The first ruthenium-silsesquioxyl complexes – synthesis, structure and mechanistic implications in silylative coupling†

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The first ruthenium-silsesquioxyl complexes have been synthesised and characterized via spectroscopic and X-ray methods. Mechanistic studies were performed and the complexes obtained were proved to be intermediates in the catalytic cycle of silylative coupling of olefins with vinylsilsesquioxane. Moreover, a mechanism for silylative coupling of styrene with vinylsilsesquioxane was proposed.

Introduction

Polyhedral oligosilsesquioxanes (POSS) of the general formula $(RSiO_{3.2})_n$ and particularly those $(n = 8)$ containing the inorganic cubic core of Si–O–Si bonds form a class of versatile building blocks for the production of inorganic–organic materials thanks to the three-dimensional highly symmetrical nature of the POSS core.¹ The properties of the core permit a wide range of technological applications of silsesquioxanes that can be used as nanofillers for the preparation of nanostructured composites, catalysts,²–⁴ and dendrimers,⁵ as precursors for optoelectronic materials,⁶–⁹ as dry resists in microelectronics¹⁰ and as precursors for SiO deposition.¹¹

Mono- and octa-functionalized silsesquioxanes and spherolitiles have been synthesized by TM catalyzed reactions such as hydrosilylation by Si–H¹² substituted POSS and silylative coupling¹³–¹⁶ by vinyl-substituted POSS with olefins, which occurs via intermediates containing a TM–silicon bond. The mechanism of silylative coupling of olefins with vinyl-substituted silicon compounds was the subject of our earlier study [see ref. 17; for reviews see ref. 18, 19]. So far only two reports have been published on the isolation and crystal structure of the complexes containing a TM–silicon bond (exactly cobalt–silicon bond).²⁰,²¹

This paper reports the synthesis of the first silylative coupled ruthenium complexes and describes the crystal structure of one of them. Moreover, the catalytic activity of this complex in the silylative coupling of vinylpentamethylsilsesquioxane with styrenes is analysed.

Results and discussion

When a toluene solution of the ruthenium hydride complex $[\text{RuHCl(CO)}(\text{PPh}_3)_3]$ ¹ was heated in the presence of 1 equiv. of vinylheptaisobutylsilsesquioxane at 110 °C, the color of the solution gradually turned yellow-brown within 48 h. The $^1$H NMR spectrum of the post-reaction mixture revealed disappearance of signals at $\delta = -6.60$ (dt) ppm characteristic of the Ru–H bond, and formation of a new singlet at $\delta = 5.25$ ppm which can be assigned to ethylene. Moreover, the appearance of a new singlet at $\delta = 37.49$ ppm was observed in the $^{31}$P NMR spectrum. On the basis of these observations, a general scheme for the synthesis of ruthenium-silsesquioxyl complexes was proposed (Scheme 1).

Further experiments led to the development of efficient synthetic procedures for the synthesis of silsesquioxyl complexes ³ and ⁴ (see ESI†). The compounds were isolated in high yields (89% and 86% respectively) as pale yellow powders. The obtained products were fully characterized by $^1$H, $^{13}$C, $^{29}$Si, and $^{31}$P NMR spectroscopy and high resolution mass spectrometry (see ESI†). Moreover the structure of complex ³ was confirmed via X-ray analysis (Fig. 1). Single crystals were...
obtained by slow evaporation of a pentane solution. Ru is 5-coordinated in a square-pyramidal fashion. Four atoms, P1, P2, Cl1 and Cl, are almost coplanar (maximum deviation from the least-squares plane is 0.053(3) Å) while the fifth one, Si1, is 2.485 Å out of this plane. The Ru atom is also slightly displaced, by 0.0219 Å, from the basal plane towards the apical Si1. Because of static disorder, in approximately one of six molecules the CO and Cl ligands are interchanged (cf. Experimental part) without a significant influence on the complex geometry. The Si8O12 moiety is quite regular, with deviations from the ideal symmetry caused mainly by the presence of Ru-coordination. The Si–O distances involving the coordinated silicon atom are systematically longer than the other Si–O bonds; the mean values are 1.636(4) Å for Si1 and 1.620(5) Å for all other ones. In fact, Si1–O distances are the three longest Si–O distances in the molecule. Also, the majority (8 of 12) of Si–O–Si angles are within quite a narrow range, with the mean value of 145.4(16)°, while the other four (O4, O8, O9, O10) are significantly larger, 154°–161°.

**Stoichiometric reactions of the ruthenium-silsesquioxyl complex with styrene**

As we previously demonstrated for the silylative coupling mechanism of vinylsilicon compounds and olefins, complexes 3 and 4 can be treated as intermediate ones in the catalytic cycle. In order to better understand the mechanism of silylative coupling of styrenes with vinylsilsesquioxanes an equimolar reaction between ruthenium-silsesquioxyl complex 3 and styrene was performed (Scheme 2).

The tests were monitored by 1H NMR spectroscopy. Addition of 1 equiv. of styrene and 5 equiv. of CuCl to the solution of complex 3 followed by heating the reacting mixture at 110 °C resulted in formation of the cross-coupling product 6.

Although spectroscopic examination does not confirm the formation of a complex 9 containing a Ru–H bond, we observed signals at δ = 6.47 ppm (d, 1H, J = 19.2 Hz, =CHSi) and δ = 7.56 ppm (d, 1H, J = 19.2 Hz, =CHPh) characteristic of product 6. The intermediate complex containing a Ru–H bond was also not observed in the reaction of Ru–Si complexes with olefins, which was previously reported by our group. Moreover, GC–MS analysis confirmed the formation of the desired coupling product 6. Formation of compound 6 is evidence for insertion of styrene into the Ru–Si(POSS) bond in complex 3 followed by β–H elimination and evolution of E-phenyl(silyl)-ethene.

In the above stoichiometric reaction, the desired product was formed only when the reaction was conducted in the presence of CuCl. Moreover, complex 3 exhibited almost no reactivity when the reaction was performed without Cu(i) salt. To better understand the role of CuCl in the reaction mixture we performed a series of stoichiometric reactions monitored by 1H and 31P NMR spectroscopy. When a toluene solution of the ruthenium-silsesquioxyl complex [Ru(POSS)Cl(CO)(PPh3)2] was heated in the presence of 5 equiv. of CuCl at 110 °C for 24 h we observed disappearance of the signals at δ = 37.49 ppm characteristic of complex 3 and formation of a new singlet at δ = 25.95 ppm in the 31P NMR spectrum (Fig. 2).

The formation of this new complex 8 was accompanied by precipitation of an insoluble grey-brown [CuPPh3] complex, whose appearance proves the dissociation of phosphine in the

**Scheme 2** Stoichiometric reactions of styrene with ruthenium-silsesquioxyl complexes.
proposed reaction system. The $^1$H NMR spectrum of the post-reaction mixture revealed changes in the aliphatic region. We observed disappearance of two multiplets at 1.85–2.06 and 2.06–2.31 and formation of a new multiplet at 2.15–2.38 ppm. All attempts to isolate this new complex $\mathbf{8}$ from the post-reaction mixture failed. However, the obtained results indicate that as a consequence of addition of CuCl to the reaction system, phosphine undergoes dissociation, as proved by [CuClPPh$_3$] complex precipitation, and a new four-coordinated intermediate complex $\mathbf{8}$ is formed. The addition of CuCl is needed because of a large steric hindrance in the intermediate complex having the Ru-Si(POSS) bond. The presence of the steric hindrance in the intermediate complex is supported by the fact that in the analogous silyl complex [RuSiMe$_3$Cl(CO)(PPh$_3$)$_2$] the insertion of intermediate complex is supported by the fact that in the analogous silyl complex [RuSiMe$_3$Cl(CO)(PPh$_3$)$_2$] the insertion of styrene is effective without the addition of CuCl.$^{17}$

Isotopic labelling experiments

In order to confirm that the coupling process in the presence of $\mathbf{3}$ proceeds via the insertion–elimination mechanism which is characteristic of a silylative coupling reaction,$^{17-19}$ we performed a series of labeling studies. In these experiments monitored by $^1$H NMR spectroscopy, a solution of equimolar amounts of styrene-d$_8$ and complex $\mathbf{3}$ was heated. The insertion–elimination mechanism should afford the formation of silylstyrene-d$_8$ and ethylene-d$_1$ (Scheme 3), at least in the initial stage of the reaction. $^1$H NMR spectroscopic analysis of the products formed in the equimolar reaction carried out in the presence of $\mathbf{3}$ revealed the formation of E-1-phenyl-2-(silyl)ethene-d$_1$ and ethylene-d$_1$.

The results obtained in the experiments with deuterium labelled styrene clearly demonstrate that functionalisation of vinylsilsesquioxanes with styrenes proceeds according to the silylative coupling mechanism involving the activation of =C–H and Si–C= bonds.

Catalytic examination and mechanistic implications

The ruthenium-silsesquioxyl complex $\mathbf{3}$ was examined in the silylative coupling of selected styrenes with vinylheptaisobutylsilsesquioxane and its catalytic activity was compared to that of its parent complex [RuHCl(CO)(PPh$_3$)$_3$] $\mathbf{1}$ and the previously studied more reactive complex [RuHCl(CO)(PCy$_3$)$_2$] $\mathbf{7}$ (Scheme 4).$^{15}$

![Scheme 3](image1.png)

**Scheme 3** Labeling experiments in the presence of complex $\mathbf{3}$.

![Scheme 4](image2.png)

**Scheme 4** Silylative coupling of vinylheptaisobutylsilsesquioxane with styrenes.

Table 1 Silylative coupling of vinylheptaisobutylsilsesquioxane with styrenes in the presence of $\mathbf{1}$, $\mathbf{3}$ and $\mathbf{7}$ with a co-catalyst (CuCl).

| $X$  | Catalyst | Yield of silylstyrene [%] |
|-----|---------|--------------------------|
| H   | 3       | 33                       |
|     | 1       | 6                        |
|     | 7       | 100*                     |
| OMe | 3       | 35                       |
|     | 1       | 7                        |
|     | 7       | 100*                     |
| Cl  | 3       | 32                       |
|     | 1       | 6                        |
|     | 7       | 100*                     |
| Br  | 3       | 34                       |
|     | 1       | 5                        |
|     | 7       | 100*                     |

**Reaction conditions**: CH$_2$Cl$_2$, reflux, argon, 5 h, [Ru]: [H$_2$C–CH(SiR$_3$)$_3$]:[styrene] = 10:2:1; [Ru]: CuCl = 1:5. *Ref. 15.

Selected data for $\mathbf{1}$ and $\mathbf{3}$ are presented in Table 1, entry 1. As indicated in this table, the silsesquioxyl catalyst $\mathbf{3}$ exhibits higher catalytic activity than catalyst $\mathbf{1}$. For example after five hours in the reaction of vinylsilsesquioxane with styrene catalysed by $\mathbf{3}$, the conversion of the silsesquioxane reached 33%, while the reaction catalysed by $\mathbf{1}$ revealed almost no conversion (6%). The activity of catalysts $\mathbf{1}$ and $\mathbf{3}$ was examined in a wide temperature range. Our experiments demonstrated that increasing the temperature of the reacting mixture from 40 °C to 110 °C did not give a higher yield of the coupling product. We compared the catalytic activity of complexes $\mathbf{1}$ and $\mathbf{3}$ with that of complex $\mathbf{7}$ bearing PCy$_3$ ligands. In all cases, complex $\mathbf{7}$ was the most active in silylative coupling of vinylheptaisobutylsilsesquioxane with styrenes; however, because of its high reactivity we were unable to isolate ruthenium-silsesquioxyl complex bearing PCy$_3$ ligands.

**Mechanistic implications**

On the basis of stoichiometric experiments and labelling studies we proposed a mechanism for the coupling of styrenes with vinylsilsesquioxanes in the presence of [RuHCl(CO)-(PPh$_3$)$_3$] $\mathbf{1}$ and CuCl as co-catalysts (Fig. 3).

Our studies have shown that the coupling reaction proceeds according to the well-recognized insertion–elimination mechanism$^{17}$ in which, in the first step, vinylsilsesquioxane reacts with hydride complex $\mathbf{1}$ to give β-silylethyl complex. This compound decomposes via β-silsesquioxyl group migration to ruthenium and evolution of ethylene to give Ru-Si[POSS] complex $\mathbf{3}$. Addition of CuCl to complex $\mathbf{3}$ causes dissociation of phosphine to produce a less sterically hindered four-coordinate complex $\mathbf{8}$ which reacts with styrene. The next step of the catalytic cycle involves the migratory insertion of styrene into a Ru–Si bond, followed by β-H elimination to give E-phenyl(silyl)ethene $\mathbf{6}$.

**Experimental**

**General methods and chemicals**

Unless mentioned otherwise, all operations were performed by using standard Schlenk techniques. $^1$H- and $^{13}$C-NMR spectra
were recorded on a Varian 400 operating at 402.6 and 101.2 MHz, respectively. $^1$P NMR spectra were recorded on a Mercury 300 operating at 121.5 MHz. $^{29}$Si NMR spectra were recorded on a Varian 400 operating at 402.6 and 101.2 MHz, respectively. $^{31}$P NMR spectra were recorded on a Varian Avance 600 operating at 119.203 MHz. All FT-IR spectra were recorded with IFS 113v FT-IR and VERTEX 70 spectrophotometers (Bruker, Karlsruhe). GC analyses were carried out on a Varian CP-3800 (column: Rtx-5, 30 m, I.D. 0.53 mm) equipped with TCD. Mass spectrometry analyses were performed using a Synapt G2-S mass spectrometer (Waters) equipped with an electrospray ion source and a quadrupole-time-of-flight mass analyzer. Acetonitrile was used as a solvent. The measurements were performed in positive ion mode with the desolvation gas flow 200 L h$^{-1}$ and capillary voltage set to 5000 V with the flow rate of 20 µl min$^{-1}$.

The chemicals were obtained from the following sources: vinyltrichlorosilane from ABCR, dichloromethane, acetone, $n$-pentane, ethanol, dichloromethane-d$_2$, benzene-d$_6$, toluene-d$_8$, styrene-d$_8$, decane, dodecane, styrene, 4-chlorostyrene, 4-bromo styrene, 4-methoxystyrene, 2-methoxethanol, triphenylphosphine, formaldehyde, copper(I) chloride, anthracene, calcium hydride and anhydrous magnesium sulphate from Aldrich, triethylamine and silica gel 60 from Fluka, ruthenium(III) chloride hydrate from Lancaster, trisilanloisoxybutyl POSS from Hybrid Plastics, and tolune and $n$-hexane from Chempur. [RuHCl(CO)(PPh$_3$)$_3$] was prepared according to the literature procedure. Monovinylheptaisobutylsilsesquioxane was prepared according to the literature procedure. Monovinylheptacyclopentylsilsesquioxane was prepared using the same procedure. All solvents were dried prior to use over CaH$_2$ and stored under argon. CH$_2$Cl$_2$ was additionally passed through a column with alumina and after that it was degassed by repeated freeze–pump–thaw cycles.

**Synthesis and characterization of ruthenium-silyl complexes**

**Synthesis and characterization of complex 3.** A 50 mL high-pressure Schlenk vessel connected to gas and vacuum lines was charged under argon with 0.2 g ($2.1 \times 10^{-4}$ mol) of [RuHCl(CO)(PPh$_3$)$_3$] and 20 mL of dry toluene. The mixture was warmed in an oil bath to 110 °C and stirred for 15 minutes. Subsequently, monovinylheptaisobutylsilsesquioxane (0.177 g, $2.1 \times 10^{-4}$ mol) was added to the reacting mixture and the suspension was refluxed for 48 h to give a yellow-brown solution. After this time the solvent was evaporated under vacuum, and $n$-hexane (5 mL) was added to the remaining contents to form a yellow precipitate. Then the resulting solution was decanted from the precipitate and the solid obtained was dried under vacuum (isolated yield = 89%).

Analytical data: $^1$H NMR (C$_6$D$_6$, δ, ppm): 0.35–0.53 (m, 14H, CH$_2$), 0.81–1.27 (m, 42H, CH$_3$), 1.85–2.06 (m, 3H, CH), 2.06–2.31 (m, 4H, CH), 6.99–7.30 (m, 12H, Ph), 7.72–7.93 (m, 18H, Ph); $^{13}$C NMR (C$_6$D$_6$, δ, ppm): 23.05, 23.25, 23.44, 23.62, 23.81, 24.72, 24.75, 24.78, 26.15, 26.20, 26.27, 26.38, 26.43, 30.45, 128.78 (t, $J = 5.0$ Hz), 130.49, 131.83 (d, $J = 2.5$ Hz), 132.61, 132.70, 132.85, 133.07, 133.29, 135.59 (t, $J = 5.8$ Hz), 200.70; $^{29}$Si NMR: δ = −66.05 (Ru-Si), −67.32, −69.60 (core); $^{31}$P NMR: (C$_6$D$_6$, δ, ppm): 37.49; IR (υ, cm$^{-1}$): 1931 (CO); MS (FD): m/z (%): 1231.24 (31), 1262.20 (61), 1265.20 (99), 1270.20 (100), 1268.20 (67), 1269.20 (50), 1275.23 (30), 1287.25 (42), 1288.24 (63), 1289.25 (97), 1290.24 (80), 1292.24 (55), 1299.09 (35), 1300.09 (30), 1465.13 (33), 1466.13 (40), 1467.13 (45), 1469.13 (42), 1470.13 (30), 1525.29 (38), 1526.29 (60), 1527.29 (95), 1529.29 (100), 1530.29 (77), 1531.29 (60), 1532.29 (33), 1543.29 (31), 1551.32 (39), 1552.32 (47), 1553.32 (62), 1554.32 (51), 1555.32 (47), 1556.32 (34), 1565.34 (38), 1566.33 (61), 1567.32 (92), 1568.33 (81), 1570.34 (57), 1571.33 (37), 1715.03 (33), 1717.03 (38), 1719.03 (34), 1803.38 (34), 1804.38 (54), 1805.37 (91), 1806.38 (93), 1807.38 (100), 1808.37 (82), 1809.38 (62), 1810.37 (37); HRMS (FD) for C$_6$H$_9$_5ClO$_2$P$_3$RuSi$_8$Na: calc. 1527.2875; found: 1527.2871.

**Synthesis and characterization of complex 4.** Complex 4 was prepared using the same procedure and the following reagents: [RuHCl(CO)(PPh$_3$)$_3$] (0.2 g, $2.1 \times 10^{-4}$ mol) and monovinylheptacyclopentylsilsesquioxane (0.1948 g, $2.1 \times 10^{-4}$ mol). Finally, a yellow precipitate was obtained (isolated yield = 86%).
Analytical data: $^1$H NMR (CD$_2$Cl$_2$, δ, ppm): 0.64–2.2 (m, 72H, C$_2$H$_{12}$–), 6.68–7.95 (m, 30H, Ph); $^{13}$C NMR (CD$_2$Cl$_2$, δ, ppm): 23.10, 23.19, 27.21, 27.56, 27.80, 27.83, 27.86, 27.89, 28.07, 28.31, 128.7 (t, $J$ = 4.5 Hz), 130.07, 131.59 (d, $J$ = 2.5 Hz), 132.34, 132.44, 135.44 (t, $J$ = 5.8 Hz), 199.82; $^{31}$P NMR: (CD$_2$Cl$_2$, δ, ppm): -48.47 (Ru–Si), -66.14, -68.75 (core); $^{31}$P NMR: (CD$_2$Cl$_2$, δ, ppm): 38.47; IR (ν, cm$^{-1}$): 1932 (CO); MS (FD): m/z (%): 1550.32 (24), 1551.33 (65), 1553.33 (100), 1554.33 (85), 1555.33 (82), 1556.33 (60), 1557.33 (37); HRMS (FD) for C$_2$H$_2$O$_3$P$_2$RuSi$_3$: calc. 1553.3289; found: 1553.3273.

Procedure for equimolar reactions of ruthenium-silyl complex 3 with styrene. The stoichiometric reactions were performed in J-Yong valve NMR tubes and controlled by $^1$H NMR spectroscopy. In a typical procedure ruthenium complex 3 with CuCl. (internal standard) were dissolved in 0.65 mL of toluene-d$_8$. Then the $^1$H NMR spectrum was recorded and 0.0032 g of 4-chlorostyrene was added under argon by a microliter syringe. Then the reaction mixture was warmed in an oil bath to 110 °C and after this time the $^1$H NMR spectrum of the reaction mixture was taken. The corresponding reactions with styrene-d$_8$ were performed using the same procedure.

Procedure for stoichiometric reactions of ruthenium-silyl complex 3 with styrene. The stoichiometric reactions were performed in J-Yong valve NMR tubes and controlled by $^1$H NMR spectroscopy. In a typical procedure, ruthenium complex 3 with CuCl. (internal standard) were dissolved in 0.65 mL of toluene-d$_8$. Then the $^1$H NMR spectrum was recorded and 1.2 mmol of a mixture of equal amounts of styrene and 4-chlorostyrene were performed using the same procedure.

General procedure for the catalytic examination. The oven dried 5 mL glass reactor equipped with a condenser and a magnetic stirring bar was charged under argon with 2 mL of CH$_2$Cl$_2$, monovinylsilsesquioxane (0.1 g, 1.19 × $10^{-4}$ mol), styrene (41 µL, 3.56 × $10^{-4}$ mol) and 20 µL of an internal standard (decane or dodecane). The reaction mixture was placed in an oil bath and preheated to 40 °C. Then ruthenium complex 1 or 3 (0.0011 g (complex 1) or 0.0017 g (complex 2), 1.19 × $10^{-4}$ mol) was added under argon. After 5 min of the reaction, copper(I) chloride (0.0006 g, 5.93 × $10^{-6}$ mol) was added. The reaction mixture was heated at 40 °C under a gentle flow of argon. Reaction yields were calculated on the basis of the $^1$H NMR spectra of the reaction mixture. The corresponding reactions with 4-methoxystyrene, 4-bromostyrene and 4-chlorostyrene were performed using the same procedure.

X-ray analysis (complex 3). X-ray diffraction data were collected at 100(1) K by the ω-scan technique, on an Agilent Technologies four-circle Xcalibur diffractometer equipped with an Eos detector with a MoK$_\alpha$ radiation source ($\lambda$ = 0.71073 Å). The temperature was controlled with an Oxford Instruments Cryosystem device. The data were corrected for Lorentz-polarization effects as well as for absorption (multiscan). Accurate unit-cell parameters were determined by a least-squares fit of 12 093 reflections of highest intensity, chosen from the whole experiment. The calculations were performed within the WinGXY program system. The structures were solved with SIR92 and refined with the full-matrix least-squares procedure on $F^2$ using SHELXL97. Scattering factors incorporated in SHELXL97 were used. The function $\Sigma w|F_o|^2 - |F_c|^2$ was minimized, with $w^{-1} = [\sigma^2(F_o)^2 + (0.0313P)^2 + 2.2712P]$, where $P = \max(F_o^2, 0) + 2E^2/3$. All non-hydrogen atoms were refined anisotropically, hydrogen atoms from methyl groups were placed geometrically, in idealized positions, and refined as riding group with their $U_{iso}$ set at 1.2 (1.5 for methyl groups) times $U_{eq}$ of the appropriate carrier atom. Disorder was detected during structure refinement: Cl and CO ligands are partially exchanged (site occupation factors 0.83/0.17), and in one iPr group the methyl groups are disordered over two positions (s.o.f.'s 0.70/0.30). Crystallographic data (excluding structure factors) for the structural analysis have been deposited with the Cambridge Crystallographic Data Centre, no. CCDC-919730.

Crystal data: C$_{65}$H$_{65}$ClO$_{12}$P$_2$RuSi$_8$, $M_r$ = 1505.57, triclinic, $P1$, $a$ = 13.4910(3) Å, $b$ = 14.8072(3) Å, $c$ = 19.0764(4) Å, $α$ = 89.338(2)°, $β$ = 81.060(2)°, $γ$ = 77.729(2)°, $V$ = 3677.57(13) Å$^3$, $F(000)$ = 1580, $d_c$ = 1.36 g cm$^{-3}$, $\mu$(MoK$_\alpha$) = 0.48 cm$^{-1}$, 33 625 reflections measured, 14 864 unique ($R_{int}$ = 0.024), 12 507 with $I > 2\sigma(I)$. Final $R$(obs) = 0.031, $R$(all) = 0.042, $wR$_2(obs) = 0.071, $wR$_2(all) = 0.075, $S$ = 1.04, max/min $Δρ$ in the final $ΔF$ map: 0.49/-0.62 e Å$^{-3}$.

Conclusions

New silesquioxyl ruthenium complexes (3 and 4) have been synthesized and their structures were confirmed by spectroscopic and X-ray methods. These complexes were proved to be intermediates in the silylative coupling of vinylsilsesquioxane with styrene. The reaction between vinylsilsesquioxanes and styrenes in the presence of ruthenium hydride complex 1 was confirmed to proceed via the insertion–elimination mechanism. Moreover, the obtained complex 3 exhibits higher catalytic activity than its parent hydride complex 1.

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