptor to disrupt any favorable interactions between the vinylogous urethanes and PnBA esters. The resulting annealed vitrimer (Block-48-Trioxa-Net) was cloudy, suggesting macrophase separation (Figure S-15).
Nonetheless, the SAXS pattern in Figure 6 again exhibited a single peak, which was narrower than that of Block-48-Net but still devoid of higher-order scattering peaks, consistent with a disorganized morphology. Therefore, simply increasing the polarity of the cross-linker is insufficient to drive organization in block copolymer vitrimers.

**Dissolving a vitrimer.** The vitrimers were formed by removing solvent from a solution of cross-linked nanoparticles, an example of architectural transformation from a hyperbranched polymer to an infinite network.\(^{35}\) The persistence of the block prepolymer-derived nanoparticles in solution at room temperature suggests that they are thermodynamically stable under these conditions. In principle, therefore, it should be possible to perform the reverse architectural transformation, from an infinite network to soluble hyperbranched polymers, simply by diluting the vitrimer. Recently, Nicolaÿ,\(^{36}\) Avérous,\(^{17b}\) and Smulders\(^{37}\) have shown that boronic ester- and imine-based vitrimers can be readily dissolved under mild conditions. While these chemistries are susceptible to hydrolysis from adventitious water, Avérous and Smulders obtained \(^1\)H NMR spectra of the dissolved vitrimers showing that imines were retained. These observations are in stark contrast to the prevailing wisdom that vitrimers cannot be dissolved because of the associative exchange mechanism.

As our gel fraction data (Table 2, 94–99%) demonstrate, the block copolymer-derived vinylogous urethane vitrimers remain largely intact in THF at 55 °C, suggesting that there are kinetic barriers to dissolution. Therefore, we conducted experiments at higher temperature, with rigorous exclusion of oxygen and moisture to minimize side reactions that could introduce permanent cross-links\(^{38,39}\) or hydrolyze the network. Indeed, we observed that Block-24-Net swelled and fragmented in THF when heated in a sealed vessel to 110 °C over the course of 48 hours (Figure 7a,b). DLS confirms the presence of nanoscale particles in the sol, analogous to the hyperbranched polymers obtained during solution cross-linking (Figure 7c). Nevertheless, we did not observe quantitative degradation of the
vitrimer even under these forcing conditions, consistent with the associative cross-links enabling local bond rearrangement but resisting macroscopic remodeling. Additionally, slight coloration of the material suggests trace amine oxidation occurred under these forcing conditions despite our precautions.

Figure 7. (a,b) Photographs of Block-24-Net (labeled 1) and the same material “capped” with methyl acetoacetate (labeled 2) to prevent reconfiguration, before (a) and after (b) heating in THF at 110 ºC in sealed vessels. (c) DLS of the sol fractions recovered from the vials in (b).

To confirm that associative dynamic bond rearrangement to form higher-order loops was responsible for partial dissolution of the vitrimer rather than undesired hydrolysis or bond-breaking, we swelled Block-24-Net with methyl ketoacetate to consume the excess primary amines (Block-24-Net-Capped). Leibler has shown that this small-molecule reaction prevents further rearrangement of
the vinylogous urethanes, “freezing” the vitrimer topology.\textsuperscript{20c} Indeed, the capped vitrimer remained intact after heating in THF at 110 °C for 4 days (Figure 7b), and no polymer or particles were observed in the sol by DLS (Figure 7c).

**Discussion**

The properties of block copolymer-derived vitrimers arise from the subtle interplay of phase separation and network defects. Below, we discuss how these interrelated features influence the formation, thermomechanical properties, and structure of vitrimers derived from phase-segregating block copolymers.

**Gelation: the formation of hyperbranched polymers in solution reflects loop defects.** The importance of network defects in these materials is most obvious when the block copolymers are cross-linked in solution to form stable 10-nm particles. Even at a relatively high concentration (170 mg/mL) in THF, a good solvent for both blocks, intra-strand cross-linking (i.e., primary loops) and cross-linking among a limited number of block polymer unimers are favored over long-range elastically-effective linkages. The resulting soluble hyperbranched polymers can only exist if a significant fraction of the cross-links is engaged in loop defects. In permanent networks, loop defects increase as the concentration of network precursors decreases, consistent with an intramolecular cross-linking process.\textsuperscript{40}

Based on the DLS and DOSY NMR studies, we propose that in solution the block copolymers react rapidly with TREN, consuming ketoester bonding sites and forming hyperbranched polymers rich in primary and higher-order loops. As solvent is removed, bond rearrangement continues to occur until a sufficient number of elastically effective cross-links are forged. A schematic for the proposed process is shown in Figure 8. Flory-Stockmayer and Carothers equations, which do not account for network defects, predict that **Block-13-Net** (average functionality of 5.6) should be well over the critical gel
point at full conversion of the ketoesters. Nevertheless, Block-13-Net did not form the expected elastomeric vitrimer even after removal of solvent.

![Diagram of polymer formation and transformation](image)

**Figure 8.** Schematic of the proposed formation of cross-linked hyperbranched polymers and topological transformation to networks upon removal of solvent.

We hypothesize these loop defects are more favorable when the functional monomers are closely spaced, as is the case in the PnBA-\(b\)-PAEAA prepolymers. Consistent with this hypothesis, statistical poly(nBA-\(co\)-AEAA) copolymers with 18 or 28 wt\% AEAA (Stat-18 and Stat-28) gel rapidly when TREN is added to 170 mg/mL THF solutions, based on the flow inversion test and oscillatory shear rheology measurements (see SI, Figure S-18).

This work demonstrates that sequence can affect the connectivity of dynamically cross-linked polymers in solution. This insight can also be useful for processing: block copolymer vitrimers display critical gel points that are a function of concentration, rather than conversion of reactive sites. The solutions of hyperbranched polymers can be exploited in situations where low viscosity is required, then solvent can be removed to form the vitrimer without any additional reagents or stimuli. While associative exchange reactions have been used to transform the architecture of dynamic covalent polymers between linear, lightly cross-linked, and network states, these transformations relied on the
addition of competitive nucleophiles.\textsuperscript{41} The sensitivity of this vinylogous urethane block copolymer vitrimer system to concentration at ambient temperature more closely resembles supramolecular polymers than typical dynamic covalent systems, which require heat, light, or reagents to trigger reconfiguration.\textsuperscript{35}

**Thermomechanical properties and swelling ratios:** the dry vitrimers have more network defects than the swollen networks. The two $T_g$ transitions of Block-24-Net and Block-34-Net clearly reflect microphase separation. The lower $T_g(\sim 26 ^\circ C)$ is assigned to the PnBA-rich domain and the higher $T_g(79–96 ^\circ C)$ is assigned to the cross-linked PAEAA-rich domain. In Block-48-Net and Block-66-Net, solely the high-temperature transition is clearly evident by DMTA. The increase in $T_g$ for the PAEAA-rich domain of the vitrimers compared to the prepolymers is expected because cross-links restrict segmental mobility.\textsuperscript{42,43,44} However, the $T_g$’s corresponding to both the PnBA- and PAEAA-rich domains increase after cross-linking, suggesting that significant mixing of the blocks has occurred, positioning vinylogous urethane cross-links in the PnBA-rich domains.\textsuperscript{45}

Cross-link densities $u_{\text{modulus}}$ for the dry vitrimers can be calculated from the rubbery plateau modulus, $E'$, at 120 $^\circ C$ using eq 2:

$$u_{\text{modulus}} = \frac{E'}{3RT} \quad (\text{eq 2})$$

We observe a non-monotonic trend in cross-link density with increasing % AEAA (Figure 9). Specifically, Block-34-Net exhibits an anomalously low cross-link density, with approximately half as many elastically effective cross-links per unit volume as Block-24-Net, despite having approximately 50\% higher functionality in prepolymers with similar molecular weight and dispersity. The low crosslink density in dry Block-34-Net is also supported by its second $T_g(79 ^\circ C)$, which is lower than that of the other vitrimers (95–99 $^\circ C$).
Figure 9. Comparisons of cross-link density calculated for the dry network, calculated from the rubbery plateau modulus using the equation $\nu = E'/3RT$ (teal), and the cross-link density calculated for the network swollen in THF, calculated from the swelling ratio using the Flory–Rehner equation (orange).

For permanently cross-linked networks, the cross-link densities calculated from rubbery plateau moduli correspond, albeit imperfectly, to those calculated from swelling ratios because the network topology is the same in the dry and swollen states.\(^{46}\) In vitrimers, however, if associative cross-link exchange occurs under the conditions of swelling, the swollen networks can adopt different topologies in the dry and swollen states. The cross-link densities of equilibrated swollen networks, $\nu_{\text{swelling}}$, can be calculated using the Flory–Rehner equation (eq 3):\(^{47}\)

$$\nu_{\text{swelling}} = - \frac{\ln(1-v_2)+v_2+\chi_{12}v_2^2}{V_1(v_2^{1/3}-2v_2/f)} \quad (\text{eq } 3)$$

where $v_2$ is the volume fraction of polymer network in the swollen sample at equilibrium calculated from the swelling ratio, $V_1$ is the molar volume of THF, $f$ is the average functionality of the prepolymer and TREN, and $\chi_{12}$ is the Flory-Huggins polymer-solvent interaction parameter, which we approximate using the literature value for PnBA in THF\(^ {48}\) (see SI for details; $\chi$ values for PAAEA and PAAEA condensed with TREN are not known).
Diluting a vitrimer by swelling is expected to decrease the cross-link density relative to a dry network because dilution favors intramolecular crosslinks. Indeed, this trend is observed for other vitrimer chemistries and for our statistical copolymer control vitrimers, Stat-18-Net and Stat-28-Net (Table S-7). It is therefore surprising that the calculated cross-link densities for swollen Block-24-Net, Block-34-Net, and Block-48-Net are higher than cross-link densities for the same materials in their dry form (Figure 9). Additionally, Block-34-Net does not display anomalously low cross-link density once swollen. Even considering the errors in Flory–Rehner theory and the approximations we have made, the discrepancies between $\mu_{\text{modulus}}$ and $\mu_{\text{swelling}}$ in both their absolute values and their trends suggest that the network topologies of block copolymer vitrimers are altered in the swollen and dry states. While this discrepancy could arise from the loss of low-functionality strands to the sol fraction, which was previously observed in phase-separated polyethylene/dioxaborolane vitrimers with gel fractions <40%, the block copolymer vitrimers described here have gel fractions $\geq 94\%$.

The persistence of cross-linked hyperbranched polymers in solution shows that the block copolymer architecture encourages the formation of network defects in solution; we expect that this trend extends to the network. While directly comparable Stat and Block vitrimers have not been synthesized, Stat-18-Net and Stat-28-Net both have higher rubbery plateau moduli (1.9 and 4.8 MPa, respectively) than Block-34-Net (1.2 MPa), despite having fewer AAEA groups in the prepolymer (Figure S-20). Both TREN and the prepolymer have functionality $\geq 3$, but TREN molecules having 1 or 2 free amines can form primary loops and dangling ends, respectively, since these vitrimers are prepared with 33% excess amine. A schematic depicting relevant types of network defects is shown in Figure 10. Interestingly, in the case of poly(methacrylate) vinylogous urethane vitrimers, Epps and Sumerlin did not observe a significant difference in the rubbery plateau moduli for vitrimers derived from statistical and diblock copolymers. The prepolymer in that study featured equimolar incorporation of non-functional (nBMA) and functional (AAEMA) monomers, corresponding most closely to
**Block-66-Net.** Despite the many similarities of that system to the present one, our use of more flexible acrylic backbones and TREN cross-linker may encourage the formation of loops. Furthermore, the low $M_n$ values of our prepolymers rule entanglements as contributors to the rubbery plateau moduli. In a clear demonstration of sequence effects on network topology, Alabi prepared homogeneous, permanently cross-linked thermosets from sequence-defined oligomers.\textsuperscript{51} A blocky AAAmmm sequence (where A is the cross-linkable unit) yielded significantly lower rubbery plateau moduli than an alternating mAmAmA sequence. This difference was ascribed to the combined effects of dangling chain ends and primary loops.

**Figure 10.** Depiction of different types of network defects that can occur in the block copolymer vitrimers.

Based on evidence of microphase separation from the $T_g$ and SAXS data, we propose that the dry vitrimers have more loop defects than their swollen counterparts because the dry state is microphase separated. This separation creates cross-link-rich PAAEA domains and cross-link-poor PnBA domains. In order to percolate an infinite network, “tie strands” of cross-linked PAAEA blocks must mix with the PnBA-rich phase to link cross-link-rich domains. The increase in the $T_g$ of the PnBA phase observed for **Block-24-Net** and **Block-34-Net** are consistent with this scenario. Furthermore,
the cross-link-rich phase can achieve an infinite network within a bicontinuous morphology; this scenario is more likely for Block-48-Net and Block-66-Net, which have higher fractions of the cross-linked block. Unfortunately, SAXS of the vitrimers does not provide insight into the morphology, and TEM imaging was not possible.

**Phase separation: dynamic covalent cross-links hinder organization.** The domain spacing of the block copolymers before and after associative cross-linking is consistent with loop-rich domains in the vitrimers. For Block-13 through Block-48, the domain spacing shrinks by 1.8–11.6% after cross-linking. Isono and Satoh have shown that intramolecularly cross-linking one block of a diblock copolymer under dilute conditions significantly reduces the chain dimensions, providing significantly (9–22%) downsized features after bulk self-assembly relative to the non-cross-linked copolymers. In that case, the topology of the polymers can be considered entirely primary loops and no elastically effective cross-links; indeed, in our study, Block-13-Net has the fewest elastically effective crosslinks and the largest reduction in domain spacing. In contrast, Block-66-Net displays an increase in domain spacing after cross-linking, suggesting the presence of fewer network defects and, possibly, the influence of added TREN, which comprises 17.5 wt% of the vitrimer.

As described in the previous section, the thermomechanical data provides some evidence for phase separation, particularly for Block-24 and Block-34, which clearly possess two $T_g$’s. In contrast to the prepolymer, which form well-defined morphologies depending on their composition, the vitrimers display a single broad scattering peak by SAXS. This disorganization can arise from thermodynamic origins if the lowest-energy state of the system is disorganized, and/or and kinetic origins, if the system experiences insurmountable barriers to achieving an organized state. Condensation of prepolymer with $N,N$-dimethylethlenediamine results in a similar loss of organization, supporting the hypothesis that the vinylogous urethane blocks have a lower $\chi$ parameter with PnBA compared to PAAEA.
Nevertheless, a more polar diamine cross-linker also resulted in disorganized phase-separated morphologies, suggesting that kinetic barriers also play a role in hindering high-order organization of block copolymer-derived vitrimers that have been cross-linked in the disordered state. In order for hyperbranched polymers such as those shown in Figure 8 to arrange themselves into a lamellar or cylindrical structure, the PnBA must be separated from the cross-linked PAAEA. Evolution of a regular periodic structure from a disorganized one requires the cooperative rearrangement of many domains at the same time, which would be impeded by the presence of the cross-links, even if they are dynamically active on the molecular level. We envision that such a reorganization would require considerable energetically unfavorable molecular distortion to pack into a regular array that optimizes chain stretching and interfacial tension as required for a regular periodic morphology.

Permanently cross-linking a block copolymer in the disordered state, either in a melt above $T_{ODT}$ or in a good solvent, can permit or prevent re-organization of the resulting network depending on cross-link density. Lightly cross-linked block copolymer networks reassemble into organized morphologies upon cooling or solvent removal, but over a critical cross-link density, the networks are trapped in a disordered or disordered-like state.\textsuperscript{53,54,55,56} Dynamic cross-links can also interfere with long-range order; there are several studies in which phase-separating block copolymers were cross-linked by H-bonds,\textsuperscript{4,5} hindered urea dynamic covalent bonds,\textsuperscript{57} or metal-ligand complexation.\textsuperscript{6,7,58,59,60} In these examples, the cross-linked networks generally exhibit a single broad SAXS peak, consistent with disorganized phase-separated morphologies; corresponding transmission electron micrographs (TEMs), when available, reveal micellar or bicontinuous morphologies.\textsuperscript{57,6,7} Interestingly, Noro and Matsushita could regain long-range order in their H-bonded block copolymer networks by adding a salt that interacted selectively with one block, raising the effective interaction parameter.\textsuperscript{5} However, the kinetic barriers to reorganizing the H-bonded network are significantly lower than the barriers to reconfiguration in a vitrimer.
Most relevant to the present study, Epps and Sumerlin studied the assembly of methacrylic diblock copolymer vitrimers formed from bulk solution cross-linking and from solvent vapor annealing of self-assembled thin films with the cross-linker. While the thin films maintained the lamellar morphology after cross-linking, as shown by AFM and SAXS, the vitrimers formed from bulk cross-linking exhibited a single broad SAXS peak. Presumably, cross-linking pre-assembled block copolymers in the solid state circumvents the kinetic barriers associated with reorganizing the network; however, transport limitations may restrict this strategy to thin films.

Based on our data and examples in the literature, we conclude that there are several challenges to achieving organized morphologies in pre-cross-linked block copolymer-derived vitrimers. Consideration must be given to the thermodynamics of mixing the blocks once the cross-linker is incorporated, particularly when the volume fraction of the cross-linked block is significant. In some CANs formed from homopolymer or small-molecule precursors, incorporation of the dynamic cross-links promotes phase separation. The extent of cross-linking may also affect the barriers to reconfiguration, with lightly cross-linked networks more readily organized than highly cross-linked ones. It remains to be determined whether the mechanism of dynamic covalent exchange, associative vs. dissociative, affects a network’s ability to achieve organized morphologies. For applications that require mechanically robust materials with transport through a continuous phase, such as filtration membranes and solid-state polymer electrolyte membranes, the disorganized morphology of solution-annealed block copolymer vitrimers may in fact be advantageous.

**Stress relaxation:** relaxation times and activation energies do not increase with cross-linking. In the majority of reported vitrimers, increasing the cross-link density increases the relaxation time $\tau^*$ and the flow activation energy $E_a$. These trends are convoluted by variations in the content of free nucleophile, which is why all our vitrimers were prepared with amounts of TREN that result
in a constant 33 mol% excess free amine. Our lab has previously shown that changing the structure of the exchangeable group in the cross-linker can modulate stress relaxation rates through changes to $\tau_0$ without affecting $E_a$.\textsuperscript{16} In this work, vitrimers derived from block copolymers display a tunable modulus based on cross-link density over a wide range, but have nearly constant $E_a$. \textbf{Block-66-Net}, the most highly cross-linked material studied here, has slightly lower $E_a$ (97 kJ/mol) compared to \textbf{Block-24-Net} through \textbf{Block-48-Net} (112–116 kJ/mol). The block copolymer vitrimers also have nearly identical relaxation times at 160 °C, with the exception of \textbf{Block-24-Net}, which has the lowest fraction of functional block but approximately 5 times slower stress relaxation. This work demonstrates that block copolymer composition offers a strategy to increase the stiffness of vitrimers without increasing reprocessing times.

To rationalize the observations in this paper, we turn to predictions by Ciarella and co-workers, based on molecular dynamics simulations of star polymer-based vitrimers. They hypothesized that loops would cause stress relaxation to be more facile.\textsuperscript{66} Swapping an elastically-effective bond for another stress-bearing bond over short distances will not relax stress very much. On the other hand, swapping an elastically-effective bond for a loop removes the ability of that linkage to bear stress, causing relaxation. In complementary work, Ricarte and Shanbhag used an inhomogeneous Rouse model to show that vitrimers with blocky distributions of cross-linkers will relax stress faster than those with uniform cross-link distribution.\textsuperscript{67} These theoretical studies, however, did not take microphase separation into account. Experimentally, like Epps and Sumerlin,\textsuperscript{23} we do not observe that block copolymer vitrimers relax stress faster than compositionally similar statistical copolymers, and in fact retain a small amount of stress at long time scales (Figure 5c). Fortunately, Torkelson has shown that reprocessing does not require a vitrimer to be capable of full stress relaxation.\textsuperscript{33}
Stress relaxation rates in microphase-separated vitrimers represent a combination of the cross-link exchange rate, which is the rate-limiting step in homogeneous vitrimers above $T_g$, and the chain diffusion rate. Associative exchange events that occur within isolated cross-link-rich domains will not contribute significantly to stress relaxation. Ricarte and Leibler have proposed that in polyethylene/dioxaborolane vitrimers, the cross-linked segments of chains must disengage from cross-link-rich aggregates and diffuse through the cross-link-poor matrix to reduce topological stress, significantly hindering stress relaxation in self-assembled vitrimers compared to homogeneous vitrimers.68 These barriers to chain diffusion similarly explain the absence of terminal relaxation in block copolymer vitrimers.

In our microphase-separated system, diffusion on the length scales required to relax stress is expected to be faster through a continuous cross-link-rich phase compared to chain hopping between isolated cross-link-poor domains. Thus, increasing the volume fraction of cross-link-rich domains facilitates bond exchange across longer length scales, counterbalancing the effect of increasing cross-link density. The uniquely slow relaxation in Block-24-Net suggests that the cross-link-rich domains in this vitrimer are discontinuous (micellar or spherical-like, Figure 11a), while the other block copolymer vitrimers have continuous domains of the cross-link-rich phase (Figure 11b). Furthermore, following Ricarte and Leibler’s logic, a strand bearing loops or dangling ends should disengage from the cross-link-rich domain and diffuse through the majority PnBA phase more readily than an elastically effective strand.68 From rubbery plateau moduli, we know that Block-24-Net has fewer network defects than Block-34-Net. We propose that the combination of lower volume fraction of the cross-linked block and fewer network defects dramatically hinder stress relaxation in Block-24-Net.
**Figure 11.** Illustration of different factors that contribute to stress relaxation times in (a) **Block-24** and (b) **Block-34.** The yellow regions are the cross-linked PAAEA-rich phase, and the red matrix is the PnBA-rich phase. The green arrows indicate the path traversed by a strand to relax stress; for clarity, only selected strands are shown.

**Conclusion**

In this contribution, we explored the composition of block copolymer precursors as a design variable in vitrimers. Fundamentally, we show that the properties of this class of vitrimers are influenced by the interplay of microphase separation and network defects. The block sequence introduces more defects as a result of the adjacent cross-linking moieties, an effect further exacerbated by microphase separation in the dry state. Loops, in turn, shrink the domain size of the disorganized morphologies. We propose that rearrangement of these block copolymer vitrimers into organized morphologies is prevented by a combination of thermodynamic and kinetic factors. Microphase
separation also prevents direct comparison of these experimental results to theories describing the role of loops and block sequence on stress relaxation in vitrimers, which assume homogeneity.

Two experimental directions are recommended to provide clearer connections to theory. First, measurements of systems with decreased segregation strength, or of gels homogeneously swollen by a good solvent, are needed to precisely determine the effect of sequence on vitrimers in the absence of microphase separation. Secondly, quantification of the loop defects in vitrimers is required. Our lab is currently pursuing this latter goal using Network Disassembly Spectrometry.\textsuperscript{69,70,71}

This study also presents several practical lessons for the design of vitrimers. First, block copolymers can be cross-linked in relatively concentrated (130–170 mg/mL) solutions to obtain soluble hyperbranched polymers. These solutions may be preferable for processing techniques that require lower viscosity. Upon solvent removal, vitrimers with high ($\geq$94\%) gel fraction are formed. Network defects must also be considered when targeting the appropriate functionality for prepolymers, as gel points significantly lower than calculated by Flory–Stockmayer or Carothers analysis are observed (Block-13). The moduli of the resulting materials will also reflect more network defects than in a homogeneous vitrimer, and may not increase monotonically by increasing prepolymer functionality. While these block copolymer vitrimers produced disorganized microphase-separated morphologies, we present considerations for future work seeking to achieve better organized morphologies. Block copolymers provide a convenient platform to increase the stiffness of vitrimers by increasing cross-link density, without affecting the flow activation energy or stress relaxation time. Finally, vinylogous urethane block copolymer vitrimers can be partially dissolved by heating in a good solvent. Further work is required to achieve complete solubilization of the vitrimer into hyperbranched polymers, which could present an alternative route to “chemical” recycling of hydrolytically stable vitrimers without any added nucleophilic reagents.
Supporting Information

Experimental procedures; sample preparation; NMR, FTIR, and UV-vis spectra; additional stress relaxation, DMTA, and SAXS data; details of cross-link density calculations

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