

Z-Selective Ethenolysis with a Ruthenium Metathesis Catalyst: Experiment and Theory

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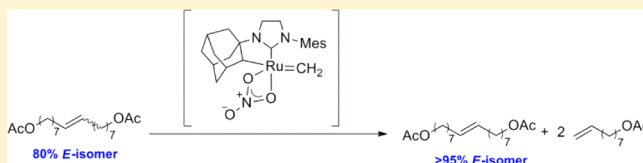
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S Supporting Information

ABSTRACT: The Z-selective ethenolysis activity of chelated ruthenium metathesis catalysts was investigated with experiment and theory. A five-membered chelated catalyst that was successfully employed in Z-selective cross metathesis reactions has now been found to be highly active for Z-selective ethenolysis at low ethylene pressures, while tolerating a wide variety of functional groups. This phenomenon also affects its activity in cross metathesis reactions and prohibits crossover reactions of internal olefins via trisubstituted ruthenacyclobutane intermediates. In contrast, a related catalyst containing a six-membered chelated architecture is not active for ethenolysis and seems to react through different pathways more reminiscent of previous generations of ruthenium catalysts. Computational investigations of the effects of substitution on relevant transition states and ruthenacyclobutane intermediates revealed that the differences of activities are attributed to the steric repulsions of the anionic ligand with the chelating groups.



INTRODUCTION

The discovery of transition metal alkylidene catalysts has allowed olefin metathesis to permeate the literature in a wide variety of fields including green chemistry,¹ natural product synthesis,² and polymer chemistry,³ since its discovery in the 1950s. The development of catalysts that exhibit preference for kinetically versus thermodynamically controlled products, such as the generation of terminal olefins from internal olefins, termed ethenolysis, and the formation of Z-olefins in cross metathesis (CM) reactions, is a particular challenge in the field.

For a metathesis catalyst to be active in ethenolysis reactions, it must exhibit high activity and stability as a propagating methylidene. However, many known metathesis catalysts are unstable as methylidene complexes and undergo rapid decomposition, thus exhibiting poor ethenolysis reactivity.⁴ The desired ethenolysis catalytic cycle is depicted in Scheme 1. Initial reaction of an internal olefin with a metal methylidene proceeds via a 1,2-metallacycle and produces a terminal olefin and the corresponding substituted metal alkylidene. Further reaction with ethylene forms a second equivalent of terminal olefin and regenerates the catalytically active methylidene.

For an ethenolysis catalyst to show high selectivity at appropriate ethylene pressures, formation of terminal olefin products must be favored over back reactions and side reactions that produce internal olefins (Scheme 1). Side reactions that reduce selectivity for the desired terminal olefin products include self-metathesis and secondary metathesis. Self-metathesis

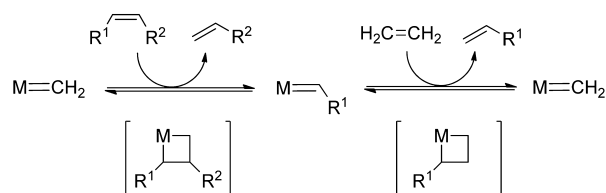
is when a metathesis reaction occurs between two substrate molecules instead of between a substrate molecule and ethylene, and secondary metathesis involves the CM of two terminal olefins to generate an internal olefin and ethylene (Scheme 1). Industrially, the ethenolysis of seed oil derivatives affords chemically desirable products with applications in cosmetics, detergents, polymer additives, and renewable biofuels.⁵

The ability to selectively form the kinetically preferred Z-olefin products in CM reactions is another significant challenge in metathesis research, as catalysts have generally been observed to favor formation of the thermodynamic E-isomer. The Z-olefin motif is prevalent in a variety of small molecules, including many natural products and pharmaceutical targets. The first example of a catalyst-controlled system capable of predominantly forming the Z-isomer in CM reactions was reported by the Hoveyda and Schrock laboratories. The Z-selectivity of the reported tungsten and molybdenum catalysts was attributed to the difference in the size of the two axial ligands. This size difference influences the orientation of the substituents on the forming metallacyclobutane intermediate and leads to productive formation of Z-olefins.⁶ These catalysts have shown great utility in the synthesis of complicated natural products and stereoregular polymers.⁷ A particular Z-selective

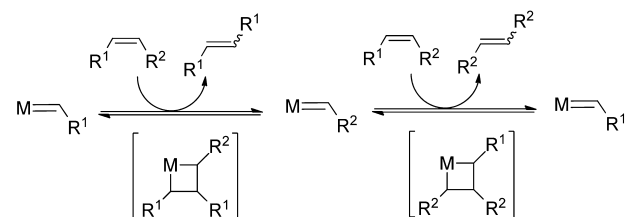
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Scheme 1. Ethenolysis and Related Side Reactions

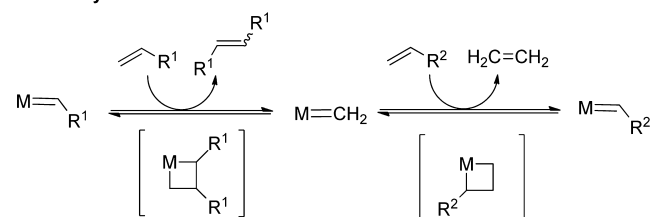
Ethenolysis Reaction



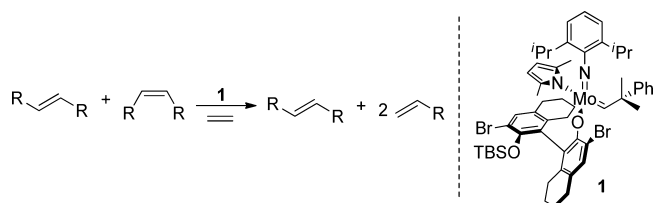
Self-Metathesis



Secondary Metathesis



molybdenum catalyst (**1**) was shown to be effective for the *Z*-selective ethenolysis of internal olefins.⁸ In this process, the corresponding molybdenum methylidene reacts preferentially with *cis*-olefins to produce two terminal olefins, while *trans*-olefins react to a significantly smaller extent (Scheme 2). Since

Scheme 2. *Z*-Selective Ethenolysis Reaction Using Molybdenum Catalyst **1**

ethenolysis is the reverse of cross metathesis, the same 1,2-disubstituted metallacyclobutane complex must be formed as an intermediate in both reactions. Hence, if a catalyst is highly *Z*-selective when forming cross products, it is expected to also be able to selectively degrade *cis*-olefins by ethenolysis, assuming that the corresponding metal methylidene complex is stable.

A family of functional group tolerant *Z*-selective ruthenium-based catalysts has recently been reported that contain a chelating *N*-heterocyclic carbene (NHC) ligand derived from an intramolecular carboxylate-driven C–H bond insertion of an *N*-bound substituent (Figure 1).⁹ Catalyst **3** is derived from C–H activation of the benzylic position of the *N*-mesityl substituent of complex **2** and thus contains a six-membered chelated structure that imparts slightly improved *Z*-selectivity compared to previous generations of ruthenium catalysts. Catalysts **4** and **5** contain five-membered chelates derived from

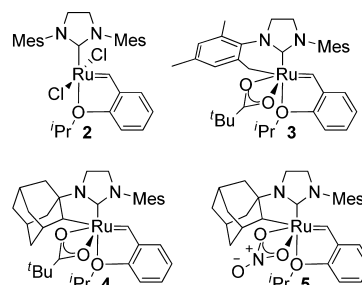


Figure 1. Prominent ruthenium metathesis catalysts (Mes = 2,4,6-trimethylphenyl).

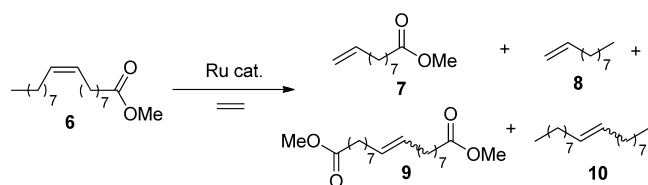
C–H activation of an *N*-adamantyl substituent, and exhibit activity and *Z*-selectivity rivaling the aforementioned group IV systems. The identity of the anionic ligand has been found to have great consequence on reactivity and selectivity, as replacement of a carboxylate on **4** for a nitrate-type ligand (**5**) results in greater stability, *Z*-selectivity, and close to 1000 turnovers for homodimerization reactions. *Z*-selective CM, macrocyclic ring closing metathesis (RCM), and ring-opening metathesis polymerization (ROMP) reactions have been reported using this family of chelated catalysts.¹⁰

Our groups and others have used density functional theory (DFT) calculations to elucidate important information about the mechanism of action, origin of *Z*-selectivity, and stability of chelated ruthenium catalysts **4** and **5**.¹¹ The metathesis reaction occurs via a side-bound mechanism, different from that with nonchelated ruthenium catalysts. The olefin approaches *cis* to the NHC ligand on the catalyst.¹² The *N*-adamantyl chelating group positions the *N*-mesityl substituent directly over the forming metallacyclobutane, thus causing its substituents to be oriented away to avoid steric repulsions and leading to high *Z*-selectivity of the metathesis products.

To design better catalysts for *Z*-selective metathesis, a more thorough understanding of this family of chelated ruthenium catalysts is required. The goal of this study is to explore the *Z*-selectivity of these catalysts for ethenolysis reactions and concurrently investigate how this can help us better understand their CM reactivity. The stability and structure of metallacyclobutane intermediates greatly influences metathesis reactivity and selectivity; thus, we sought to study the effects of substitution on relevant ruthenacyclobutane intermediates using experimental and theoretical techniques. Both catalysts **3** and **5** were tested so that the effects of chelate size could be investigated.¹³ Herein, we report a method for the functional group tolerant *Z*-selective ethenolysis of internal olefins and explore other unique reactivity of chelated ruthenium catalysts, providing hypotheses of observed behavior based on ruthenacyclobutane stability and structure.

RESULTS AND DISCUSSION

Ethenolysis Experimental Investigations. The ethenolysis activity of chelated ruthenium complexes **3** and **5** was initially investigated in order to directly compare our chelated catalysts to previously reported catalysts.¹⁴ We first explored their activity and selectivity for the ethenolysis of the completely *cis*-olefin substrate, methyl oleate (Table 1). The ethenolysis of methyl oleate is a standard assay used to compare ethenolysis reactivity and selectivity of metathesis catalysts. It should be noted that selectivity here refers to the formation of the desired ethenolysis products, terminal olefins **7** and **8**, and

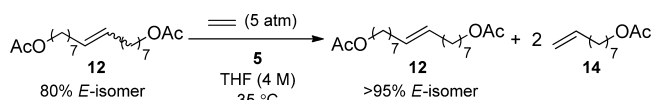
Table 1. Ethenolysis Reactions of Methyl Oleate Catalyzed by Catalysts 3 and 5

entry ^a	catalyst	mol %	yield ^b	selectivity ^c	TON ^d
1	3	0.1	0%	-	0
2	3	0.01	0%	-	0
3	5	0.1	80%	>95%	800
4	5	0.01	12%	>95%	1160

^aThe reactions were run in a minimal amount of CH₂Cl₂ for 1 h at 40 °C and 10.2 atm of ethylene. ^bYield = (moles of ethenolysis products 7 + 8) × 100%/(initial moles of 6). ^cSelectivity = (moles of ethenolysis products 7 + 8)/(moles of total products 7 + 8 + 9 + 10). ^dTON = yield × [(initial moles of 6)/(moles of catalyst)].

not to the catalyst's *E/Z* selectivity. Although catalyst 3 showed no reactivity at the catalyst loadings tested, catalyst 5 was able to catalyze the transformation with high turnovers and high selectivity at low loadings.¹⁵ The fact that 5 is highly active as an ethenolysis catalyst and previously exhibited high *Z*-selectivity in CM reactions strongly suggests that it would exhibit high selectivity for *Z*-olefins in ethenolysis reactions.

We tested chelated catalysts 3 and 5 in the ethenolysis of ~4:1 mixtures of the *trans*- and *cis*-isomers of two internal olefins, 5-decene and the acetate-substituted substrate 12, to determine if these catalysts exhibited any selectivity for *Z*-olefins. We were pleased to find that under the optimized conditions depicted in Scheme 3, catalyst 5 was able to enrich

Scheme 3. Z-Selective Ethenolysis Reaction of Substrate 12**Table 2. E-Isomer Enrichment by Z-Selective Ethenolysis of Various Functionalized Symmetrical Internal Olefins with the Formula R(CH₂)_nCH=CH-(CH₂)_nR Using Catalyst 5**

entry	compound	R; n	initial %E	mol % 5	pressure (atm)	time (h)	final %E
1	11	CH ₃ ; 3	79	0.5	1	4	90
2			79	0.5	5	4	>95
3			52	0.5	5	4	90
4	12	OAc; 7	78	0.5	1	4	93
5			78	0.5	5	4	>95
6	15	OH; 4	82	0.5	1	4	92
7			82	0.5	5	4	>95
8			68	0.5	5	4	90
9	16	CO ₂ Me; 6	80	0.5	1	6	88
10			80	0.5	5	6	>95
11	17	NHPh; 3	80	0.5	1	4	92
12			80	1.0	5	4	>95
13			60	1.0	5	6	86
14	18	C(O)Me; 2	72	0.5	1	4	90
15			72	0.5	5	4	>95

both internal olefin mixtures (~80% *E*) to >95% of the *E*-isomer at 5 atm of ethylene and 0.5 mol % catalyst loading for the two substrates (Scheme 3); the products of both reactions were recovered by flash column chromatography.¹⁶ For 5-decene, the purely *E*-isomer (>95% *E*) was isolated in 90% yield based upon initial *E*-content.¹⁷ For 12, the purely *E*-internal olefin (>95% *E*) and 8-nonenyl acetate produced by ethenolysis of the *Z*-olefins were both quantitatively recovered (Scheme 3). Exposure of catalyst 3 (0.5 mol %) to a ~4:1 mixture of the *trans*- and *cis*-isomers of 12 at 5 atm of ethylene led to a very small amount of ethenolysis (<3% conversion) and no observable selectivity. Additionally, only olefin migration of the starting material 12 was observed when the reaction was carried out at 1 atm of ethylene at higher loadings of catalyst 3 (5 mol %).

With the knowledge that 5 was an effective *Z*-selective ethenolysis catalyst, we sought to investigate its functional group compatibility in *E*-isomer enrichment reactions.¹⁸ A variety of functional groups were tolerated under the reaction conditions, including acetates, alcohols, esters, amines, and ketones (Table 2). Reaction performed with 5 atm of ethylene for all *E*-dominant substrates (72–82% *E*) led to enrichment of the internal olefins with >95% of the *E*-isomer as monitored by ¹H NMR. Although the same reactions performed under 1 atm of ethylene proceeded with high selectivity, reaction at 5 atm was necessary to push the ~80% mixtures to >95% of the *E*-isomer.

We next attempted to quantify the ethenolysis selectivity of catalyst 5 by investigating the relative rates of degradation of *E* and *Z* internal olefins using ¹H NMR spectroscopy. 5-Decene was chosen as a substrate since the stereopure *E*- and *Z*-isomers are commercially available. The rate of ethenolysis was found to be first-order in substrate⁸ and the relative rates of 5-decene ethenolysis were determined under 1 atm of ethylene (see Supporting Information). Neither the ethenolysis of *Z*-5-decene nor *E*-5-decene proceeded to completion under the reaction conditions.¹⁹ Nevertheless, log plots of substrate concentration versus time at early reaction times were found to be linear (*E*-5-decene, *R*² = 0.93; *Z*-5-decene, *R*² = 0.98). From

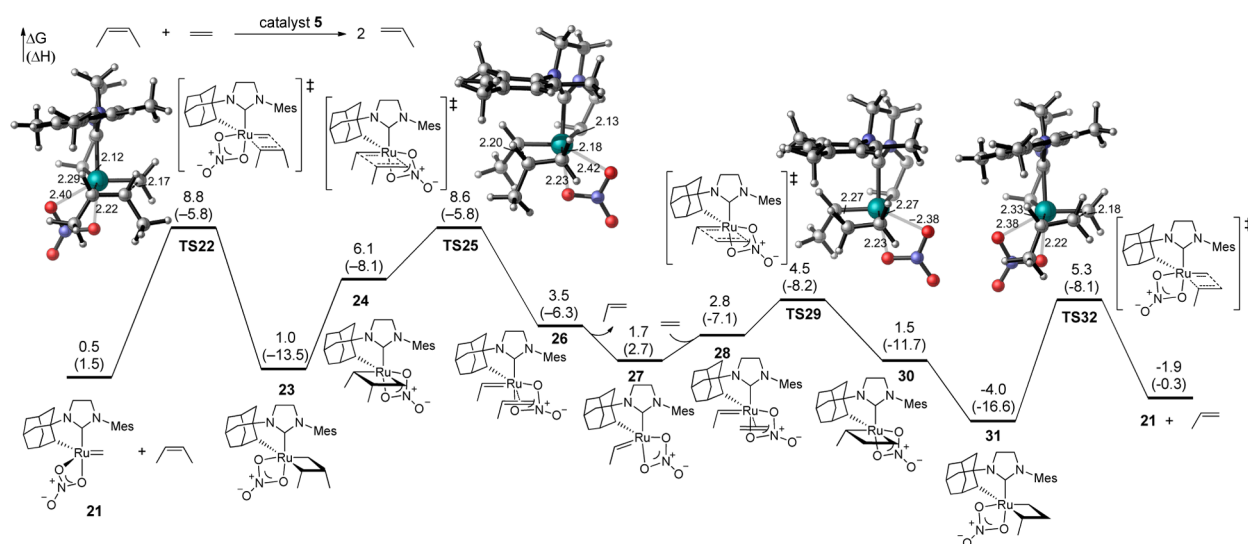


Figure 2. The most favorable pathway of ethenolysis of *cis*-2-butene with catalyst 5. Gibbs free energies and enthalpies (in parentheses) are in kcal/mol and with respect to the most stable ruthenium ethylidene complex 47 (an isomer of 27, see Figure 4). For clarity, the chelating adamantyl group is not shown in the 3D transition state structures.

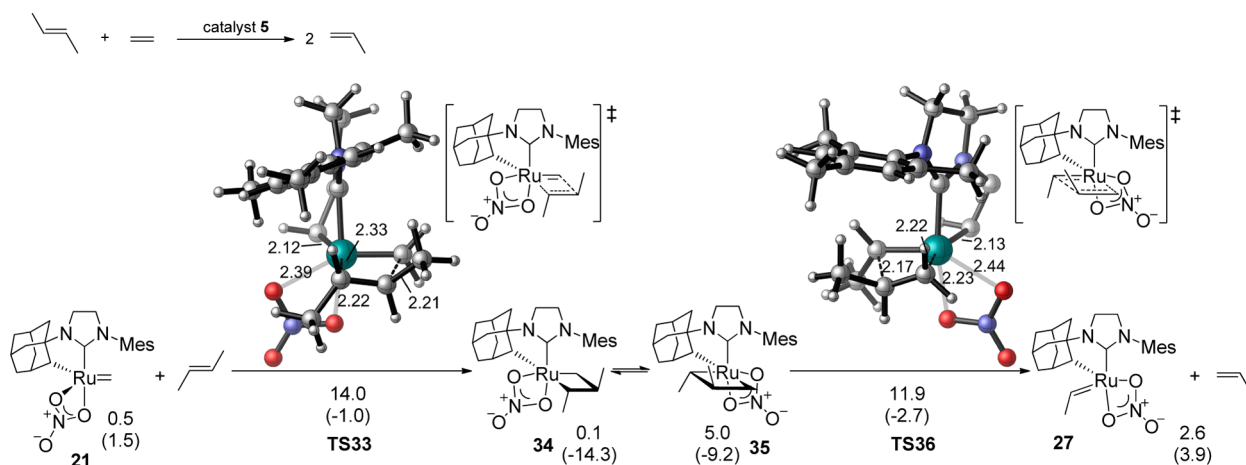


Figure 3. The most favorable pathway of ethenolysis of *trans*-2-butene with catalyst 5. Gibbs free energies and enthalpies (in parentheses) are in kcal/mol and with respect to the most stable ruthenium ethylidene complex 47 (an isomer of 27, see Figure 4). For clarity, the chelating adamantyl group is not shown in the 3D transition state structures. In subsequent steps, 27 reacts with ethylene to regenerate 21. This is identical to the second half of the catalytic cycle in the reaction with *cis*-2-butene (see Figure 2).

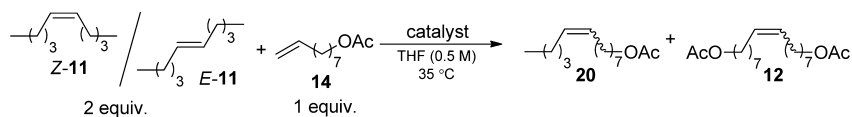
the slopes of these plots, the ratio of the rate constants for ethenolysis of *Z*-5-decene and *E*-5-decene (k_Z/k_E) was found to be ca. 4.5. The corresponding k_Z/k_E value reported for molybdenum catalyst 1, 30 ± 5 , is significantly higher, implying that catalyst 1 is inherently more selective than 5. However, all reactions catalyzed by 1 were conducted under high ethylene pressures not suitable for benchtop reactions (4–20 atm).²⁰ The functional group tolerance of catalyst 5 and the *Z*-selectivity at ethylene pressures as low as 1 atm highlights advantages of this particular ruthenium-based system for the preparation of terminal olefins from internal olefins and for the purification of *E/Z* mixtures.²¹ The further development of chelated catalysts with increased *Z*-selectivity will lead to ruthenium catalysts with increased k_Z/k_E values.²²

Ethenolysis Computational Investigations. To understand the mechanism of ethenolysis, and the origin of *Z*-selectivity with catalyst 5, we computed the ethenolysis reaction pathways and the *Z/E*-selectivity with density functional theory (DFT). The calculations were performed using Gaussian 09²³

with a theoretical level found to be satisfactory in our previous computational studies of chelated ruthenium catalysts.¹¹ Geometries were optimized in the gas phase with B3LYP²⁴/LANL2DZ–6-31G(d). Single point calculations were performed with M06²⁵/SDD-6-311+G(d,p) and the SMD²⁶ solvation model with THF solvent.

Reaction pathways initiated from both ruthenium methylidene and alkylidene complexes were investigated, since these interconvert during the ethenolysis reaction. The most favorable pathway of the ethenolysis of *cis*-2-butene with catalyst 5 involves the side-bound approach of the internal olefin to the ruthenium methylidene complex 21 (Figure 2). Formation of the ruthenacyclobutane intermediate 23 requires an activation free energy of 8.8 kcal/mol (TS22). In 21, TS22, and 23, the nitrate is *syn* to the α -H on the chelating adamantyl group. Ruthenacyclobutane 23 isomerizes to form a less stable ruthenacycle 24, in which the nitrate is *anti* to the adamantyl α -H.²⁷ Cleavage of the ruthenacycle 24 via TS25 requires a comparable activation energy as TS22 ($\Delta G^\ddagger = 8.6$ kcal/mol),

Table 3. Internal–Terminal Cross Metathesis Reaction of 5-Decene (11) and 8-Nonenyl Acetate (14) Catalyzed by 3 or 5



entry	catalyst	mol %	substrate	time (h)	yield of 20 ^a	yield of 12 ^a	% Z of 20 ^b	% Z of 12 ^b
1	5	0.2	Z-11	6	57%	21%	91%	83%
2	5	0.2	E-11	24	<1%	19%	-	87%
3	3	2.5	Z-11	2	69%	14%	23%	22%
4	3	2.5	E-11	2	53%	17%	25%	33%

^aDetermined by gas chromatography. ^bDetermined by ¹H NMR.

and generates a ruthenium–propene π complex **26**. In contrast, productive cleavage of **23** without isomerization to **24** requires a much higher barrier ($\Delta G^\ddagger = 20.9$ kcal/mol) and forms an unstable ruthenium ethylidene complex in which the ethylidene is *trans* to the chelating Ru–C bond. Thus, the isomerization to **24** is necessary before cleaving the ruthenacyclobutane. Decoordination of propene from **26** yields ruthenium ethylidene **27**, which then binds to an ethylene molecule to form π complex **28**. Subsequent steps involve the formation and cleavage of monosubstituted ruthenacyclobutane intermediates via **TS29** and **TS32**, respectively, and eventually regeneration of the ruthenium methylidene complex **21**. The monosubstituted transition states in the second half of the catalytic cycle (**TS29** and **TS32**) are both 3–4 kcal/mol more stable than the disubstituted transition states **TS22** and **TS25**. Thus, the reaction of ruthenium methylidene with the internal olefin is the rate-limiting step in the catalytic cycle (**TS22**), while the productive cleavage of the disubstituted metallacycle (**TS25**) requires essentially identical activation energy.

The anionic nitrate ligand binds bidentate to the ruthenium in all four transition states in the catalytic cycle, although the Ru–O bond *trans* to the alkylidene is significantly longer than the Ru–O bond *trans* to the NHC (~ 2.4 versus ~ 2.2 Å). The transition states with monodentate nitrate are five-coordinated with trigonal bipyramidal geometries and 1–4 kcal/mol less stable than the corresponding bidentate transition states (see Supporting Information). The small energy differences between mono- and bidentate nitrate complexes suggest that the monodentate transition structures might become favorable with bulkier olefin substituents and/or bulkier anionic ligands. Structures containing both mono- and bidentate binding modes are considered in the following computations and only the most favorable structures are shown.

In the analogous reaction with *trans*-2-butene, both transition states **TS33** and **TS36** are less stable than the corresponding transition states **TS22** and **TS25** in the reaction with *cis*-2-butene (Figure 3). In **TS33** and **TS36**, one of the olefin substituents is pointing toward the *N*-mesityl group and leads to significant steric repulsions. The overall activation barrier is 5.2 kcal/mol higher than the ethenolysis of *cis*-2-butene. This explains the observed *Z*-selectivity in ethenolysis reactions.²⁸ Interestingly, the ruthenacyclobutane intermediates **34** and **35** in the reaction with *trans*-2-butene are slightly more stable than the corresponding *cis*-substituted metallacycles **23** and **24**. Unlike the metathesis transition states, in which the olefin and the Ru–alkylidene are almost in the same plane, the four-membered metallacycle intermediates are puckered. The methyl substituents on the metallacycles in **34** and **35** are not directly pointing toward the *N*-mesityl group on the ligand (see Supporting Information for the 3D structures of the

metallacycle intermediates). The ligand–metallacycle repulsions in the metallacycle intermediates are smaller than in the transition states.

Crossover Experimental Studies. Research presented in a previous report led us to believe that ethenolysis plays a major role in CM reactions catalyzed by **5**.^{10c} The CM reaction between a *cis*-internal olefin and a terminal olefin was monitored over time and revealed that internal olefins must be broken down by ethenolysis before a heterocross product can be generated; it was proposed that the required methylidene complex was generated by homodimerization of the terminal olefin substrate. In addition to this, no crossover was observed when two *cis*-internal olefins were reacted in the presence of catalyst **5**, and it was suggested that this was due to high steric demands associated with forming trisubstituted ruthenacycles using this particular catalyst.²⁹ Since CM between two internal olefins is a common occurrence for previous generations of metathesis catalysts including molybdenum-based *Z*-selective catalysts,⁶ we desired to further probe this unique reactivity of catalyst **5**. Previously, species **20** was synthesized by reacting the two terminal olefins 1-hexene and 8-nonenyl acetate in the presence of catalyst **5** (0.5 mol %) and proceeded with high yield (67%) and *cis*-selectivity (91% *Z*-olefin). We explored whether catalysts **5** and **3** were able to form substrate **20** from (1) an internal olefin and a terminal olefin, or (2) two internal olefins. Catalysts **5** and **3** were both investigated in order to elucidate differences in reactivity, activity, and selectivity between the complexes with different chelate sizes.

The reaction of 5-decene (**11**) and 8-nonenyl acetate (**14**) to form compound **20** was initially probed (Table 3). When catalyst **5** was employed, use of the *cis*- and *trans*-isomers of 5-decene greatly affected its metathesis activity (entries 1 and 2). Reaction with *cis*-5-decene led to formation of **20** with 57% yield and 91% *Z*-isomer at 0.2 mol % of **5**. The analogous reaction under the same conditions with *trans*-5-decene led to only trace amounts of **20**. In both cases, the undesired homodimer of 8-nonenyl acetate (compound **12**) was also formed in similar quantities and *Z*-selectivities, regardless of the isomer of 5-decene used. Conversely, catalyst **3** was able to form compound **20** with both isomers of 5-decene (entries 3 and 4). The % *Z* values of **20** and **12** were notably low compared to the reactions catalyzed by **5**; however, this is attributed to extensive *Z/E* isomerization by secondary metathesis processes at the long reaction times.

The unique behavior of catalyst **5** gives important insight into the reactivity of this chelated catalyst.³⁰ Since **5** is effective for the *Z*-selective ethenolysis of internal olefins, its inability to react with *trans*-olefins in CM reactions further suggests that all internal olefins must undergo ethenolysis first to generate

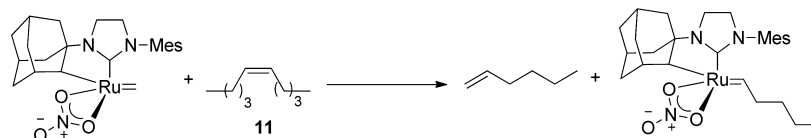
Scheme 4. Reaction of a Methylidene with *cis*-5-Decene To Produce 1-Hexene and the Corresponding Substituted Alkylidene

Table 4. Internal–Internal Cross Metathesis Reaction of 5-Decene (11) with 12 Catalyzed by 3 or 5

entry	catalyst	mol %	substrate	ethylene exposure ^a	time (h)	yield of 20 ^b	yield of 14 ^b	% Z of 20 ^c	% Z of 12 ^c
1	5	1.0	Z-11	-	24	<1%	<1%	-	76%
2	5	1.0	E-11	-	24	<1%	<1%	-	75%
3	5	1.0	Z-11	+	4.5	21%	8%	95%	70%
4	5	1.0	E-11	+	24	<1%	2%	-	76%
5	3	2.5	Z-11	-	2	37%	<1%	31%	46%
6	3	2.5	E-11	-	2	30%	<1%	31%	60%

^a+ = 1 atm of ethylene was introduced into the headspace of the reaction vessel for 2 h prior to reaction. ^bDetermined by gas chromatography. ^cDetermined by ¹H NMR.

terminal olefins, and the productive CM reaction occurs between two terminal olefin molecules. In the reaction shown in Table 3 catalyzed by 5, it is proposed that a methylidene is initially formed by homodimerization of 8-nonenyl acetate (14) to form 12. This methylidene can then react with *cis*-5-decene to form 1-hexene and the corresponding substituted alkylidene (Scheme 4), both of which can react further with the terminal olefin 8-nonenyl acetate to generate cross product 20. Since 8-nonenyl acetate must initially be homodimerized for productive CM to occur, larger amounts of 12 will be generated compared to other catalysts, as was observed. It is also interesting to note that because catalyst 3 is not particularly active as an ethenolysis catalyst, CM reactions catalyzed by 3 seems to proceed through different pathways that are more similar to previous generations of ruthenium catalysts, like 2. To further test these hypotheses, the reactions of two internal olefins in the presence and absence of ethylene were attempted.

Exposure of catalyst 5 to a mixture of the internal olefins 5-decene (11) and 12 (75% Z) under the conditions shown in Table 4 led to no formation of cross product 20 regardless of which isomer of 5-decene was employed (entries 1 and 2).^{31,32} However, addition of 1 atm of ethylene into the headspace of the reaction vessel for 2 h followed by stirring for 4.5 h did lead to formation of product 20 with *cis*-5-decene (entry 3). No crossover was observed under these conditions when the *trans*-isomer was used (entry 4). This again supports the hypothesis that productive CM reactions involving internal olefins first proceed via Z-selective ethenolysis. In contrast, under the same conditions depicted in Table 4, catalyst 3 catalyzes the CM of two internal olefins in the absence of ethylene regardless of which isomer of 5-decene is employed (entries 5 and 6). Thus, CM with catalyst 3 proceeds through a completely different pathway and with this catalyst, trisubstituted ruthenacyclobutane intermediates are accessible. The low *E/Z* ratio is again attributed to extensive *Z/E* isomerization by secondary metathesis processes as evidenced by the degradation of 12 from 75% to 46% of the *Z*-isomer.

Crossover Computational Studies. We employed computations to determine the activation energies to form

and cleave the di- and trisubstituted ruthenacycle intermediates involved in reactions catalyzed by chelated catalysts 5 and 3, and the nonchelated catalyst 2. We first investigated the metathesis reactions of two internal *cis*-olefins with catalysts 2, 3, and 5.³³ To simplify the calculations, we used trimethyl substituted ruthenacyclobutanes (i.e., the reaction of ruthenium ethylidene with *cis*-2-butene) in the calculations as a model of the long conformationally mobile substrates used experimentally. Figure 4 shows the reactions of alkylidenes formed from catalysts 2, 3, and 5 with *cis*-2-butene. In these reactions, *trans*-2-butene is formed preferentially with catalyst 2, and catalysts 3 and 5 give *cis*-2-butene product.

Reaction with the unchelated catalyst 2 (Figure 4a) forms *trans*-olefin product via the bottom-bound mechanism, i.e., the olefin approaches *trans* to the NHC ligand.^{34,12} In the reactions with chelated catalysts 3 and 5, the most favorable pathway involves the side-bound mechanism (Figure 4b,c).¹¹ The activation barrier of the reaction catalyzed by 5 is 5.6 and 3.7 kcal/mol higher than that with catalysts 2 and 3, respectively. The activation energy of the rate-determining transition state TS51 is 14.1 kcal/mol with respect to the ruthenium alkylidene complex 47. In the reactions with catalysts 2 and 3, the activation energies are 8.5 and 10.4 kcal/mol, respectively (TS40 and TS45). The overall barrier of the reaction with catalyst 5 is likely to be higher than 14.1 kcal/mol, since the catalyst resting state may be more stable than the energy zero in the calculations (47). This suggests that formation of trisubstituted ruthenacyclobutanes with catalyst 5 is much more difficult than with catalysts 2 and 3, in agreement with the observed low crossover reactivities of 5 (Table 4).

The low crossover reactivity of catalyst 5 is attributed to one particular trisubstituted transition state, TS51, which is 4.4 kcal/mol less stable than the other trisubstituted transition state TS48. In TS51, the ethylidene is *syn* to the α -hydrogen on the chelating adamantyl group, while in TS48 the ethylidene is *anti* to the α -hydrogen. Interestingly, TS51 is the only transition state involving a nitrate ligand bound monodentate among all the transition states investigated in this study. Its bidentate isomer TS51' is 0.8 kcal/mol less stable, which is in contrast to

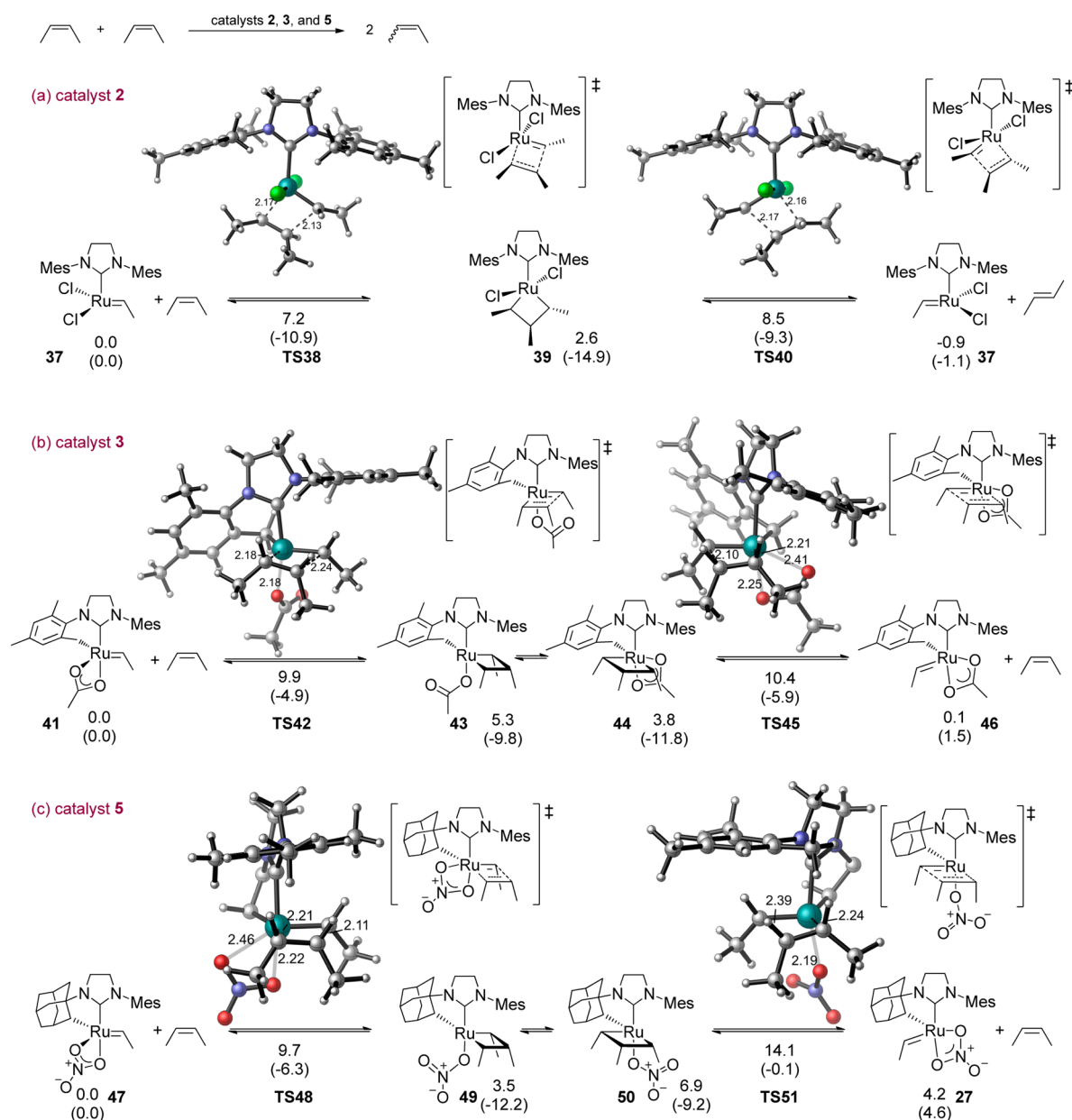


Figure 4. Reactions of ruthenium ethylidene complexes with *cis*-2-butene. These are the rate-determining steps in the metathesis of two *cis* internal olefins in the absence of ethylene. Free energies and enthalpies (in parentheses) are given in kcal/mol with respect to the ruthenium alkylidene complexes (37, 41, and 47, respectively). For clarity, the chelating adamantyl group is not shown in the 3D structures of TS48 and TS51.

other bidentate nitrate transition states that are typically 3 kcal/mol more stable than corresponding monodentate nitrate TS. To better illustrate the steric interactions with the nitrate ligand, “side-views” of the di- and trisubstituted transition states with catalyst **5** are shown in Figure 5. In the high energy trisubstituted transition state **TS51'**, the nitrate is located between the bulky chelating adamantyl group and one of the methyl substituents on the olefin. The distances between the nitrate and the olefin and between the nitrate and the adamantyl group are both significantly shorter than the sum of the van der Waals radii (the N–H distances are 2.57 and 2.51 Å, respectively, compared to the sum of van der Waals radii of N and H, 2.75 Å). The steric repulsions of the anionic nitrate ligand with the chelating adamantyl group and the substituent on the olefin clearly destabilize the bidentate transition state **TS51'** and force the trisubstituted reaction to

proceed via a generally less favorable monodentate transition state (**TS51**). As described earlier, the isomerization of the metallacyclobutane intermediate is necessary for a productive turnover. In the other trisubstituted metathesis transition state **TS48**, the nitrate is located on the less crowded side that is *syn* to the α -adamantyl hydrogen. In **TS48**, the nitrate–adamantyl and nitrate–olefin distances are both longer than corresponding distances in **TS51'**. With the diminished steric repulsions with the nitrate, **TS48** is 5.4 kcal/mol more stable than **TS51'**. Thus, the rate-limiting step in the trisubstituted catalytic cycle is **TS51**.

In the presence of ethylene, ruthenium methylidene complexes are formed by the reaction of ethylene and ruthenium alkylidenes. The reaction of ruthenium methylidene **21** with *cis*-2-butene proceeds through disubstituted transition states **TS22** and **TS25** (Figure 2). **TS22** and **TS25** are both

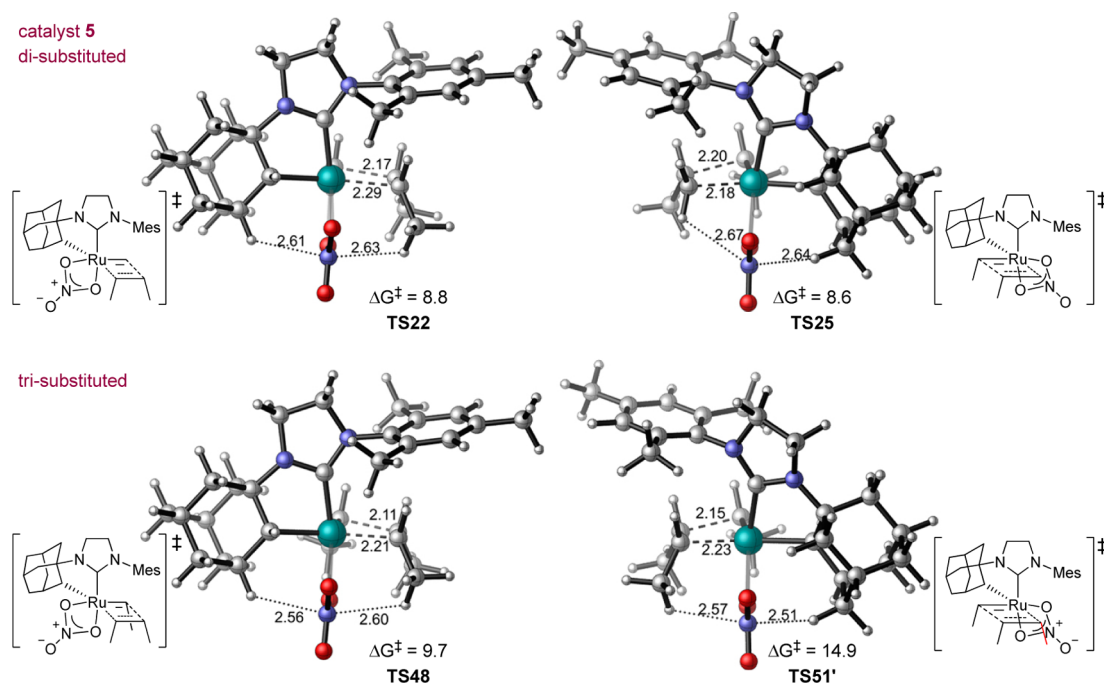


Figure 5. Side view of the dimethyl substituted transition states (TS22 and TS25) and the trimethyl substituted TS48 and TS51'. TS51' is destabilized due to steric repulsions of the nitrate with the chelating adamantyl group and the methyl group on the olefin.

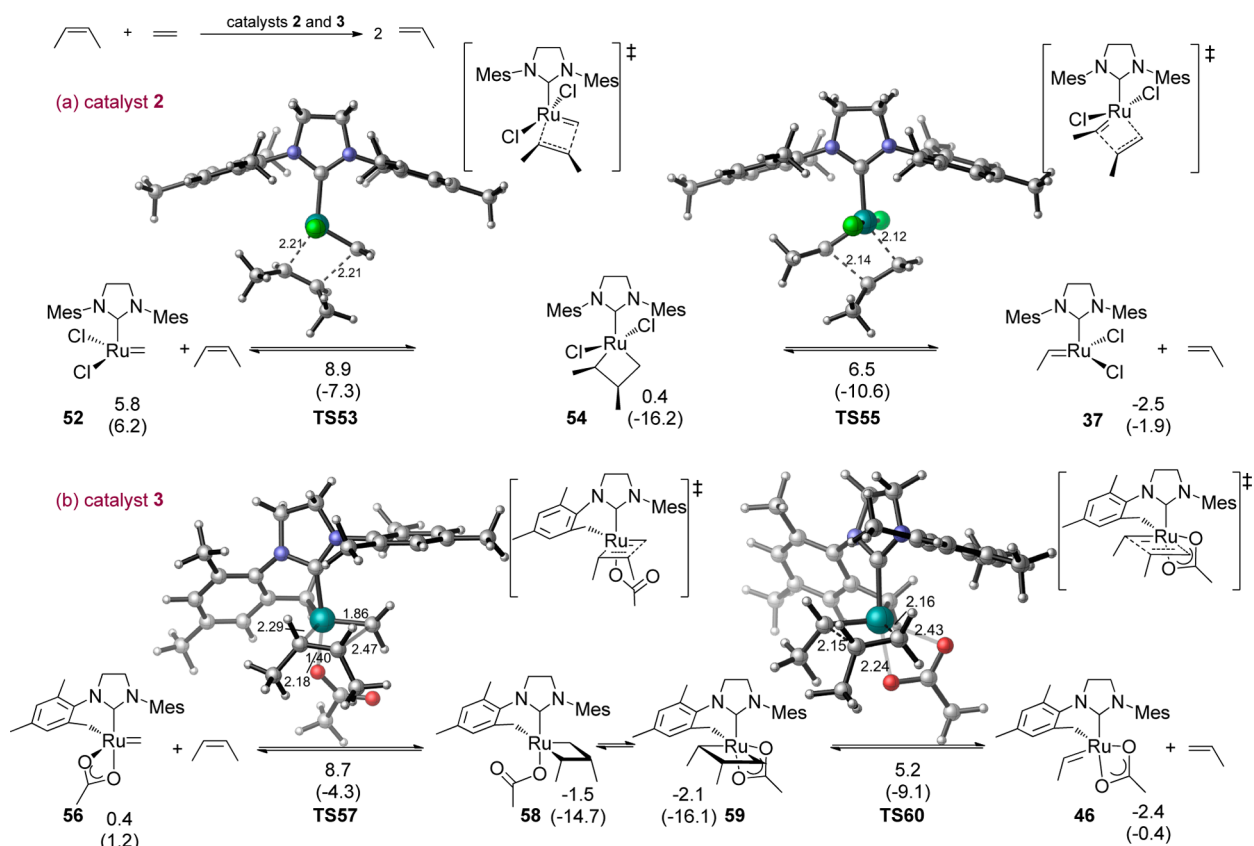


Figure 6. Free energies and enthalpies (in parentheses) of the reaction of ruthenium methylenide complexes with *cis*-2-butene catalyzed by (a) catalyst 2 and (b) catalyst 3. These are the rate-determining steps in the ethenolysis of internal olefins. All energies are with respect to the ruthenium ethylenide complexes (37 and 41, see Figure 4) and are given in kcal/mol. See Figure 2 for the reaction catalyzed by catalyst 5.

more stable than the corresponding trisubstituted transition states: TS22 is only 0.9 kcal/mol more stable than TS48 because of smaller steric repulsions of the internal olefin with

the methylenide than with the ethylenide (Figure 5). Replacing the methyl substituent with hydrogen, TS25 is dramatically stabilized by 6.3 kcal/mol compared to TS51', due to

alleviation of steric repulsions with the nitrate (see Figure 5 for direct comparison of the structures of di- and trisubstituted TS). The activation energy of the reaction of *cis*-2-butene with this methylidene is 5.3 kcal/mol lower than the corresponding pathway to form the trisubstituted ruthenacycle ($\Delta G^\ddagger = 8.8$ kcal/mol, TS22 compared to 14.1 kcal/mol, TS51). Thus, in the presence of ethylene, crossover products are formed, as was observed experimentally, from the reaction of internal olefins with the methylidene complex of catalyst 5 rather than with corresponding alkylidene complexes.

The trisubstituted metathesis pathways with catalysts 2 and 3 were also calculated and shown in Figure 4a,b. In contrast to the high activation barrier of the trisubstituted pathway with catalyst 5, catalysts 2 and 3 both have lower activation barrier in the trisubstituted reaction ($\Delta G^\ddagger = 8.5$ and 10.4 kcal/mol, respectively, compared to $\Delta G^\ddagger = 14.1$ kcal/mol with 5). This is attributed to less steric demand in the trisubstituted transition states with these catalysts; the chelating mesityl group in catalyst 3 is less bulky than the chelating adamantyl in 5. With the unchelated catalyst 2, the olefin approaches from the bottom, *trans* to the NHC ligand and thus there are no unfavorable ligand–substrate steric repulsions in the transition states.

As a comparison with the disubstituted pathway of catalyst 5 (Figure 2), we also computed the activation barriers of the reactions of *cis*-2-butene and the ruthenium methylidenes derived from catalysts 2 and 3 (Figure 6, panels a and b, respectively). For the unchelated catalyst 2, the reaction of *cis*-2-butene with methylidene 52 requires an activation energy of 8.9 kcal/mol. This is slightly higher than the barrier in the reaction with corresponding ethylidene 37 ($\Delta G^\ddagger = 8.5$ kcal/mol, Figure 4a). This is attributed to fact that ruthenium methylidene 52 is 5.8 kcal/mol less stable than corresponding ethylidene 37 as well as the absence of unfavorable ligand–substrate steric repulsions in the trisubstituted transition states. The reaction of *cis*-2-butene with the methylidene complex of catalyst 3 (56) requires a slightly lower activation energy than corresponding ethylidene (41) ($\Delta G^\ddagger = 8.7$ kcal/mol compared to 10.4 kcal/mol). With both catalysts 2 and 3, the differences between di- and trisubstituted activation barriers are within 1–2 kcal/mol. This suggests that the rate of crossover of internal olefins will not be significantly affected by the exposure to ethylene.

To complete the computational investigations, another two scenarios involving disubstituted ruthenacyclobutanes, homodimerization and nonproductive metathesis of terminal olefins, were also computed and the detailed results are provided in the Supporting Information. The homodimerization pathways of propene to form *E*- or *Z*-2-butene with catalysts 2, 3, and 5 are shown in Figure S1, and the competing nonproductive reactions of propene and ruthenium ethylidenes are shown in Figure S2.³⁵ The computations predicted that the nonproductive equilibration with catalysts 2 and 3 both requires lower activation barrier than the corresponding productive homodimerization pathway. In contrast, with catalyst 5, the nonproductive pathway requires 1.5 kcal/mol higher activation energy than homodimerization (12.7 kcal/mol compared to 11.2 kcal/mol, see Supporting Information for details). Similar to the trisubstituted transition state, the 1,3-disubstituted nonproductive transition state is destabilized by steric repulsions with the nitrate, which is also located between the adamantyl and the methyl substituent on the olefin.

CONCLUSION

In summary, we have investigated the ethenolysis behavior of a new class of ruthenium metathesis catalysts, 3 and 5, containing chelating NHC ligands. Catalyst 5 was found to catalyze *Z*-selective ethenolysis reactions at low ethylene pressures (1–5 atm) with substrates containing a wide variety of functional groups. DFT calculations showed that the *Z*-selectivity in ethenolysis reactions catalyzed by 5 is a result of steric effects that prohibit *E*-olefins from productively reacting with the corresponding methylidene. Two internal olefins could not undergo CM in the presence of 5 to form cross products but must first react with ethylene to form terminal olefins. In addition to this, no crossover was observed when *trans*-internal olefins were employed as substrates in CM reactions. This implies that the *Z*-selective ethenolysis behavior of 5 plays a large role not only in its ethenolysis reactivity, but also in its CM reactivity. In contrast, catalyst 3 containing a six-membered chelate exhibited poor ethenolysis reactivity and was capable of catalyzing the crossover of two internal olefins in the absence of ethylene, and thus reacts by a different pathway compared to 5. DFT calculations revealed the origins of the different reactivities of catalysts 3 and 5 in the crossover of internal olefins. The low crossover reactivity of two internal olefins with catalyst 5 is attributed to the steric repulsions of the nitrate anionic ligand with the chelating adamantyl group and the olefin substituent in the trisubstituted metathesis transition state. In ethenolysis reactions with catalyst 5, similar steric control also prevents the ruthenium alkylidene from reacting with internal olefins. In contrast, the most favorable ethenolysis pathway catalyzed by 5 involves the reaction of an internal olefin with a ruthenium methylidene to avoid trisubstituted metathesis transition states. Catalyst 3 has a smaller mesityl chelating group, and thus the steric repulsions with the anionic ligand are diminished, making it capable to productively form and cleave trisubstituted metallacycles.

The elucidation of a functional-group-tolerant ruthenium metathesis catalyst capable of performing *Z*-selective ethenolysis at ethylene pressures as low as 1 atm should enable the widespread use of this technology in academic and industrial settings. In addition to providing important insight into the ethenolysis behavior of catalyst 5, a better understanding of its CM reactivity has been gained and should provide important information for researchers planning on using this catalyst for a variety of applications. It is envisioned that the further development of new *Z*-selective catalysts should make this *Z*-selective ethenolysis methodology even more selective and efficient.

ASSOCIATED CONTENT

Supporting Information

Experimental details, NMR spectra, optimized Cartesian coordinates and energies, details of computational methods, and complete reference of Gaussian 09. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

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- (14) Basic olefin metathesis reactions are near thermoneutral and at equilibrium produce a statistical mix of products. Fortunately, the pre-equilibrium mixture of products can be controlled by the catalysts and by removal of one of the products. In RCM, the ethylene product can be removed and the structure of the catalyst can be used to control the initial product ratio and the rate of approach to equilibrium. In the present paper, the standard metathesis reaction is driven backward by the addition of ethylene and the structure of the catalyst controls the rate of reaction of the components of the reaction mixture.
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- (17) It should be noted that the volatility of the generated 1-hexene prevented it from being recovered and it was thus removed *in vacuo*.
- (18) The reactions depicted in Table 2 were performed to showcase the functional group tolerance of this method and the final %E. Isolated yields of highly %E products were obtained for two of the substrates (see Supporting Information), **12** (96% yield, >95%E) and the relatively volatile **11** (90% yield, >95%E).
- (19) The catalyst loading for E-5-decene was five times higher than for Z-5-decene.
- (20) The reactivity differences between **5** and **1** can be partially explained by the fact that **5** is not soluble in the 5-decene substrate, meaning that neat reactions could not be performed as with catalyst **5**. The necessary addition of solvent to these reactions seemingly reduced the activity of catalyst **5**.
- (21) It is envisioned that this Z-selective ethenolysis method can be used to purify products from reactions other than metathesis that produce E-olefins but that are not perfectly selective for their formation.
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- (27) Isomerization of **23** to **24** occurs via monodentate nitrate complexes. See Supporting Information for details.
- (28) Theoretical calculations predicted higher *Z*-selectivity than what the observed relative rates of *E* and *Z* olefins would suggest.
- (29) It should be noted that catalyst **4** was also not able to catalyze the CM of two internal olefins and, as such, we believe that trisubstituted metallacycle intermediates are also unfavorable.
- (30) For catalyst **4**, it is thought that formation of a methyldiene is possible and that the reaction proceeds via the same mechanism as with **5**, but reaction with a large excess of ethylene leads to decomposition.
- (31) Raising the temperature from 35 to 80 °C did not affect reactivity as formation of **20** was still not observed.
- (32) This suggests that the self-metathesis processes outlined in Scheme 1 are prevented, helping explain the high selectivity observed in the ethenolysis of methyl oleate by catalyst **5**.
- (33) To simplify calculations, an analog of catalyst **3** where the pivalate ligand is replaced by an acetate ligand was used for modeling.
- (34) The competing pathway to form *cis*-2-butene requires a barrier of 8.7 kcal/mol, only 0.2 kcal/mol higher than the *trans* pathway, indicating poor *Z/E*-selectivity from kinetic control.
- (35) Computational studies of olefin homodimerization with catalysts **2**, **4**, and **5** have already been reported (see ref 11). Here we repeated some of the computations to make direct comparisons between different catalysts and between productive and nonproductive metathesis.