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Controlled Cationic Ring-Opening Polymerization of Monothiocarbonate

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Cyclic carbonates undergo ring-opening polymerization with both cationic and anionic initiators, which have been extensively studied by several research groups.¹ We have also found that cyclic carbonates show volume expansion during polymerization, which can be accounted for by the difference in strength of the intermolecular interaction between monomers and polymers.² Recently, sulfur-containing polymers such as polythioethers³ and polythioesters⁴ are gaining importance due to their excellent optical and thermal properties. In the course of the study of the ring-opening polymerization of cyclic carbonates, we have designed a sulfur analogue of cyclic carbonate, i.e., cyclic monothiocarbonate. We have reported the first anionic ring-opening polymerization of a six-membered cyclic monothiocarbonate, 1,3-oxathian-2-one, to afford poly(trimethylene thiocarbonate).⁵ Polythiocarbonate with the similar structure can be obtained by the cationic ring-opening polymerization of 1,3-dioxane-2-thione as reported by Kricheldorf et al.⁶ Unfortunately, the obtained polythiocarbonate is poorly soluble in common organic solvents; therefore, the detailed polymerization behavior is not sufficiently reported. This communication deals with the controlled cationic ring-opening polymerization of a six-membered cyclic monothiocarbonate, 5,5-dimethyl-1,3-dioxan-2-thione (**1**), as described in Scheme 1.

The monomer **1** was synthesized by the reaction of 2,2-dimethylpropane-1,3-diol with an equivalent amount of thiophosgene in the presence of 2 equiv of 2,3-dimethyl-1-phenyl-3-pyrazolin-5-one in chloroform at 50 °C for 12 h in 45% yield.⁷ The cationic polymerization of **1** was carried out with 2 mol % of trifluoromethanesulfonic acid (TfOH), methyl trifluoromethanesulfonate (TfOMe), boron trifluoride etherate (BF₃OEt₂), or triethyloxonium tetrafluoroborate (Et₃OBf₄) as an initiator at 30–80 °C under a dry nitrogen atmosphere as summarized in Table 1. The consumption of the monomer was achieved within 8 h in runs 1 and 2 and within 3 h in runs 3–5. Figure 1 shows the GPC profiles at the different polymerization time. There is no significant difference between the GPC profiles at 8 h and that at 24 h (conditions: monomer, 2.00 mmol; initiator, 0.040 mol; solvent, 2.00 mL of CH₂Cl₂; temperature, 30 °C), supporting the absence of backbiting reaction. In all cases in Table 1, polythiocarbonates with the narrow molecular weight distribution could be obtained, which

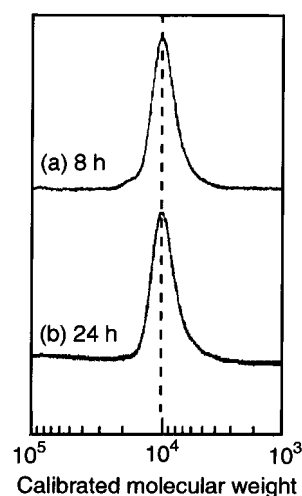
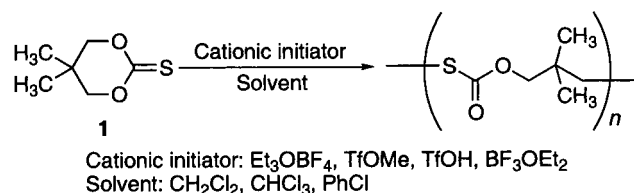


Figure 1. GPC profiles of the obtained polythiocarbonate at different polymerization times: (a) 8 h, (b) 24 h. Conditions: monomer, 2.00 mmol; initiator, 0.040 mol; solvent, 2.00 mL of CH₂Cl₂; temperature, 30 °C.

Scheme 1



exhibited good solubility in common organic solvents such as tetrahydrofuran (THF), chloroform, dichloromethane, and acetone. The conversion estimated by ¹H NMR spectroscopy was nearly equal to the yield of the polymer. The difference in the observed *M_n*'s among the initiators used would be due to the difference in initiation efficiency. The difference between the observed and expected *M_n*'s would be owing to the low initiation efficiency and/or the different solvation state in THF between the polythiocarbonate and the polystyrene standards. The obtained polymer could be isolated by precipitation with hexane, and the structure was examined by ¹H NMR, ¹³C NMR, and IR spectroscopy. Figure 2 illustrates the ¹H and ¹³C NMR spectra of **1** and the polymer obtained in the polymerization in run 3 in Table 1. The ¹H NMR spectrum of the polymer (Figure 2B) showed three singlet signals at 1.01, 2.94, and 4.07 ppm, which were assignable to the methyl group and the α- and γ-methylenes of the thiocarbonate sulfur, respectively. The ¹³C NMR spectrum of the polymer (Figure 2D) showed the corresponding three methylene carbon signals at 35.5, 39.5, and 73.6 ppm, the methyl carbon one at 23.4 ppm, and the carbonyl carbon one at 171.0 ppm. The assignments of the observed signals were confirmed by DEPT experiment. The ¹³C NMR spectrum of **1** (Figure 2C) showed the thiocarbonyl carbon signal at 189.1 ppm, indicating the progress of the cationic ring-opening polymerization accompanying isomerization of the thiocarbonyl group into the carbonyl group. The isomerization was supported by a C=O absorption peak at 1717 cm⁻¹ in the IR spectrum of the polymer. Scheme 2 illustrates a plausible mechanism for the ring-opening polymeriza-

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Table 1. Cationic Ring-Opening Polymerization of **1**^a

run	solvent	initiator	time (h)	temp (°C)	conv ^b (%)	M_n^c	M_w/M_n^c
1	CH ₂ Cl ₂	Et ₃ OBf ₄	24	30	>99	13 200	1.04
2	CHCl ₃	Et ₃ OBf ₄	20	40	>99	16 800	1.04
3	PhCl	TfOMe	12	80	>99	11 200	1.15
4	PhCl	TfOH	12	80	>99	13 900	1.13
5	PhCl	BF ₃ OEt ₂	12	80	>99	31 000	1.08

^a Monomer, 0.25 mmol; solvent, 0.25 mL; initiator, 0.005 mmol. ^b Determined by ¹H NMR spectroscopy measured in CDCl₃. ^c Estimated from GPC eluted with THF based on polystyrene standards.

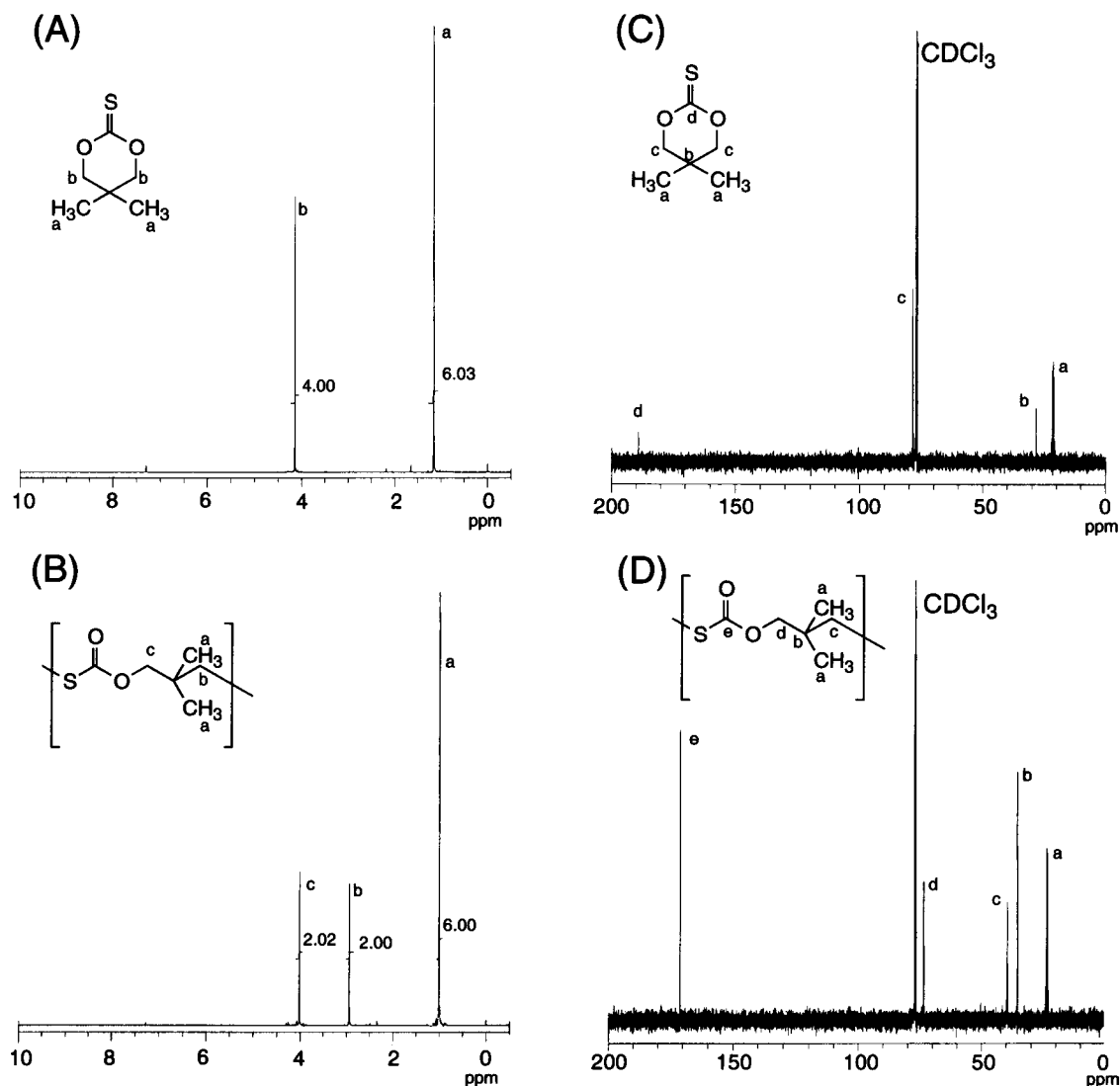
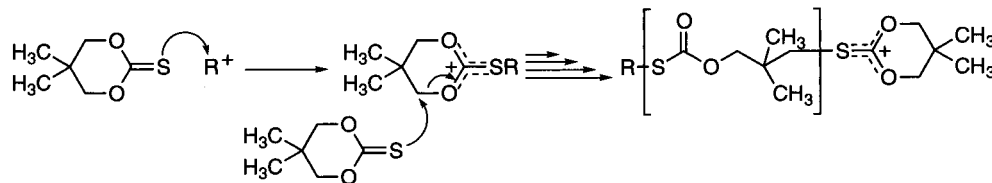


Figure 2. (A) ¹H NMR spectra (solvent; CDCl₃, 500 MHz) of **1**, (B) polymer obtained in the polymerization in run 3 in Table 1, (C) ¹³C NMR spectra (solvent; CDCl₃, 125 MHz) of **1**, and (D) polymer obtained in the polymerization in run 3 in Table 1.

Scheme 2



tion, which may proceed via the nucleophilic attack of a thiocarbonyl sulfur to a cationic initiator, followed by the nucleophilic attack of a thiocarbonyl sulfur to the α -position of an ether oxygen in a cyclic carbenium ion.

The relationship between the M_n of the polymer and feed ratio of $[1]/[Et_3OBf_4]$ was linear, and the M_w/M_n was around or less than 1.1 as shown in Figure 3. The

living character of the present polymerization was confirmed by postpolymerization. After the first stage polymerization of **1** (20 equiv to Et_3OBf_4) for 12 h at 0 °C, the second monomer (40 equiv to Et_3OBf_4) was fed to react continuously for an additional 12 h at 30 °C. The M_n of the obtained polymer increased by the second monomer feed as shown in Figure 4, though a small part

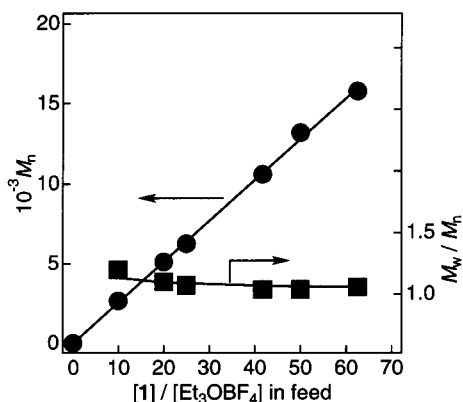


Figure 3. Relationships between M_n , M_w/M_n , and feed ratio of $[1]/[Et_3OBF_4]$. Conditions: monomer, 0.25 mmol; solvent, 0.25 mL of CH_2Cl_2 ; temperature, 30 °C.

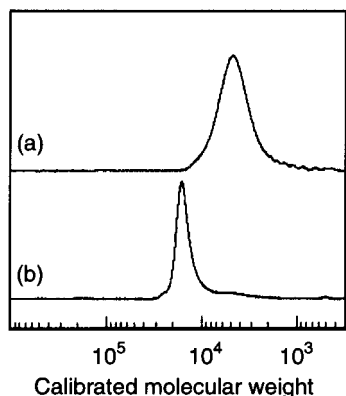


Figure 4. GPC profiles of the obtained polythiocarbonate in postpolymerization: (a) after prepolymerization of **1** (20 equiv to Et_3OBF_4), $M_n = 4200$ ($M_w/M_n = 1.16$), (b) after postpolymerization of **1** (40 equiv to Et_3OBF_4), $M_n = 12\,400$ ($M_w/M_n = 1.15$).

of the prepolymer chains did not completely grow, supporting quasi-living character of the polymerization.

In summary, the polymerization of 5,5-dimethyl-1,3-dioxane-2-thione (**1**) by several cationic initiators afforded polythiocarbonate with good solubility in common organic solvents and narrow molecular weight distribution. The molecular weight of the obtained polymer could be controlled by the feed ratio of the monomer to

initiator and increased when the second monomer was added to the polymerization mixture after quantitative consumption of **1** in the first stage, supporting that the cationic ring-opening polymerization of **1** proceeded via a quasi-living process.

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- (7) Mp 99–100 °C. 1H NMR δ ($CDCl_3$, 500 MHz): 1.15 (s, 6H, $-CH_3$), 4.18 (s, 4H, $-CH_2-O-$) ppm. ^{13}C NMR δ ($CDCl_3$, 125 MHz): 21.4 ($-CH_3$), 28.5 ($C-(CH_3)_2$), 78.5 ($-O-CH_2-$), 189.1 ($C=S$) ppm. IR (KBr disk): 2965, 2905, 2880, 1472, 1404, 1381, 1319, 1244 ($C=S$), 1196, 1057, 978, 905, 828, 748, 656, 627, 529 cm^{-1} . EI-MS (m/e): 146 (M^+ , 41%), 105 (3%), 91 (8%), 69 (23%), 57 (43%), 41 (100%).

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