Effects of Branching and Crystallization on Rheology of Polycaprolactone Supramolecular Polymers with Ureidopyrimidinone End Groups

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ABSTRACT: The rheology of supramolecular polycaprolactone polymers with two ureidopyrimidinone (UPy) end groups and unimer molecular weights between 600 and 4000 was compared with that of branched tri- and tetrafunctional analogs. Above the melting point of polycaprolactone, the previously observed low-frequency plateau in storage and loss moduli during oscillatory frequency sweep of lower molecular weight bifunctional unimers was shown to persist up to 130−170 °C, where it gradually disappeared but reappeared upon slow cooling. Even though they are supramolecularly cross-linked, the tri- and tetrafunctional materials showed no plateau. This counterintuitive behavior was further investigated with optical microscopy, WAXS, and DSC experiments, which indicated that the plateau is closely connected to the presence of crystalline domains in the lower molecular weight fraction of bifunctionalized unimers. Because the formation of crystallites is prevented by branching, and because the network formed by the tri- and tetrafunctional unimers has a short lifetime, the branched materials do not show a low-frequency plateau.

Introduction

Over the last two decades, supramolecular polymers have gained increased interest because of their properties that combine features from high- and low-molecular-weight materials. As opposed to fully covalently bonded classical polymers, supramolecular polymers consist of unimers of low-to-medium molecular weight, which interact with each other through noncovalent ligand interactions, van der Waals, hydrophobic, or hydrogen-bonding interactions. Because of the dynamic character of the supramolecular bonds, supramolecular polymers often show much better solubility and lower melt viscosity and hence display better processability than classical long chain thermoplastics. The physics of supramolecular polymers, particularly their unique rheology, has received attention since about 20 years ago. The foundations of stress-relaxation and rheology of supramolecular polymers in solution were laid by Cates and rheology of supramolecular polymers was pioneered by Stadler. Compared with single hydrogen bonds, multiple hydrogen bonds have higher strength and directionality, and many systems with arrays of three, four, or more hydrogen bonds have been investigated. Initial theoretical and exploratory studies have more recently been complemented by work related to potential applications.

A particularly useful building block in supramolecular polymers is the ureidopyrimidinone (UPy) scaffold (Scheme 1), which was developed in our laboratory more than a decade ago. It features a linear array of four hydrogen bonding sites, and strongly dimerizes in most organic solvents ($K_{\text{dim}} = 6 \times 10^7 \text{M}^{-1}$ in CHCl$_3$ at 25 °C). Although at any time most UPy units are in their dimerized state, their strongly temperature-dependent exchange dynamics results in thermorheological behavior that is distinctive for supramolecular polymers.

In our laboratory, many types of polymers have been functionalized with UPy units, one of the most widely investigated ones being poly-ε-caprolactone (PCL). Living polymerization of the ε-caprolactone monomer can be initiated with a wide variety of alcohols and produces alcohol-terminated polymers with narrow polydispersities (Scheme 2a). In the absence of any particular functionalization, PCL is semicrystalline with a melting point at 60 °C and a glass transition at −60 °C. In addition, the biocompatibility of PCL has made it a material of choice in tissue engineering research. Much attention is currently being dedicated to developing industrial applications for UPy-based materials.

We recently showed that the rheological properties of UPy-containing supramolecular polymers above the crystalline melting point of the PCL main chain are strongly affected by the tendency of dimerized UPy end groups to aggregate into

Scheme 1. Dimer of Ureidopyrimidinone (UPy)
It was found that if the UPy unit is linked to the PCL via a urethane linkage (Scheme 2b), which enhances stacking of UPy dimers, then a plateau is observed at low frequency in the rheological spectra where $G'$ and $G''$ are nearly constant over several decades of frequency. However, when the UPy groups have bulky substituents or if the urethane linker is replaced by an ester bond these fibers are absent, and the rheological behavior is much simpler.

In the current Article, we report a systematic study of the structure–rheology relationship of a series of PCLs with urethane-linked UPy units (Scheme 3). For the sake of clarity, the molecules formed by functionalization of PCL macromonomers, which are the monomeric units in the supramolecular polymerization, are denoted throughout this Article as unimers, whereas we refer to the materials, the products of supramolecular polymerization, as the (supramolecular) polymer. The topology was varied by increasing functionality to 3 or 4 in branched unimers with branches ranging in molecular weight from 300 to 2000. Scheme 3 depicts the di-, tri-, and tetrafunctional UPy–urethane unimers 2 to 4. These unimers provide a unique set of materials that allow us to compare the effects of supramolecular cross-linking on rheology. Unexpected differences in rheology between bifunctional and branched materials encountered during the course of this investigation prompted us to investigate the morphology of the linear supramolecular polymers in more detail using WAXS, DSC, and polarizing optical microscopy (POM), leading to deeper insight into the origins of the complex rheological behavior of UPy-functionalized linear PCL.

**Experimental Section**

**Chemicals.** OH-telechelic macromonomers as precursors for all unimers studied were obtained by ring-opening polymerization of $\varepsilon$-caprolactone (Aldrich) using tin(II) 2-ethylhexanoate (Sn(Oct)$_2$, Aldrich) or fumaric acid (Acros) and various initiators, designated as $R$–OH in Scheme 2a. The latter were

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**Scheme 2.** (a) Synthesis of UPy–Urethane-Functionalized PCLs and (b) Ureidopyrimidinone Linked to a Polymer via a Urethane (Top) or an Ester (Bottom) Linkage

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**Scheme 3.** Structures of the Unimers and Small Molecules Used

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*Numbers of monomer units $n$ are approximate average values.*
diethylene glycol (Acros) for 2a–d, 1,1,1-trimethylpropane (TMP, Aldrich) for 3a, 2-hydroxyethyl-1,3-propanediol (Fluka) for 3b, and pentaerythritol (Aldrich) for 4. Dibutyl tin dilaurate (DBTDL, Aldrich) was used as a catalyst for urethane formation. Solvents were purchased from Biosolv (CHCl₃, heptane, toluene) or Cambridge Isotope Laboratories (CDC₁₂, DMSO-d₆). γ-Caprolactone was distilled from CaH₂ under reduced pressure. All other chemicals were used as purchased.

**Characterization.** ¹H NMR and ¹³C NMR spectra were recorded on a Varian Mercury 400 spectrometer. ¹H and ¹³C chemical shifts are reported in ppm relative to the solvent peak.

**Differential Scanning Calorimetry (DSC).** DSC measurements were performed on a TA differential scanning calorimeter Q-2000 fitted with a refrigerated cooling system under a nitrogen atmosphere with heating and cooling rates of 10 °C/min and subsequently cooled to 20 °C/min.

**Rheology.** Films were prepared by dissolving the polymer in dry CHCl₃ and casting the solution into a silanized Petri dish, followed by slow evaporation of the solvent and drying the film in vacuo. Discs of 25 mm diameter were punched out for rheometry.

**Synthesis and Analytical Data.** Experimental details of unimers 2b–d have been previously published. All newly synthesized unimers were obtained via the following procedures.

Dry γ-caprolactone and initiator were mixed at 90 or 120 °C under nitrogen without any solvent. After the addition of the catalyst, the mixture was allowed to react overnight at 90 or 120 °C. When Sn(Ot)₂ was used, 0.01 equiv relative to the initiator was added as a 0.2 M solution in dry toluene; when fumaric acid was used, 0.4 equiv relative to the initiator was added. The degree of conversion can easily be monitored by ¹H NMR. After full conversion, the reaction mixture was diluted with CHCl₃, filtered through a glass filter when fumaric acid was used, and precipitated into an eight-fold volume of heptane. The precipitated polymer was rinsed with heptane and dried in vacuo. Subsequent functionalization with isocyanate 5 followed the reported procedures.

**Results**

**Synthesis and Molecular Characterization.** The linear and star PCLs were synthesized by ring-opening polymerization at 90 or 120 °C in bulk, as previously reported, without any solvent and using β-, γ-, or tetrafunctional initiators. The polymers were end-functionalized with (Me)₂P═C═O—hexamethylene isocyanate 5 (Scheme 2, results summarized in Table 1).
After functionalization, the reaction mixture was diluted with chloroform, filtered, and precipitated into n-pentane or n-heptane. High yields and purities were reproducibly obtained, as ascertained by 1H NMR, GPC, and ESI-MS. Sn(Oct)$_2$ was generally used as the polymerization catalyst; in a set of control experiments, unimers 2a and 2b were also synthesized using fumaric acid as a catalyst to exclude possible artifacts related to residual catalyst in the present studies.

For the sake of clarity, the unimers are referred to as ‘n × M’, where n refers to the functionality of the unimer (or the number of UPy—urethane units per molecule) and the number M refers to the approximate number-average molecular weight of one arm of the molecule, without the UPy—urethane part.

Polystyrylactone (PCL) is narrow, well-defined, and reproducible. Figure 1 shows that all oligomers of 2b 2 × 500 can be easily discerned as the adducts with H$^+$ and Na$^+$ as well as doubly charged species.

Differential Scanning Calorimetry. DSC was performed on all unimers to record their relative melting points. In our previous study,$^1$ three thermal transitions were observed below 130 °C: a glass-transition temperature ($T_g$) related to the PCL part of the unimers and two melting points ($T_{m1}$ and $T_{m2}$) due to the melting of the crystalline domains of PCL and of the stacked UPy—urethane units. The $T_g$ is in the region of ~60 to ~80 °C, and the two melting points lie between 40 and 90 °C; they are often close to each other and coincide in most unimers (Figure 2a). Although UPy groups decompose if kept for prolonged times at temperatures above 150 °C, control experiments show that they are unaffected by higher temperatures at the short time spans of DSC runs. Above 130 °C, the shorter bifunctional unimers 2a–c reproducibly display weak, broad transitions at temperatures between 130 and 170 °C (Figure 2b), whereas the other unimers, including all tri- and tetrafunctional species as well as 2d 2 × 2000, do not show any further transition (Figure 2c).

As shown in Figure 2, the endotherm at higher temperatures indicates the presence of an additional melting process. The shape of the peak varies somewhat from batch to batch and also from measurement to measurement within the same batch. Nevertheless, DSC traces reproducibly show the same features, which confirms the presence of a phase transition. (See the Supporting Information.) The shape of the peak does not allow for accurate determination of total heat values of the transitions.

The transition invariably appears in first heating runs. If the material is then cooled slowly from 180 °C (e.g., 3 °C/min), then the second heating run also displays an endotherm at similar temperatures. (See the Supporting Information.) If the supramolecular polymer is cooled quickly (20 °C/min), then no melt endotherm is observed above 130 °C in second heating runs. Moreover, cooling runs at 20 °C/min do not display any exotherm, indicating that the reverse transition is too slow or does not occur at all.

To establish the fact that the high temperature transition is not due to a fully irreversible phenomenon such as decomposition of the polymer, a sample of 2 × 500 was heated to 180 °C for 15 min and subsequently subjected to the usual film-forming treatment. The films obtained this way display a similar endotherm, confirming that the entity causing the peak remains structurally intact up to 180 °C. Furthermore, the rheological behavior and the 1H NMR spectrum in CDCl$_3$ remain unaltered. (See the Supporting Information.) Although crystallization is slow, its reversibility was established by temperature-modulated DSC (TM-DSC) using a linear heating rate of 3 °C/min and a superimposed

| Table 1. Polymerization Catalysts Used, Molecular Weights (1H NMR and GPC), and Polydispersity Index (PDI) (GPC) for the Unimers Presented Here$^a$ |
|-------------------------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| **unimer**                    | **n × M**       | **catalyst**    | **$M_n$ (NMR)** | **$M_d$ (GPC)$^b$** | **PDI** |
| 2a                            | 2 × 300         | Sn(Oct)$_2$    | 1.2 × 10$^3$    | 8 × 10$^2$          | 1.44   |
| 2b                            | 2 × 500         | Sn(Oct)$_2$    | 1.5 × 10$^3$    | 1.7 × 10$^3$        | 1.29   |
| 2c                            | 2 × 1000        | Sn(Oct)$_2$    | 2.7 × 10$^3$    | 3.7 × 10$^3$        | 1.68   |
| 2d                            | 2 × 2000        | Sn(Oct)$_2$    | 5.7 × 10$^3$    | 7.0 × 10$^3$        | 1.49   |
| 2e                            | 3 × 300         | Sn(Oct)$_2$    | 1.6 × 10$^3$    | 1.8 × 10$^3$        | 1.47   |
| 2f                            | 3 × 1000        | Sn(Oct)$_2$    | 4.5 × 10$^3$    | 5.0 × 10$^3$        | 1.50   |
| 4                             | 4 × 1000        | Sn(Oct)$_2$    | 4.6 × 10$^3$    | 5.3 × 10$^3$        | 1.50   |

$^a$Molecular weight values were measured on the UPy-telechelic oligomers.$^b$In THF, using polystyrene standards.

Figure 1. ESI-MS spectrum of 2b 2 × 500 with exact m/z values and the number of ε-caprolactone units in the oligomers. The m/z values correspond, respectively, to adducts with H$^+$, Na$^+$, and 2Na$^+$—H$^+$. 

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oscillation (period 1 min, amplitude 1 °C). The high-temperature transition clearly appears in the reversing heat flow, with only a minor component in the nonreversing trace (Figure 3). These findings imply that the transitions observed in the “plateau polymers” correspond to an endothermic transition that is essentially reversible, such as a slow melting/crystallization phenomenon.

**Polarizing Optical Microscopy.** POM is a technique that allows visualizing optically anisotropic regions in materials because of the birefringence of these domains. Anisotropy can be due to, for example, strong chain alignment, crystallinity, or liquid crystallinity and gives rise to bright iridescent spots when polarizer and analyzer are crossed, whereas isotropic regions remain dark. In 2b 2 × 500, birefringent domains of sizes below 1 μm were observed under crossed polarizers at temperatures around 130 °C, that is, in the range above the UPy and PCL melting point(s) and below the high-temperature transition (Figure 4a). These domains are irregularly shaped and solid; they do not deform under shear or compression. When the temperature is raised above the onset of the second DSC endotherm, the domains slowly disappear to give a single liquid phase devoid of any birefringence at temperatures >180 °C (Figure 4b). When the material is slowly cooled, the birefringent domains reappear (Figure 4c,d), confirming the reversible character of the observed high-temperature transition.

**Rheology.** Rheological master curves were obtained by TTS generated from frequency sweep curves measured between the melting point of the polymer (typically 70—90 °C, attributed to the melting of crystalline PCL and UPy–urethane stacks) and 130 °C. All master curves in this Article are referenced to 100 °C, given that the melting points of some polymers were significantly >70 °C.

Figure 5 summarizes the master curves of all bi-, tri-, and tetrafunctional materials. In our previous study,1 we reported that polymers 2 × 500 and 2 × 1000 display a plateau down to low frequencies or on long time scales (on the order of several hours) on both rheological shear moduli. A similar plateau is observed in 2a 2 × 300; these polymers have short oligomer lengths and high volume fractions of UPy groups. Tri- or tetrafunctional species of similar length, molecular
Two of the plateau polymers, 2a and 2c, fail to master properly on the low-frequency (high temperature) side; instead they show pronounced fanning. This failure to obey TTS is expected for a polymer that is not in a molten state or an undercooled melt, that is, not in a homogeneous glassy state.

Rheological studies confirm the reversible nature of the transition that causes the plateau. In temperature ramp experiments with a heating rate of 3 °C/min, a constant angular frequency of 1 rad/s, and a strain of 2%, the elastic and viscous shear moduli of the plateau polymers remain high up to 140 °C (Figure 6). Only around this temperature do they plummet by two to three orders of magnitude; this is the region where the endothermic transitions were observed in DSC. In rheometry, the decrease also stretches over 20 to 30 °C rather than being a sharp change. This behavior is qualitatively the same for the three “plateau polymers”, whereas the other polymers display steadily decreasing moduli in the whole temperature range above the PCL/UPy–urethane melting point (Figure 6). These experiments can be conducted up to 160–165 °C when the viscosity of the material becomes too low to give sufficient torque to the rheometer’s transducer and when the polymer starts to flow out between the plates.

When the temperature is gradually lowered again, an increase up to the initial low temperature moduli values is observed. However, the behavior shows hysteresis: the heating and cooling traces do not superimpose. In polymers 2 × 500 and 2 × 1000, the moduli at a given temperature are lower in the cooling run than in the heating run. This is the expected result for slow crystallization. Surprisingly, 2 × 300 shows a negative hysteresis (Figure 6): the cooling run is characterized by higher moduli at a given temperature than the heating run. In 2a, the structure that induces the plateau appears to form more efficiently from the melt (at the cooling rate and shear conditions used) than from solution. This indicates that starting from the melt, the formation of cross-linking regions due to either crystallization of PCL or stack formation of end groups is more effective from the rheological point of view.

Wide-Angle X-ray Scattering. WAXS shows a distinct diffraction peak corresponding to a correlation distance of 11.04 Å (Figure 7a for 2 × 500, see the Supporting Information for other polymers) at room temperature as well as
several small peaks superimposed on the alkyl halo. Other polymers studied here display the same pattern.

Crystal structures of UPy-urethane-containing polymers have not been studied by X-ray crystallography. However, a correlation distance of 11.05 Å has been found in a crystal structure25 of a small molecule containing the urethane-hexamethylene-(methyl)UPy motif. The dimerized UPy's stack on each other, with an interplanar distance of 4.56 Å. Furthermore, the stacks of UPy dimers align next to each other (Figure 7b), with a repeating distance of 11.05 Å, interacting via a zipper-like series of weak hydrogen bonds arranged in pairs, involving the “keto” oxygen and the adjacent aromatic proton of the UPy’s (Figure 7b).

We propose that the UPy stacks present in the polymers at room temperature align in a similar fashion to the small-molecule single crystal, accounting for the measured correlation distance of 11.04 Å. In addition to this, the smaller peaks superposed on the so-called alkyl halo present in the diffraction pattern (Figure 7a) can be related to crystalline PCL and stacked UPy’s (interplanar distance of 4.56 Å).

At 120 °C, however, that is, in the plateau temperature range, WAXS of 2b/C2500 shows a weak but significant peak at the same distance of 11.04 Å (Figure 7a): this implies that the ordered state of aligned UPy stacks must persist to some extent at this temperature, a finding in good agreement with the presence of the birefringent domains observed in POM. The peaks superimposed on the alkyl halo cannot be discerned any more at this temperature.

The WAXS peak was observed in the batch of 2b/C2500 in which birefringent domains were discerned. Another batch of the same material or the other plateau polymers showed neither conspicuous diffraction peaks nor birefringent domains in POM, but they displayed the same melt behavior and rheological features. We ascribe this to insufficient scattering due to small crystallite sizes as well as low volume fractions of crystallites, low ordering in crystalline domains, or both.

Discussion

Among the supramolecular polymers studied, only 2a/C2300, 2b/C2500, and 2e/C21000 show the rheological plateau; all observations of additional peaks in DSC, crystallites in POM, or WAXS peaks at temperatures above 100 °C occurred in one or more of these polymers, whereas the plateau is absent in samples such as 3a/C2300, which contain similar ratios of PCL and UPy units but are branched. The conclusion is that only linear oligomers can lead to these ordered structures. In addition, higher molecular weights of the oligomer, that is, lower volume fractions of UPy-urethane groups, also do not lead to a plateau behavior, as demonstrated with 2d/C2200. It thus appears reasonable to impute the crystallization to oligomers of short to medium size, probably on the order of 2b/C2500. Such oligomers would contain around eight ε-caprolactone units.
We speculate that although unimers of several lengths are able to crystallize, each crystalline domain consists of oligomers of a single length. This would explain the observation of very broad melting ranges, some of which even appear to consist of several “steps” (Figure 2b), each corresponding to the melting of crystallites of one particular oligomer.

Also, the melting range observed in POM appears to be narrower than the range seen in DSC, despite careful calibration of both POM and DSC. This can in turn be explained by the fact that the crystals that melt first might be too small to be observed optically. We hypothesize that unimer 2b 2 × 300 has the best volume fraction of crystallizable oligomers; this is why it would be the only one to form visible crystals.

Nevertheless, crystallization must be governed by more subtle factors. Although the control batch of 2b material displays similar analytical, rheological, and calorimetric properties, it does not show crystallinity in POM. Similarly, in the other “plateau polymers”, 2a 2 × 300 and 2c 2 × 1000, no birefringence can be seen at any temperature above the first melting point, that is, ~60 °C. However, the domain sizes below 1 μm are already close to the range of wavelengths of visible light (400–700 nm) and thus at the low end of the sizes of structures that can be discerned under an optical microscope. We thus hypothesize that crystalline domains do exist in these polymers but that they are too small for optical detection. Birefringence was not detected in any of the polymers that have no plateau.

The crystalline domains are likely connected to the matrix by hydrogen bonding between UPy groups, which bind to UPy units from molten chains. Therefore, the crystallites act as supramolecular cross-links of very high functionality, causing rheological behavior typical of a viscoelastic solid. In this sense, the crystallites can be likened to a reinforcing filling material. This behavior is at first sight counterintuitive: oligomers with a functionality higher than two are able to form a network, which is expected to show higher moduli, especially a more markedly elastic behavior. Indeed, in concentrated chloroform solutions, the tri- and tetra-functional species form very thick gel-like phases, whereas the bifunctional ones give solutions with lower viscosity. Significantly, gel formation at very low degrees of crystallinity has also been described and studied in a covalent polymer (poly-1-caprolactones) show a rheological plateau indicative of the formation of transient physical network, whereas such a plateau is absent in analogous supramolecular polymers with higher functionality. Combined microscopy, WAXS, and DSC experiments indicate that the plateau is related to the presence of crystalline domains of low-molecular-weight bifunctional unimers. Therefore, the stacking of dimerized UPy end groups that has been identified as the origin of a rheological plateau in our previous publication has now been shown to have such a pronounced effect on rheology because it leads to the formation of stable crystallites in 2a and 2b, whereas in supramolecular polymers from tri- and tetrafunctional unimers the effects of UPy dimer stacking on rheology are minor, presumably because branching prevents the formation of crystallites of UPy units. Investigating the quantitative relation between crystallinity and viscoelastic behavior of 2a and 2b was beyond the scope of the current work, but the striking analogy of our observations to early gelation during crystallization of semicrystalline polymers recommends more detailed study of crystallization in supramolecular polymers.

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Supporting Information Available: Analytical data (1H NMR, ESI-MS) and additional rheology, POM, WAXS, and DSC data. This material is available free of charge via the Internet at http://pubs.acs.org.

References and Notes
