

Scope of the Ring-Opening Metathesis Polymerization (ROMP) Reaction of 1-Substituted Cyclobutenes

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Abstract: The reactivities of a series of 1-substituted cyclobutene derivatives (carboxylate esters, carboxamides, and carbinol esters) were investigated as substrates for ring-opening metathesis polymerization (ROMP) with $[(H_2IMes)(3-Br-pyridine)_2(Cl)_2Ru=CHPh]$. Both the secondary amides of 1-cyclobutenecarboxylic acid and the esters of 1-cyclobutene-1-methanol undergo polymerization. The secondary amides provide translationally invariant polymers (*E*-olefins). Although the carbinol esters yield stereo- and regiochemically heterogeneous polymers, the 1-cyclobutenecarboxylic acid esters and tertiary amides undergo ring-opening metathesis (ROM) but not ROMP. The regio- and stereochemical outcomes of these ROMP and ROM reactions were analyzed at the B3LYP/6-31G* and LANL2DZ levels of theory. Calculations suggest that the regiochemistry and stereochemistry of the addition to the propagating carbene to form the metallocyclobutane intermediate depend on both charge distribution and steric interactions.

Introduction

A variety of strained, cyclic monomers undergo ruthenium-catalyzed ring-opening metathesis polymerization (ROMP) to provide a range of polymeric materials.^{1–7} Development of N-heterocyclic carbene-substituted ruthenium catalysts with pyridyl ligands (third-generation Grubbs catalysts),⁸ for example, **1**, provided entry into living ROMP reactions (Scheme 1). The fast initiation rate ($k_i \gg k_p$), high thermal stability, and excellent functional group tolerance of catalyst **1** have enabled the pursuit of a multitude of applications.^{6,9–13} Norbornene, oxanorbornene, and oxanorbornene-dicarboximide derivatives are the most popular ROMP monomers because of their high ring strain, easy preparation, and the facility with which functional groups are attached.¹⁴ Living ROMP of these monomers generates linear

polymers with accurate molecular weight control and low polydispersities (PDIs). However, internally heterogeneous mixtures are obtained because the polymerization reactions lack complete stereochemical and regiochemical control.

With the intention of obtaining polymers that are translationally invariant by virtue of a regioselective addition step and the absence of asymmetric centers on the backbone, we designed cyclobutenecarboxylic acid amide ROMP monomers, for example, 1-(methyl ester)glycine cyclobutenecarboxylate **2a**.¹⁰ We found that ROMP of substrate **2a** with catalyst **1** provides regioregular polymers with strictly *E*-olefin geometry and excellent PDIs (Scheme 2).¹⁰

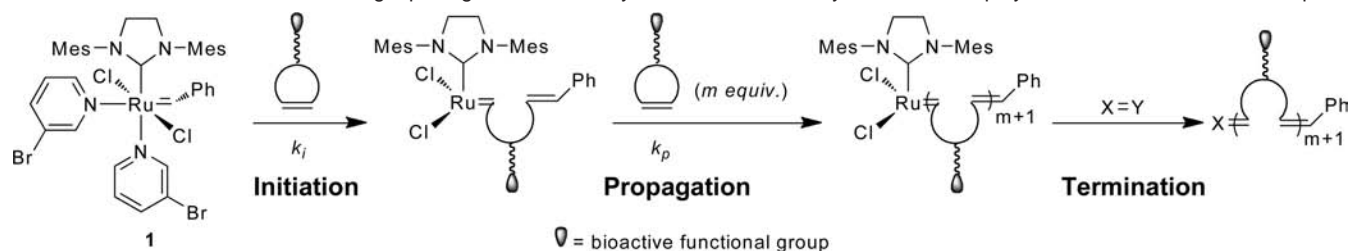
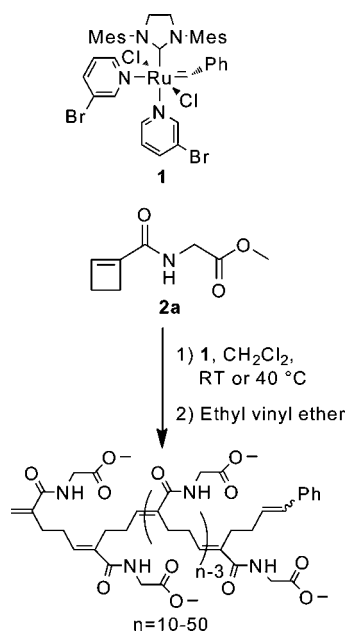
We have now investigated the reactivities of the readily available 1-substituted cyclobutenes **2b–5** (Figure 1) under ROMP conditions. All secondary amides examined (**2**) provide polymers with translationally invariant backbones and excellent PDIs, whereas ROMP of the 1-cyclobutene-1-methanol esters (**5**) is neither regio- nor stereoselective. In contrast, both 1-cyclobutenecarboxylic acid tertiary amides (**3**) and 1-cyclobutenecarboxylic acid esters (**4**) undergo only a single ring-opening metathesis cycle (ROM) without polymerization. Here, we present the kinetic data for the metathesis transformations, the computed energies of key reaction intermediates and products, and the correlation between these energies and the observed reactivities.

Results and Discussion

Preparation of Monomers. The known cyclobutene-1-carboxylic acid **6** was derivatized to provide all of the monomers (Scheme 3).^{10,15} It was converted to amides **2** by EDC coupling. Tertiary amide **3a** was prepared in useable yield by a modified EDC procedure in which DMAP was added to the reaction

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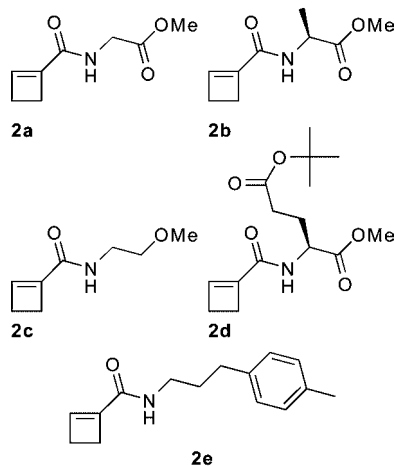
Scheme 1. General Mechanism of Ring-Opening Metathesis Polymerization with Catalyst **1** for the Display of Bioactive Functional Groups**Scheme 2.** Ring-Opening Metathesis Polymerization (ROMP) of **2a**¹⁰

mixture. Piperidine amide **3b** was obtained by a procedure based on *N,N'*-bis(2-oxo-3-oxazolidinyl)phosphinic chloride (BOP-Cl). The known methyl ester **4a** was prepared by treatment of acid **6** with *N,N'*-diisopropyl-*O*-methylisourea.^{16,17} Ester **4b** was obtained by treating the acid with methyl bromoacetate in the presence of Hunig's base. Reduction of ester **4a** with DIBAL-H gave cyclobutene-1-carbinol, which was esterified with acetyl and pivaloyl chloride to provide **5a** and **5b**, respectively.

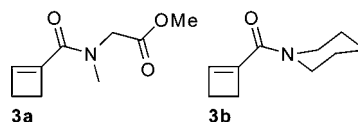
Kinetics, Regiochemistry, and Stereochemistry of 1-Substituted Cyclobutene ROMP. When a 1-substituted cyclobutene coordinates to the ruthenium center of the catalyst, there are four Ru-cyclobutane intermediates that may be formed (*cis*- and *trans*-**IM-1** and *cis*- and *trans*-**IM-2**, Scheme 4). In Pathway I, the Ru-cyclobutane ring is formed with a 1,3-relationship between the substituent-bearing carbons, whereas in Pathway II, there is a 1,2-relationship between them. Selectivity in formation of the metallocyclobutanes determines the regiochemistry and stereochemistry of chain extension (**IM** → **RP**).

1. ROMP of Secondary Amide Monomers. The ROMP kinetics of monomers **2a**–**2e** were measured by monitoring the ROMP reactions in CD₂Cl₂ at 25 °C. We have previously shown that secondary amide cyclobutenes provide at least 50-mer polymers with excellent PDIs, if required.¹⁰ Here, we studied

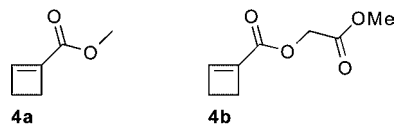
Secondary Amides



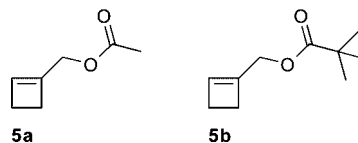
Tertiary Amides



Esters



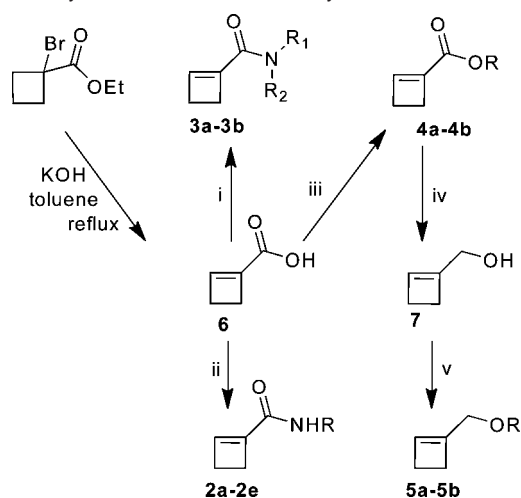
Carbinol Esters

**Figure 1.** Cyclobutene monomers subjected to ROMP.

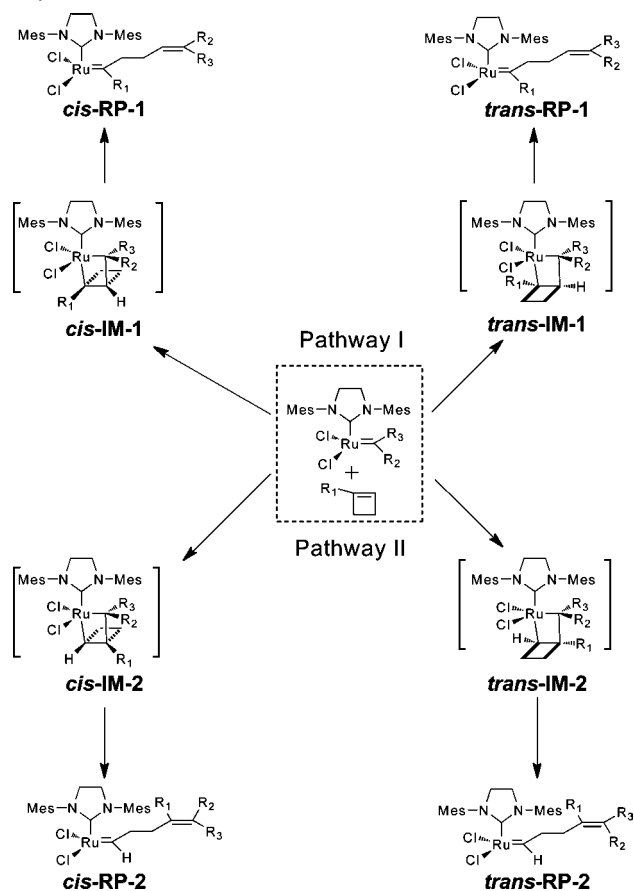
the ROMP reaction in the context of the preparation of shorter polymers so that NMR analysis would be more informative.

The ¹H NMR spectra of the ROMP reaction of monomer **2b** illustrate the method used to follow the kinetics of the reaction and the assignment of stereochemistry in the products (Figure 2). As a function of time, the integration of the peaks a, b, and c (corresponding to protons in monomer **2b**) decreased as the integration of peaks d and e (attributed to protons on the chain of the ring-opened ruthenium carbene) increased. During the course of the reaction of each monomer **2**, a single broad peak appeared at 6.2 ppm in the ¹H NMR spectrum, indicating the formation of an internal trisubstituted olefin with *E*-configuration.¹⁰ There was no evidence of disubstituted olefin (signals in the 5 to 6 ppm region). In addition, the absence of a peak at

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Scheme 3. Synthesis of 1-Substituted Cyclobutenes^a

^a (i) **3a**: H-Gly-OMe, EDC·HCl, DMAP, DIEA, CH₂Cl₂, rt; **3b**: BOP-Cl, piperidine, DIEA, CH₂Cl₂, rt. (ii) primary amines, EDC·HCl, DIEA, CH₂Cl₂, rt. (iii) **4a**: *N,N'*-diisopropyl-*O*-methylisourea, Et₂O, rt; **4b**: BrCH₂CO₂CH₃, KI, DIEA, rt. (iv) DIBAL-H, Et₂O, -78 °C. (v) Pivaloyl chloride or acetyl chloride, DIEA, DMAP, CH₂Cl₂, 0 °C to rt.

Scheme 4. Two Possible Reaction Pathways and Their Corresponding Regio- and Stereoisomeric Intermediates and Products in the Ruthenium-Catalyzed Ring-Opening Reactions of 1-Cyclobutene Derivatives^a

^a For the ROM reactions and the ROMP initiating reaction, R₂ = H, and R₃ = Ph. For the ROMP chain extending reactions, R₁ = R₂ and R₃ = the polymer chain.

18.0–19.1 ppm indicated that the ruthenium alkylidene ([Ru]=CH–CH₂R) was not formed, and implied that an amide-substituted ruthenium carbene (RP-1 in Pathway I, Scheme 4)

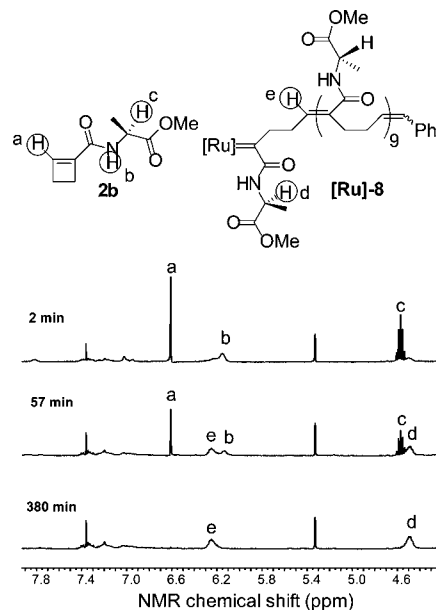


Figure 2. Time course of ROMP of monomer **2b** followed by ¹H NMR spectroscopy. Reaction conditions: CD₂Cl₂, [**2b**] = 0.1 M, [**1**] = 0.01 M, 25 °C. The peak at 5.33 ppm is CHDCl₂.

is the exclusive chain-lengthening species. The 10-mer polymers derived from secondary amides **2** are formed with high regioselectivity and stereoselectivity.

The rates of consumption of monomers **2a–2e** are very similar (*t*₅₀ entries, Table 1, Figure S1). The polymerization rates of substrates **2** are approximately 4 times slower than those of 1,2-unsubstituted, 3-substituted cyclobutenes,¹⁸ in which the olefinic bond is disubstituted and the substituents are one atom removed from the carbons that undergo metathesis.

2. ROM of Tertiary Amide Monomers. Reaction of tertiary amides **3a** and **3b** with 10 mol % catalyst **1** resulted in the ring-opening metathesis (ROM) of approximately 10 mol % of the monomer. The ring-opening reaction requires 2 h to reach this 10% conversion; no polymerization is observed (Table 1, Figure S2).

ROM of **3a** and **3b** exhibits the same regioselectivity as that of the secondary amide monomers **2**, and only resonances consistent with Pathway I were observed in the ¹H NMR spectra (Figure S3). Although secondary amides with γ -branching (i.e., **2b** and **2d**) undergo ROMP, tertiary amides, in which branching is at the β -position, yield only ring-opened monomer. We hypothesize that the increased steric bulk around the ruthenium in RP-1 hinders the binding of subsequent monomers to the tertiary amide-substituted carbene (see below).

3. ROM of Cyclobutenecarboxylic Acid Esters. We have reported that monomer **4a** undergoes ROM but not ROMP with catalyst **1**.¹⁹ Here, we describe the ROM of a second ester and the regiochemistry of the ROM reaction. As seen with monomer **4a**, incubation of ester **4b** with 10 mol % of catalyst **1** resulted in the ring-opening metathesis (ROM) of approximately 10 mol % of the monomer with no polymerization (Table 1, Figure S2); that is, the remaining 90% of the monomer does not react.

An NMR time course of the reaction of 10 equiv of monomer **4a** with 1 equiv of catalyst **1** is shown in Figure 3. The peak at

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Table 1. ROM and ROMP Results^a

monomer	[monomer]/ 1	conversion yield % ^b	rxn time/h	<i>t</i> ₅₀ /min ^c	<i>M</i> _n (theo)	<i>M</i> _n ^d	<i>M</i> _w ^e	PDI
2a	10	93	3	3	1796	1835	2220	1.21
2b	10	97	6	3	1936	1820	2522	1.39
2c	10	85	3	4	1656	1349	1653	1.23
2d	10	94	4	3	3077	3483	4796	1.38
2e	10	96	6	3	2397	2222	3047	1.37
3a	10	10	2	∞ ^f	n/a	n/a	n/a	n/a
3b	10	10	2	∞ ^f	n/a	n/a	n/a	n/a
4a	10	10	3	∞ ^f	n/a	n/a	n/a	n/a
4a ^g	1	78	21	87	n/a	n/a	n/a	n/a
4b	10	10	3	∞ ^f	n/a	n/a	n/a	n/a
4b ^g	1	72	31	333	n/a	n/a	n/a	n/a
5a	10	98	1.5	<2	1366	2057	2740	1.33
5b	10	99	1.5	<2	1786	2067	2726	1.33

^a Reaction conditions: CD₂Cl₂, 25 °C, [monomer] = 0.1 M, [**1**] = 0.01 M. ^b Determined by ¹H NMR spectroscopy. ^c Reaction time for 50% consumption of monomer. ^d Number-averaged molecular weight by GPC using polystyrene standards. ^e Weight-averaged molecular weight by GPC using polystyrene standards. ^f The reaction stops after 10% consumption of monomer. ^g [**1**] = 0.1 M.

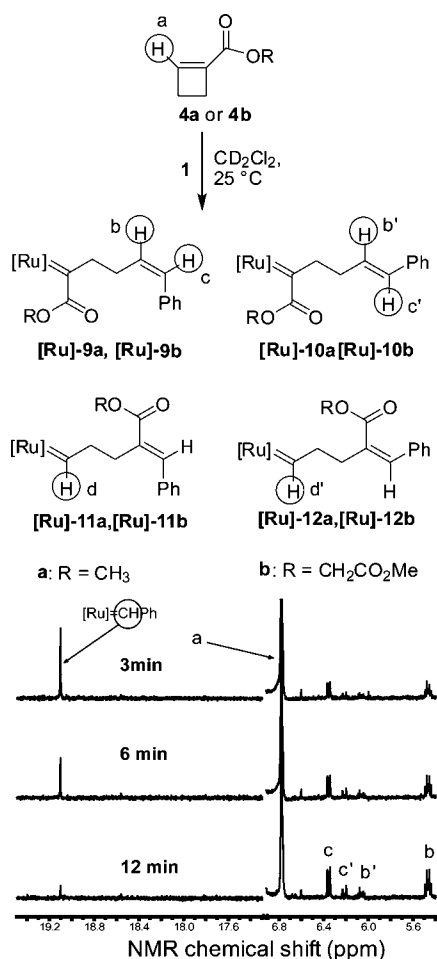


Figure 3. Time course of ROM of monomer **4a** followed by ¹H NMR spectroscopy. Reaction conditions: CD₂Cl₂, [**4a**] = 0.1 M, [**1**] = 0.01 M, 25 °C. Resonances corresponding to H (d) and H (d') are predicted to occur at 18.5 ppm. These resonances are not observed.

19.1 ppm corresponding to the carbene proton on the catalyst **1** disappears over time and no new carbene proton peak appears. At the same time, the styrenyl peaks from the ring-opened monomer appear (peak b, b', c, and c'). Therefore, **RP-1** is formed preferentially to **RP-2**, as is observed for both secondary and tertiary amides. The 1-cyclobutene ester undergoes a single ring-opening reaction with the alkylidene catalyst to form an enonic carbene that does not react with remaining 1-cyclobutene

ester. We conclude that the enonic ruthenium carbene, **RP-1** cannot undergo further metathesis with an unsaturated ester.

To determine the kinetics of the ring-opening and to analyze the structures of the products in greater detail, we examined the reaction of esters **4a** and **4b** with 1 equiv of catalyst (Figure S4). Again, the NMR spectra were monitored as a function of time. The ring-opening reactions of monomers **4a** and **4b** required more than 20 h to reach 75% consumption of monomer. The low reaction rates suggest that an ester further reduces the reactivity of the cyclobutene olefin relative to the amide-substituted cyclobutenes. In contrast to the secondary amides, these esters have no β- or γ-branching and are analogous in structure to primary amides **2**. These comparisons suggest that the failure of esters **4a** and **4b** to polymerize is the result of an electronic factor rather than steric congestion at the catalytic center.

There are four possible ROM products that may be formed from monomer **4a** or **4b** (Figure 3). As noted above, ROM of 1-substituted esters was regioselective. The primary products [Ru]-**9a** and [Ru]-**10a** are formed as a 1/2 mixture of *Z/E* styrenyl olefins (Figure 4a) through Pathway I (Scheme 4). The quenched reaction products were purified and characterized by ¹H NMR spectroscopy (Figure S6) and LC-MS (APCI) spectroscopy to establish their structures. A minor peak at 18.5 ppm in the ¹H NMR spectrum (Figure S5) was assigned to the carbene proton of species [Ru]-**11a** and [Ru]-**12a** (via Pathway II). It is reasonable to assume that [Ru]-**11a** and [Ru]-**12a** initiate ROM with monomer **4a**; the resulting carbenes would be enonic and stable to further reaction prior to quenching. The time course (Figure 4a) revealed that the ruthenium carbenes [Ru]-**11a** and [Ru]-**12a** are formed in the initial reaction and are not secondary products from cross metatheses.

High ROM regioselectivity was also observed with monomer **4b**. The ROM products of **4b** formed through the enonic carbene (Pathway I, Scheme 4) are likewise a mixture of *Z* and *E* styrene olefins ([Ru]-**9b** and [Ru]-**10b**) with a *Z/E* ratio around 2/3 (Figure 4b). At 75% consumption of monomer, the combined yield of [Ru]-**11b** and [Ru]-**12b** generated through formation of the alkylidene carbene (Pathway II, Scheme 4) is less than 4% (data not shown). Thus, Pathway I (Scheme 4) to form the enonic carbene is the energetically favorable pathway.

4. ROMP and ROM of Carbinol Esters 5. Metathesis reactions with 1 equiv of catalyst **1** and 10 equiv of monomer **5a** or **5b** were monitored by ¹H NMR spectroscopy at room temperature for 1.5 h (Table 1, Figure S7). Greater than 98% of each monomer was consumed in 1 h. The *t*₅₀ values are at

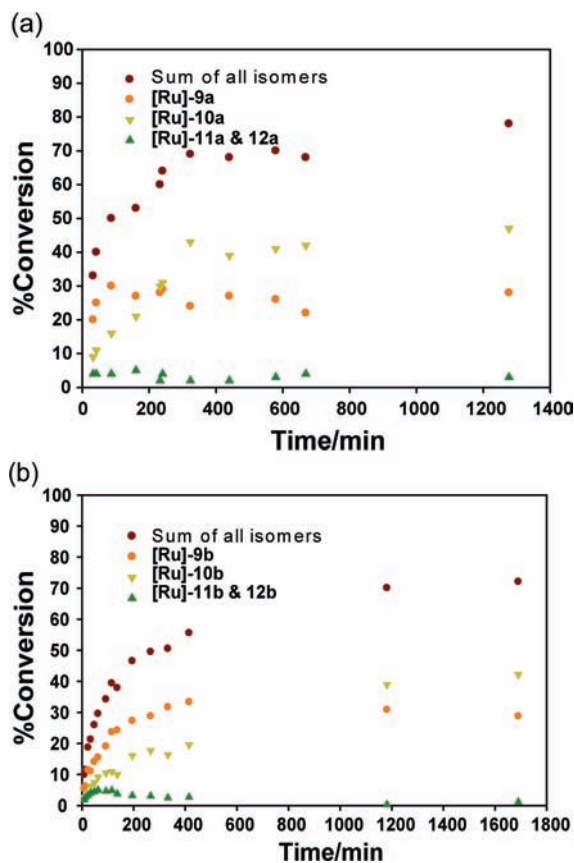


Figure 4. Rate of formation of ring-opened monomer products formed from monomers **4a** and **4b**. Reaction conditions: [**4a**] or [**4b**] = 0.1 M, [**1**] = 0.1 M, 25 °C.

least 5 times smaller than the t_{50} values for monomers **2–4**; the carbinol esters **5** undergo ROMP much faster than secondary amides **2**.

In the ^1H NMR spectrum of the [Ru]-**13** polymer, the methylene protons adjacent to the ester oxygen are observed as three broad signals centered at 4.64, 4.57, and 4.46 ppm of approximately equal intensity (Figure 5, peak d and Figure S8, peaks f, g, and h, Table S1). The assignment of these resonances to the allylic oxygen-bearing carbon was corroborated by ^{13}C NMR, ^{13}C -APT, gCOSY, and gHMQC spectroscopy of the purified 10-mer products **13** and **14** (Table S1, Figure S8). The presence of three signals indicates that at least three, and perhaps all four (if peaks are overlapping) of the possible substitution patterns on ($-\text{CH}_2\text{OR}$)-bearing double bonds are present (Figure 6). Furthermore, the ^1H NMR spectrum of the [Ru]-**13** polymer exhibits multiple broad resonances for the olefinic protons in the polymer backbone at 5.35–5.62 ppm (Figure 5, peak c), indicating that the polymer backbone is composed of different types of olefins (Figure 6). Therefore, the polymerization is neither regio- nor stereoselective.

Computational Analysis of Regio- and Stereoselectivity. To understand the effects of structure on reactivity in each of the ring-opening reactions, we analyzed the energies of the stable and metastable species along the two polymerization pathways by *ab initio* methods. The isomer formed depends on which step along the reaction coordinate is rate-determining. The rate of formation of the 14-electron ruthenium complex upon dissociation of bromopyridine from **1** is the same regardless of cyclobutene structure and will not contribute to the regio- or stereoselectivities. Likewise, during the catalytic cycle, competi-

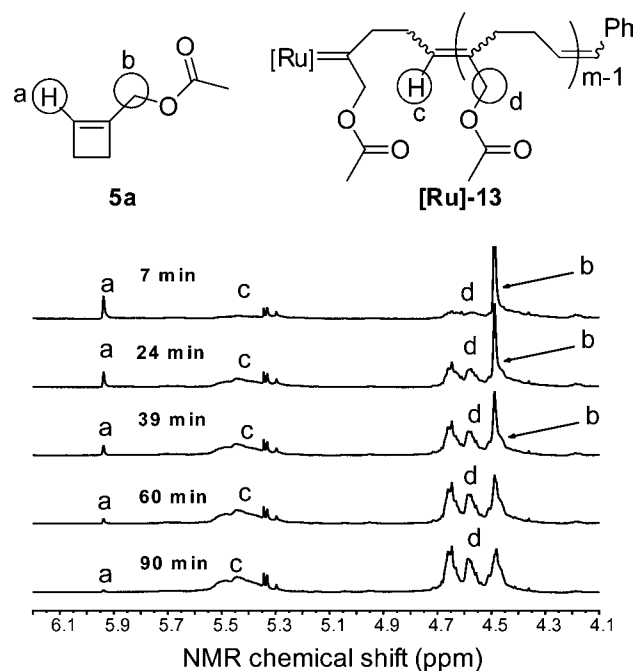


Figure 5. Time course of ROMP of monomer **5a** followed by ^1H NMR spectroscopy. Reaction conditions: CD_2Cl_2 , [**5a**] = 0.1 M, [**1**] = 0.01 M, 25 °C.

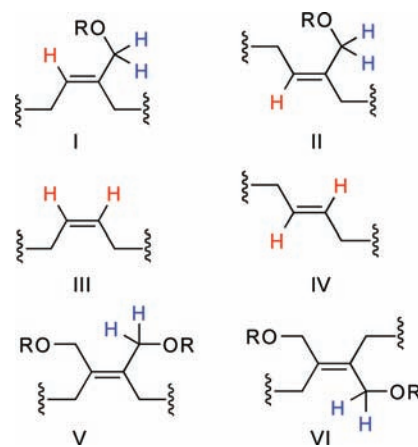


Figure 6. Six possible alkene configurations in the polymer backbone upon ROMP of monomers **5**.

tion between the cyclobutene and the bromopyridine does not alter the regio- or stereoselectivity.

Therefore, two kinetic steps after dissociation were considered: (a) formation of the metallocyclobutane intermediate, and (b) formation of ring-opened product. To understand the regioselectivity of π -complex (π -SM) formation, we calculated the natural bond orbital charge populations for the cyclobutene monomers and their respective propagating carbenes. For the polymerization of cyclobutene amide **2a**, we considered reaction with the amide substituted carbene, which is the species formed in the reaction; for the polymerization of the carbinol ester **5b**, we considered only the reaction with the substituted carbene that is the analogous isomer (Figure 7a). For the ROM monomers, **3** and **4**, we analyzed the initial reaction with the benzyldiene carbene **1** (Figure 7a). To predict the relative activation energies for metallocyclobutane formation and subsequent ring-opening, we calculated the energies of the reaction intermediates (**IM**) and products (**RP**).

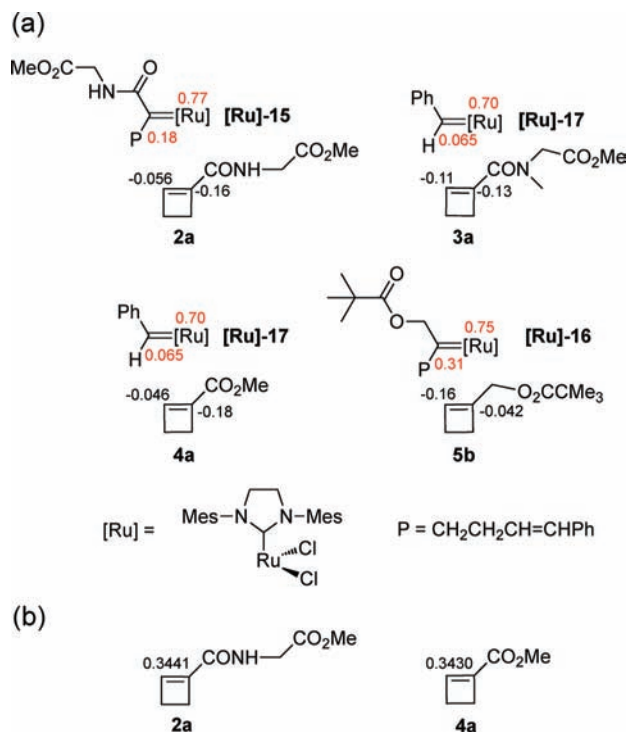


Figure 7. (a) NBO charge populations of 1-substituted cyclobutene derivatives **2a**, **3a**, **4a**, and **5b**, the corresponding ring-opened ruthenium carbenes of **2a** and **3a**, and ruthenium benzylidene. Hartree–Fock calculations were performed with the 6-311G++* basis set (for cyclobutene monomers) and the LANL2DZ basis set (for ruthenium carbenes) in Gaussian 03W. (b) AIM electron densities (atomic units) for the olefin bonds in 1-substituted cyclobutene derivatives **2a** and **4a** were calculated with AIMPAC.

1. Calculation Methods. For optimization of the geometries of the cyclobutene monomers (**2a**, **3a**, **4a**, and **5b**), we used B3LYP/6-31G*. This method is frequently employed for simple organic compounds^{20–23} because it provides accurate structural predictions with economical calculations. To limit the number of atoms included in the propagating ruthenium carbenes and thereby facilitate the calculations, we modeled the substituents as (CH₂CH₂CH=CHPh); that is, we used structures [Ru]-15 and [Ru]-16. For these carbenes, and for the ruthenium benzylidene [Ru]-17 (Figure 7a), we used B3LYP/LANL2DZ. This method has been used extensively for the geometry optimization of ruthenium carbenes. The LANL2DZ basis set has superior properties with respect to effective core potentials (ECPs) and polarization functions on the ruthenium^{24–37} and calculated results agree very well with experimental data, for example, X-ray crystal structures,^{32,33} IR and NMR spectra,³⁶ thermal parameters,^{25,26,37} or metathesis reactivities.^{24,25,28–31,34} For natural bond orbital (NBO) charge populations (Figure 7a), we used Hartree–Fock with the 6-31G++* basis set (for cyclobutene monomers) and the LANL2DZ basis set (for ruthenium carbenes) in Gaussian 03W.

The geometry optimizations and energy calculations of the intermediates (**IM**) and products (**RP**) were also performed with B3LYP/LANL2DZ in Gaussian 03W. To simplify the computation, the polymer chain and the 1-carbonyl amide group were modeled as a methyl group and a CONHCH₃ substituent, respectively (Figure S9). To prevent ring-opening and to find the local minimum of the intermediate structures, only one bond in the metallocyclobutane ring was optimized while the other three bond lengths were kept constant; the partial optimization was rotated to optimize the other three bonds, until the optimized structure changed little in energy ($\Delta E < 6.3 \times 10^{-4}$ kcal/mol). Vibrational frequency calculations using B3LYP/LANL2DZ were performed for all model compounds (Figure S9); there were no imaginary vibrational frequencies indicating that local minima had been found for each structure. Free energies computed in Gaussian 03W for structures (Figure 8) in solvent (CH₂Cl₂) include the electronic energy plus the solvation free energy from the conductor-like polarizable continuum model (CPCM) based on the United Atom Kohn–Sham (UAKS) radii.

2. NBO Charge Calculations. For the ester and amide cyclobutene monomers, the calculations showed that the electron density is higher on C-1 than on C-2 as expected for olefins bearing conjugated electron withdrawing groups (monomers **2a**, **3a**, **4a**). In contrast to the carbonyl substituents, the C-1 methanol ester substituent (monomer **5b**) is electron donating, making C-1 more electropositive than C-2. In all of the ruthenium carbenes, the metal atom is more electropositive than the adjacent carbon atom.

3. Relative Energies of Intermediates and Products. The NBO calculations allowed us to rationalize the regiochemical results of the ROM and ROMP processes. In the case of monomers **2–4**, alignment of the cyclobutene and the electron-deficient Ru to pair C-1 and the Ru to form the π -complex (π -SM) is favored by the electronic factors; however, in the case of monomers **5**, electronic factors favor a cycloaddition in which C-2 becomes bonded to the ruthenium (Figure 7). Assuming then that steric effects between the coordinating cyclobutene and the carbene are relatively unimportant (because of the long distance between the cyclobutene and Ru in the π -complex), we ordered the energies of the regioisomeric cyclobutene π -complexes (π -SM) according to the charge distribution of the cyclobutenes.

Noting energy values from the literature for the ring strain of cyclobutenes and for π -stabilization of ruthenium carbene

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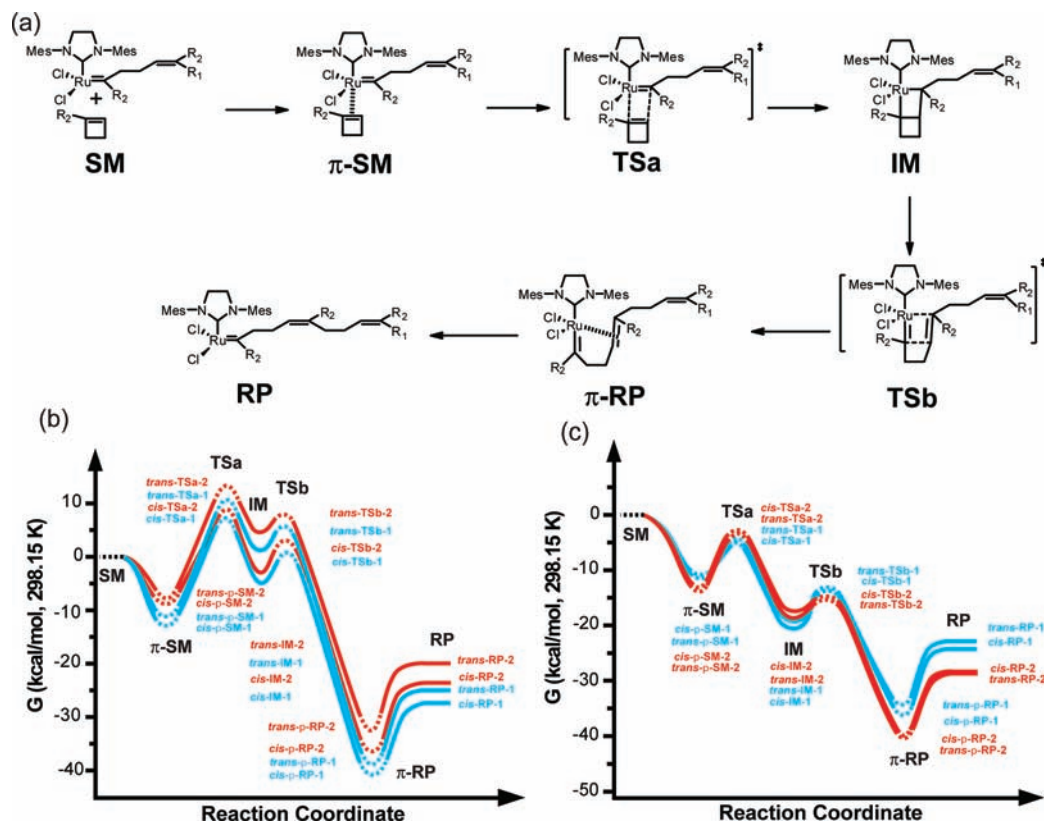


Figure 8. (a) Cyclobutene ROMP reaction scheme and relative free energy profiles for the reactions of (b) monomers **2**, $R_2 = \text{CONHMe}$, and (c) monomers **5**, $R_2 = \text{CH}_2\text{OAc}$. Structures were optimized with B3LYP/LANL2DZ in Gaussian 03W program. Free energies computed for structures in solvent (CH_2Cl_2) at 298.15 K include the electronic energy plus the solvation free energy from the CPCM solvation model based on the UAKS radii using Gaussian 03W. **SM**, starting material; π -**SM**, π -complex of starting material; **TS**, transition state; **IM**, intermediate; π -**RP**, π -complex of reaction product; **RP**, reaction product. In the potential minima and maxima, dashed lines represent energy estimates and solid lines represent calculated energies.

complexes, we developed a comprehensive picture (Figure 8) of the ROMP processes of **2** and **5**. On the basis of the calculations of Goddard and co-workers,³⁸ we estimated the strain energy of 1-substituted cyclobutene to be ~ 26 kcal/mol. We used the strain energy to plot the difference in energy between the starting 14 electron ruthenium carbene (**SM**) energy and the mean product carbene (**RP**) energy. The calculated energy differences between the metallocyclobutane intermediates (**IM**) and the product carbenes (**RP**) were used to order the isomers on the free energy profile. For the ROMP of **5**, we plotted the energies for the reaction that begins with the more substituted of the two regioisomeric ruthenium carbenes. Further, we assumed that the most favorable π -stabilized 16-electron ruthenium carbene (π -**SM**) is about 14 kcal/mol more stable than the uncomplexed carbene (**SM**) and that the most favorable π -**RP** is 14 kcal/mol more stable than the corresponding **RP**. An 18 kcal/mol stabilization energy was calculated by Zhao and Truhlar for ethylene³⁹ and measured by Chen and co-workers for norbornene;⁴⁰ however, there are additional steric interactions present in cyclobutene π -**SM** complexes (Figure

8). The **TSa** energies were ordered according to the relative stability of their corresponding metallocyclobutane intermediates (**IM**) to account for steric effects that develop in the transition state. All of the species were placed along the reaction coordinates of monomers **2** and **5** in their respective free energy profiles (Figure 8).

4. The *cis*-Pathway I Is Favored for Secondary Amide Monomers. In the case of 1-substituted cyclobutene amides, starting (π -**SM**), intermediate (**IM**), and reaction product (**RP**) complexes on the *cis*-pathway I are always the lowest energy isomers. By extension, the transition states are expected to be the lowest energy transition states and the corresponding activation energies for their formation the most favorable. This picture is consistent with the high selectivity for ring-opening metathesis of amides **2** (>99%) of the more substituted ruthenium carbene and (*E*)-olefin,¹⁰ and by extension for ROM of monomers **3** and **4**.

5. Activation Energies in Pathways I and II Are Nearly Equivalent for Carbinol Ester Monomers 5. In the case of the carbinol esters **5**, if we consider the partitioning of metallocyclobutane intermediate forward to ring-opened product versus backward to the coordinated cyclobutene, the forward reaction is favored due to the release of ring strain upon formation of π -**RP**, and π -**SM** is not in equilibrium with **IM**. Likewise by the Hammond postulate, we surmise that **TSb** is not rate-limiting. Thus, the energy of **TSa** determines the regio- and stereoselectivities. The NBO charge and energy calculations suggest that *trans*- π -**SM-2** and *cis*- π -**SM-2** are lower in energy than the π -**SM-1** regioisomers. However, upon formation of the

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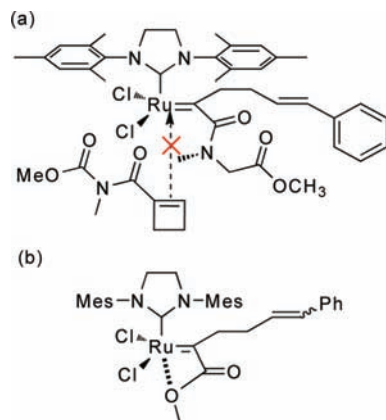


Figure 9. (a) Schematic model based on the B3LYP/LANL2DZ optimized structure (Gaussian 03W) of the new N,N-disubstituted carbonyl ruthenium carbene for monomer **3a** illustrating how the second N-substituent blocks access to the ruthenium carbene of a second monomer. (b) Model of the ester coordination observed in the B3LYP/LANL2DZ optimized structure (Gaussian 03W) of the enoic carbene formed from monomer **4a**.

metallocyclobutane intermediate, steric interactions destabilize **IM-2** relative to **IM-1** counteracting electronic effects. As these steric interactions develop in the transition state, the difference in **TSa** energies between isomers is much smaller than in either the π -**SM** or **IM** species. The consequence is that *trans*- and *cis*- π -**SM-1** react at a comparable rate to *trans*- and *cis*- π -**SM-2**. Thus, poor regio- and stereoselectivity are observed. In addition, the overall estimated activation energies for carbinol esters **5** are smaller than for those of the secondary amides **2**; this result is consistent with the higher rates of propagation observed for monomers **5** (Table 1).

6. Geometry of ROM Intermediate from Tertiary Amide Monomers 3. Kinetic studies with monomers **3a** and **3b** revealed that only ROM reactions occurred. We hypothesized that the second N-substituent may block the approach and binding of an incoming monomer at the ruthenium carbene center. To test our hypothesis, geometry optimization of the N,N-disubstituted carbonyl ruthenium carbene formed from monomer **3a** was performed with B3LYP/LANL2DZ in Gaussian 03W. In the optimized structure of the ruthenium carbene, the N-methyl group blocks the face of the metal carbene to which the incoming cyclobutene must bind (Figure 9a). Therefore, we conclude that severe steric crowding is responsible for a lack of propagation after the initial ring-opening occurs.

7. Inability of Ester Monomers 4 To Form Homopolymer. Our initial hypothesis for the lack of polymerization of enoic carbene was that the electron withdrawing nature of the ester deactivated the cyclobutene olefin to further metathesis reaction. However, the NBO calculations followed by AIM electron density analysis reveal that the charge density on the ester-substituted cyclobutene is not significantly different than the

charge density on the secondary amide-substituted cyclobutene, which does undergo ROMP (Figure 7). *Ab initio* calculations for the enoic carbene reaction product of **4a** revealed that the ester oxygen forms a chelate at the open coordination site of the 14-electron ruthenium center (Figure 9b). This chelation stabilizes the ester-substituted carbene by approximately 7 kcal/mol relative to the amide-substituted carbene. Similar ester coordination was observed in calculations by which Fomine and co-workers investigated the reactivity of the enoic carbene³⁵ and in the reaction of enoates in cross metathesis.⁴¹ Although ring-opening metathesis with the relatively electron-rich olefin of cyclohexene^{19,42,43} is still feasible, the chelated enoic carbene is kinetically trapped from further reaction with the sterically hindered 1-substituted cyclobutene ester, **4**.

Conclusion

In summary, we have found that cyclobutenes undergo stereo- and regio-regular ring-opening metathesis when substituted with an electron withdrawing carbonyl at the 1-position. Regio-regular addition to the catalyst carbene is consistent with both the calculated charge distributions for the carbene and the cyclobutene and the minimization of steric interactions. In the case of more electron-rich 1-substituents, the inverse rank order of energies between the intermediates and the π -complexed starting materials strongly suggests that the corresponding activation energies to reach the transition states may be close enough in magnitude to make ROMP of carbinol ester monomers **5** neither regio- nor stereoselective.

Neither cyclohexene carboxylic acid esters nor tertiary amides ROMP. When an enoic carbene is formed from ester monomers **4**, the formation of a ruthenium chelate with the ester oxygen traps the enoic carbene in a stabilized state that precludes the reaction of these esters with 1-substituted cyclobutene esters, for example, in homopolymerizations. This reactivity opens an avenue for preparing alternating polymers with unique functionality.¹⁹ Steric crowding prevents propagation for tertiary amide substituents. In conclusion, of the functionalized cyclobutenes studied, the secondary amides **2** exhibit the optimal level of reactivity and stereo- and regio- control for the generation of translationally invariant polymers.

Acknowledgment. This research was supported by NIH grants R01HD38519 (N.S.S.), S10RR021008 (N.S.S.), and GM074776 (K.A.P.), NYSTAR award (FDP C040076, N.S.S.) and NSF grant CHE0131146 (NMR). We thank Dr. James Marecek for his assistance with NMR spectroscopy and Dr. Derek Middlemiss for his consultation on calculations.

Supporting Information Available: Additional figures, experimental methods and results including spectroscopic data and coordinates of calculated structures. This material is available free of charge via the Internet at <http://pubs.acs.org>.

JA1037098