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STUDIES ON NEW CATALYZED REACTIONS USING HYDROSILANES AND CARBON MONOXIDE



YOSHIO SEKI

OSAKA UNIVERSITY OSAKA JAPAN JANUARY, 1978

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USING HYDROSILANES

AND CARBON MONOXIDE

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Preface

This thesis deals with the studies accomplished by the author under the guidance by Professor Noboru Sonoda at the Department of Petroleum Chemistry, Faculty of Engineering, Osaka University.

The author would like to express his sincere thanks to Professor Noboru Sonoda for his invaluable guidance and constant encouragement throughout this investigation since 1972.

The author is also deeply grateful to Associate Professor Shinji Murai for his helpful suggestion and discussions. Grateful acknowledgements are also due to the late Dr. Membo Ryang, Dr. Ilsong Rhee, Mr. Noritaka Miyoshi, and Dr. Kiyoshi Kondo for their valuable suggestions through stimulating discussions with the author.

The author expresses his gratitude to Mr. Iwao Yamamoto and Akihiko Hidaka for their collabolation in the course of experiment. The author's grateful thanks is also due to Shigeo Makino, Makoto Fukutani, and Toshikazu Kato for their assistance in the cource of experiment. The author wishes to thank all the members of Sonoda Laboratory.

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Suita, Osaka January, 1978

Yoshio Seki

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Introduction

Carbon monoxide was discovered in 1776 by Lassone and already found technical application in 19th century. In the first half of the 20th century, many important reactions using carbon monoxide were discovered in Germany. Especially, hydroformylation(oxo reaction) discovered by O. Roelen in 1938¹ and Reppe reactions reported by W. Reppe in 1953² have still been employed for technical production of useful materials. A fundamental catalyzed reaction using carbon monoxide, however, has not been discovered since the discoveries of hydroformylation(oxo reaction) and Reppe reactions except for the rhodium(III)-catalyzed reaction which gives acetic acid from methanol and carbon monoxide³.

Recently, with a high price of oil and a shortage of natural gas, the countries without abundant indigeous oil resources must make good use of the coal resources again. This is resulting in the revival of carbon monoxide chemistry. Therefore, interest in a new catalytic process using carbon monoxide has increased again.

It is well known that the chemistries of hydrosilanes and of molecular hydrogen are very similar with respect to the various mechanisms of catalytic activation. For example, in the case of oxidative addition to Ir(I) complex, both of a hydrosilane⁴ and hydrogen^{5a} react to give cis-adducts. Hydrosilylation⁶, the catalyzed addition of R₃SiH to an olefin,

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is formally analogous to catalytic hydrogenation⁵(Fig. 1). Moreover, mechanistically, these reactions seem to be very closely related.^{5,6,7a}

The well-known hydroformylation (oxo reaction)⁷ involves addition of hydrogen and carbon monoxide to olefins. The similarity between hydrogen and a hydrosilane has made us curious about the catalyzed reaction of olefins with a hydrosilane and carbon monoxide. However, no such reaction has been reported so far except for very complex results reported by Chalk and Harrod.⁸ If the catalyzed reaction of

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Н-Н	H-Si
hydrogenation	hydrosilylation
$C=C + H_2 \longrightarrow C - C$	C=C + HSi→ Ç—Ç H Si
hydroformylation	Not known
$C=C + CO + H_2 \longrightarrow \begin{array}{c} C &C \\ H & C=O \\ H \end{array}$	C=C + CO + HSi

an olefin with a hydrosilane and carbon monoxide is found, it will be expected that the discovery opens an entirely new field spanding over organosilicon chemistry, homogeneous catalysis, and synthetic organic chemistry. From these points of view, the author studied catalyzed reactions using hydrosilanes and carbon monoxide.

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This thesis consists of three chapters.

Chapter 1 deals with a catalyzed reaction of olefins with hydrosilanes and carbon monoxide. Futhermore, the mechanism is discussed on the bases of the reaction of cyclohexene with diethylmethyldeuteriosilane and carbon monoxide. In chapter 2, a catalyzed reaction of cyclic ethers with a hydrosilane and carbon monoxide is described. In chapter 3, a catalyzed reaction of aldehydes with a hydrosilane and carbon monoxide is described.

This thesis is composed of the following reports.

- (1) The Reaction of Olefins with Silane and Carbon Monoxide Catalyzed by Co₂(CO)₈
 Y. Seki, A. Hidaka, S. Murai, and N. Sonoda,
 Angew. Chem., <u>89</u>, 196(1977); Angew. Chem. Int. Ed. Engl.,
 <u>16</u>, 174(1977).
- (2) Transition Metal Reactions of Silanes II. The Reaction of Cycloalkenes with Various Hydrosilanes and Carbon Monoxide Catalyzed by Co₂(CO)₈
 Y. Seki, A. Hidaka, S. Makino, S. Murai, and N. Sonoda, J. Organomet. Chem., 140, 361(1977).
- (3) The Reaction of Cyclic Ethers with a Hydrosilane and Carbon Monoxide Catalyzed By Co₂(CO)₈
 Y. Seki, S. Murai, I. Yamamoto, and N. Sonoda, Angew. Chem., <u>89</u>, 818(1977).

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- (4) Catalyst-Dependent Product Distribution in the Reaction of 1-Hexene with a Hydrosilane and Carbon Monoxide
 Y. Seki, S. Murai, A. Hidaka, and N. Sonoda,
 Angew. Chem., in press.
- (5) The Reaction of Aldehydes with a Silane and Carbon Monoxide Catalyzed by Co₂(CO)₈-Ph₃P
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Chapter 1 A Catalyzed Reaction of Olefins with Hydrosilanes and Carbon Monoxide

1-1 Introduction

There have been several studies dealing with the catalyzed reaction involving a hydrosilane and carbon monoxide or the reaction between the complexes containing silicon-metal bonds and carbon monoxide. Chalk and Harrod reported that no direct carbonylation of triethylsilane occurred at temperatures up to 300°C and carbon monoxide pressures up to 1700 psi in the presence of $Co_2(CO)_8$.¹ It has been reported that attempts to induce carbonyl insertion into the Si-Co², Si-Mn³, and Si-Rh⁴ bond by direct reaction with carbon monoxide were unsuccessful. Hagen and co-workers carried out the reaction of MeSiCl₂Co(CO)₄ with 4000 atm of carbon monoxide at 90°C, but no insertion of carbon monoxide into the silicon-cobalt bond was observed.⁵ Insertion of carbon monoxide into the carbon-cobalt bonds is very easy⁶. This difference suggests that silicon-cobalt bonds are similar to a hydrogen-cobalt bond with respect to the insertion reaction of carbon It has been reported that the use of carbon monoxide, monoxide. as well as of phosphines, inhibited dicobalt octacarbonylcatalyzed addition of dimethylchlorosilane to acrylonitrile'.

Chalk and Harrod have reported an interesting reaction in 1967¹. They studied the dicobalt octacarbonyl-catalyzed reaction of olefins with triethylsilane and carbon monoxide. The reaction of 1-pentene and cyclohexene was reported to be "very complex" and no detail were given. In the case of ethylene, however, a complex mixture of the reaction products

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with boiling points ranging from 90°C to 300°C was fractionated into thirty three fractions and one of them was found to contain unsaturated alkoxy silane(s) of the formula $C_{3}H_{5}OSi(C_{2}H_{5})_{3}$ which gave the 2,4-dinitrophenylhydrazone of propionaldehyde on treatment with the hydrazine.

In contrast to this previous work, the author has found that carbon monoxide reacts very smoothly with cycloalkenes and hydrosilanes in the presence of $\text{Co}_2(\text{CO})_8$. The catalytic reaction gave remarkably clean results. From cycloalkenes, only one product, the trialkylsiloxymethylenecycloalkane, has been obtained in most cases studied. In this chapter, the results of the catalyzed reaction of olefins with hydrosilanes and carbon monoxide under various reaction conditions are described.

1-2 Results and Discussion

1-2-1 Reaction of Cycloalkenes with Diethylmethylsilane and Carbon Monoxide

Contrary to the "very complex" results reported by Chalk and Harrod for the reaction of cyclohexene with triethylsilane and carbon monoxide in the presence of $\text{Co}_2(\text{CO})_8$, as mentioned above, we have found that this reaction gave triethylsiloxymethylenecyclohexane(Id) as a sole product in a good yield.

Unfortunately, it is not possible to explain the difference in the results of Chalk and Harrod and ours, since no experimental details were given in their paper. See ref. (1).

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For diethylmethylsilane, a similar result was found. If one assumes that the reaction proceeds in a formally similar way as hydroformylation, two types of products, (II) and (III), may be expected. However, neither of these compounds was found. The stoichiometry of the reaction can be presented as shown in eq. 1, and all these reactants are incorporated into the product(I). While it is obvious that the carbon and oxygen atoms of carbon monoxide are found in the oxymethylene group of the product, it will be discussed in this chapter which hydrogen in the product(I) has come from the hydrosilane used.

The results obtained for the reaction using cycloalkenes (3 equivalents) and diethylmethylsilane(l equivalent) are given in the Table I. All the catalytic reactions were carried



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Olefin	Catalysts	Product	Yield(%) (GLC)
	Co ₂ (CO) 8	OSiEt ₂ Me(IV)	48
	co ₂ (co) ₈	OSiEt ₂ Me(Ic)	71
,	Ru ₃ (CO) 12		1.4
	RhCl (PPh ₃) ₃		0.5
 \bigcirc	co ₂ (co) ₈	OSiEt ₂ Me(V)	74
\bigcirc	Co ₂ (CO) ₈	OSiEt ₂ Me(VI)	69
	Co ₂ (CO) ₈	OSiEt ₂ Me(VII)	22
	^{Co} 2 (CO) 8	OSiEt ₂ Me (VIII	45)

Table I Reaction of Various Olefins with Et, MeSiH and CO^a

a) Reaction conditions: cat. (0.2 mmol), olefin(30 mmol), Et₂MeSiH (10 mmol), CO(80 kg/cm²), 140°C, 20 h, in benzene(20 ml)

out using a 100 ml stainless steel autoclave. Generally, a solution of a cycloalkene(30 mmol), Et, MeSiH(10 mmol), and Co2(CO)8(0.2 mmol) in benzene(20 ml) was heated under a pressure of carbon monoxide at 140°C for 20 h to give diethylmethylsiloxymethylenecycloalkane(Ic and IV-VIII). The pressure of carbon monoxide was initialy 80 kg/cm² at 25°C, then reached 105-108 kg/cm² at 140°C, and it was finally 76-79 kg/cm² at 25°C after the reaction had been completed. Simple cyclic olefins(cyclohexene, cycloheptene, and cyclooctene), except for cyclopentene, gave the expected product in good yield, while the reaction of 1-methylcyclohexene, a trisubstituted olefin, and norbornene, a bicyclic olefin, gave the expected product only in moderate yield. In the case of 1-methylcyclohexene, 2-diethylmethylsiloxyethenyl-l-cyclohexane was not obtained. All the reactions were accompanied by the formation of the corresponding disiloxane in 5-15% yield.

When $RhCl(PPh_3)_3$ and $Ru_3(CO)_{12}$ were used as a catalyst for the reaction of cyclohexene, only traces of the enol silyl ether(Ic) were obtained. This result is interesting, since hydroformylation of cyclohexene has been known to give good yields of cyclohexanecarboxaldehyde in the presence of $Ru_3(CO)_{12}$.⁸

1-2-2 Reaction of Terminal Olefins with Diethylmethylsilane and Carbon Monoxide

The reaction of terminal olefins with diethylmethylsilane and carbon monoxide gave four isomers(regio- and stereoisomers)

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as shown in eq. 2. The results in the reaction of 1-hexene and methyl acrylate were summarized in Table II and III.

$$R$$
 + Et₂MeSiH + CO cat.

 $(R=n-C_4H_9, COOMe)$



Table II	Catalyzed	Reaction of	1-Hexene	a)		
Cataly	st	Total yield(%)	Product (Z)-(IXa	distributi a) (E)-(IXa)	on(%) (Z)-(Xa)	(E)-(Xa)
Co ₂ (CO) ₈	57	16	16	43	25
RhCl(P	^{Ph} 3 ⁾ 3	41	9	9	58	24
Ru ₃ (CO) ₁₂	40	4	7	44	45

a) Reaction conditions; cat.(0.4 mmol), olefin(30 mmol), Et₂MeSiH (10 mmol), CO(50 kg/cm²), 140°C, 20 h , benzene(20 ml).

			`				
Catalyst	MA	HSi	Total	Product	distri	bution	(
	(mmol)	(mmol)	yield(%)	(Z-IXD)	(E-IXD)	(Z-XD)	(E-XD)
Co ₂ (CO) ₈	30	10	49	0	12	59	29
	10	30	6	0	5	72	23
RhCl (PPh3)3	30	10	6	42	58	0	0
-	10	30	18	11	89	0	0
Ru ₃ (CO) 12	30	10	9	43	57	0	0
	10	30	22	36	64	0	0

Table III Catalyzed Reaction of Methyl Acrylate^{a)}

a) Reaction conditions; $Co_2(CO)_8(0.2 \text{ mmol})$, RhCl(PPh₃)₃ (0.1 mmol), Ru₃(CO)₁₂(0.05 mmol); methyl acrylate=MA $CO(80 \text{ kg/cm}^2)$, 140°C, 20 h, benzene(20 ml).

In the reaction of cyclohexene, $RhCl(PPh_3)_3$ and $Ru_3(CO)_{12}$ were not effective catalysts, but the reaction of terminal olefins was catalyzed by these metal complexes. In the reaction of methyl acrylate, straight chain isomers(X) were main products in the case of $RhCl(PPh_3)_3$ and $Ru_3(CO)_{12}$, while branched isomers(IX) were main products in the case of $Co_2(CO)_8$. These results show that the rate-limiting step differs between the case of $RhCl(PPh_3)_3$ and $Ru_3(CO)_{12}$ and that of $Co_2(CO)_8$. When a three fold excess of hydrosilane to methyl acrylate was used in the reaction catalyzed by $RhCl(PPh_3)_3$ or $Ru_3(CO)_{12}$, the yields increased to about 20%.

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1-2-3 Reaction of Cyclohexene with Various Hydrosilanes

Cyclohexene reacted with various hydrosilanes and carbon monoxide in the presence of $Co_2(CO)_8$ to give the corresponding siloxymethylenecyclohexanes(I) as shown in eq. 1. The results were summarized in Table IV.

		•	· · · ·		
Hudrogilano	Yield o	f	Recovered	(p ai) o ^b	
nyulostiane	the pro	oduct(I)	hydrosilane(%)	⁽¹ 3 ³¹⁾ 2 ⁰	
Me ₃ SiH	(Ia)	4	88	0	
EtMe ₂ SiH	(Ib)	57	36	5	
Et2 ^{MeSiH}	(Ic)	71	5	8	
Et ₃ SiH	(Id)	74	0	23	
PhMe ₂ SiH	(Ie)	51	29	18	
Ph_2^{MeSiH}	(If)	26	57	6	
Ph ₃ SiH		0	88	0	
Ph2SiH2		0	39	18	
(MeO) ₂ MeSiH		0	0	0	

Table IV Reaction of Cyclohexene with Various Hydrosilanes and Carbon Monoxide^a

a) Reaction conditions; Co₂(CO)₈(0.2 mmol), cyclohexene
 (30 mmol), hydrosilane(10 mmol), CO(80 kg/cm²), 140°C,
 20 h, in benzene(20 ml).

b) Disiloxanes corresponding to hydrosilanes used.

Trialkylsilanes, except for Me_3SiH , yielded the corresponding siloxymethylenecyclohexanes(Ib-Id) in good yield. In the case of hydrosilanes having a phenyl substituent, the yield of the product(Ie and If) decreased in the order $PhMe_2SiH$ Ph_2MeSiH Ph_3SiH . The following hydrosilanes did not give the corresponding siloxymethylenecyclohexane: Ph_3SiH , Ph_2SiH_2 , and (MeO) 2MeSiH. The reaction of (MeO) 2MeSiH gave very complex results without recovery of the hydrosilane. These results show that hydrosilanes with electron-donating substituents have good reactivity in the present reaction. The reason for low yield in the case of Me_3SiH is not clear. It may be intrinsic or due to the different experimental procedure(see Experimental Part).

The reaction with dimethylchlorosilane was also examined. Two products were obtained in about 10% yields, respectively. One product was cyclohexanecarboxaldehyde and the other has not been identified yet. White solid precipitated from the liquid which was obtained by distillation(b.p. 83-95°C/15 torr) of the reaction mixture. The white solid was identified with 2,4,6-tricyclohexylparaldehyde. The expected product, dimethylchlorosiloxymethylenecyclohexane, may be very unstable to moisture and hydrolyzed during work-up.

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1-2-4 Mechanism of the Reaction of Olefins with Hydrosilanes and Carbon Monoxide

Effects of the reaction conditions such as the pressure of carbon monoxide, the temperature, and the ratio of methyl acrylate to Et₂MeSiH on the total yield and the product distribution were examined in the reaction of methyl acrylate. The results are summarized in Table V

Table	V Rea	ction of	E Methy	yl Acryl	ate with	Et2 ^{MeS}	iH and	l co ^a
MA	SiH	co ^c	Temp.	Total	Product	distri	butior	b
(mmol)	(mmol)	(kg/cm^2)	(°C)	yield(%)(Z-IXb)	(E-IXb)	(Z-Xb)	(E-Xb)
30	10	80	140	49	0	12	59	29
30	10	50	140	50	0	16	58	26
30	10	30	140	49	0	12	61	27
30	10	30	70	29	8	55	30	7
10	10	80	140	44	0	9	68	23
10	30	80	140	6	0	5	72	23

a) All reactions were carried out in 20 ml of benzene for 20 h.

b) The yields and the product distribution were determined by GLC.

c) Initial pressure at room temperature.

When the other reaction conditions were fixed, the influence of the pressure of carbon monoxide on the total yields and the percentage of straight chain isomers may be neglected.

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The total yield increased with increase in the ratio of methyl acrylate to diethylmethylsilane. Branched isomers(IXb) were main products at 70°C. It has already been reported that branched isomers increased at lower temperature in hydroformylation(oxo reaction) of ethyl acrylate catalyzed by $\text{Co}_2(\text{CO})_8$.⁹ The results suggest that the reaction path in the reaction of methyl acrylate with diethylmethylsilane and carbon monoxide is similar to that in hydroformylation(oxo reaction) of ethyl acrylate at the initial stage.

Similarly, effects of the reaction conditions on the total yield and the product distribution were examined in the reaction of cyclohexene. When the reaction were carried out at higher temperature or using equimolar amounts of cyclohexene and a hydrosilane, another product(XI) was obtained in addition to Ic (eq. 3). The product(XI) may be formed by catalyzed double-bond isomerization of the product(Ic).



The yield and distribution of the products were found to be affected by the reaction conditions such as molar ratio of reactants, the temperature and the pressure of carbon monoxide. The results obtained for the reaction of cyclohexene with diethylmethylsilane and carbon monoxide are shown in the Table VI.

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Tak	ole	VI	Effects	of	the	Read	ction	Condi	tions	in	the	Reaction	L.
										÷.,		· •	
of	Cvo	lob	nexene wi	th :	Et_Me	∋SiH	and	co ^a	÷.,	· ·			

	Reac	tion co	onditions	Total	b •P	roduct	t ^b
Olefin	SiH	Temp.	CO pressu	re ^C yield	(%) d:	istril	oution
(mmol)	(mmol)	(°C)	(kg/cm^2)	(Ic +	XI)	Ic :	XI(%)
10	10	140	30	12		99 :	1
10	10	140	60	36		97 :	3
10	10	140	80	37		98 :	2
20	10	140	80	51		98 :	2
30	10	140	80	71	-	100 :	0
30	10	200	80	84		80 :	20

- a) All reactions were carried out in 20 ml of benzene for20 h.
- b) The yields and the product distribution were determined by GLC.
- c) Initial pressure at room temperature.

The total yield of Ic and XI increased with an increase in the pressure of carbon monoxide, but the difference was very small at 60-80 kg/cm². The total yield increased remarkably with increasing molar ratio of cyclohexene to diethylmethylsilane. The total yield was best at 200°C, but the yield of the initial product(Ic) decreased and that of isomerization product(XI) increased.

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The fact that the use of a higher molar ratio of cyclohexene to the hydrosilane has resulted in a higher yield of the product suggested the importance of hydrocobaltcarbonyl $HCo(CO)_4$, as the active catalyst in the present reaction. The reaction of trialkylsilane with $Co_2(CO)_8$ has been known to give $HCo(CO)_4$ and $R_3SiCo(CO)_4^{1,10}$ (eq. 4). The hydrocobaltcarbonyl may react with olefins to initiate a catalytic cycle(eq. 5). However, when a sufficient amount of the olefin is not present in the system, it reacts with the hydrosilane to give $R_3SiCo(CO)_4$ and hydrogen 1,10 (eq. 6). This reaction (eq. 6) would stop the catalytic cycle. The alkylcobalt





complex formed in eq. 5 may give an acylcobalt complex which then reacts with a hydrosilane to give the product with regeneration of hydrocobaltcarbonyl. The reaction path(eq. 5) is the same as the initial stage of hydroformylation.

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As mentioned above, an acylcobalt complex is formed at the initial stage. Next, the reaction path from the acylcobalt complex to the products was investigated. To' begin with, the deuterium position in the products was determined in the reaction of cyclohexene with diethylmethyldeuteriosilane and carbon monoxide. The reaction gave the corresponding siloxymethylenecyclohexane in 87% yield as shown in eq. 7. The amount of deuterium incorporation at the vinyl position of the product was 91%.



From these results, the following two reaction paths were proposed. The reaction path via an acylsilane or hydroformylation(oxo reaction) was completely excluded.

1)
$$\operatorname{RCH}_{2}\operatorname{CH}_{2}^{\circ}\operatorname{CCo}(\operatorname{CO})_{3} \xrightarrow{\operatorname{HSi}} \operatorname{RCH}_{2}^{\circ}\operatorname{CH}_{2}^{\circ}\operatorname{C}_{0}^{\circ}(\operatorname{CO})_{3} \xrightarrow{\operatorname{CO}} \xrightarrow{\operatorname{CO}} \operatorname{RCH}_{2}^{\circ}\operatorname{CH}_{2}^{\circ}\operatorname{C}_{0}^{\circ}(\operatorname{CO})_{3} \xrightarrow{\operatorname{CO}} \operatorname{RCH}_{2}^{\circ}\operatorname{CH}_{2}^{\circ}\operatorname{C}_{0}^{\circ}(\operatorname{CO})_{3} \xrightarrow{\operatorname{CO}} \operatorname{RCH}_{2}^{\circ}\operatorname{CH}_{2}^{\circ}\operatorname{C}_{1}^{\circ}\operatorname{CO} \operatorname{RCH}_{2}^{\circ}\operatorname{C}_{1}^{\circ}\operatorname{CO} \operatorname{RCH}_{2}^{\circ}\operatorname{C}_{1}^{\circ}\operatorname{CO} \operatorname{RCH}_{2}^{\circ}\operatorname{C}_{1}^{\circ}\operatorname{CO} \operatorname{RCH}_{2}^{\circ}\operatorname{C}_{1}^{\circ}\operatorname{RCH}_{2}^{\circ}\operatorname{C}_{1}^{\circ}\operatorname{CO} \operatorname{RCH}_{2}^{\circ}\operatorname{C}_{1}^{\circ}\operatorname{RCH}_{2}^{\circ}\operatorname{RCH}_{2}^{\circ}\operatorname{C}_{1}^{\circ}\operatorname{CO} \operatorname{RCH}_{2}^{\circ}\operatorname{RC$$

$$\xrightarrow{-HCo(CO)_3}$$
 RCH₂CH=CHOSi

(

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RCH, CH=CHOSi

In the route(1), initially, a hydrosilane adds oxidatively to the acylcobalt complex to give the intermediate(XII) and then an aldehyde reductively eliminates. The aldehyde reacts with silylcobaltcarbonyl to give the intermediate(XIII). The last reaction probably involves the β -elimination of hydrocobaltcarbonyl and the formation of the product.

In the route(2), the silyl group shifts to the oxygen of the acyl group in the intermediate(XII) produced to give the siloxycarbene complex(XIV) and then the carbene inserts to the cobalt-hydrogen bond to give intermediate(XIII). The subsequent path is the same as the route(1).

It is very difficult to determine which route is real. To begin with, the possibility of an aldehyde as an intermediate was investigated. Experimentally, a solution of n-heptanal (10 mmol), diethylmethylsilane(10 mmol), and $\text{Co}_2(\text{CO})_8(0.4 \text{ mmol})$) in benzene(20 ml) was heated at 140°C for 20 h under carbon monoxide(50 kg/cm²) in 100 ml of autoclave. The results were shown in Table VII and eq. 8.

The reaction of the aldehyde(XV) with the hydrosilane (Run 1 and 2) gave almost alkoxysilane(XVI), while the reaction of olefins with the hydrosilane and carbon monoxide (Run 3)

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Table VI Reaction of an Aldehyde and Reaction of an Olefin							
Run	Reactants	Yield of (X)+(XVI)	(X : XVI)				
	XV : HSi (1:1)	40	18:82				
2	XV : HSi : 1-Octene (0.1:1:1)	69	3:97				
3	HSi : 1-hexene(1:3)	39	100 : 0				

gave only the enol silyl ethers(X). These results suggest that the aldehyde(XV) is not an intermediate. But it is possible that hydrosilylation of the aldehyde(XV) occurres for one hour, since it takes one hour that the temperature rises from room temperature to 140°C. Therefore, the direct reaction of the aldehyde(XV) with the hydrosilane at about 140°C was tried. Experimentally, a solution of an aldehyde (2 mmol) in 2 ml of xylene was gradually added to the mixture of diethylmethylsilane(4 mmol) and $\text{Co}_2(\text{CO})_8(0.04 \text{ mmol})$ in 4 ml of xylene at about 140°C. The enol silyl ethers(X) and the alkoxysilane(XVI) were obtained in 79% and 21% yields, respectively. This result suggests that the enol silyl ethers(X) can be produced selectively under proper reaction conditions.¹¹ Therefore, it is reasonable that the reaction of an olefin with a hydrosilane and carbon monoxide proceeds via an aldehyde as an intermediate. The plausible mechanism was shown below.

$$HSi + Co_2(CO)_8 \longrightarrow HCo(CO)_4 + SiCo(CO)_4 (a)$$

$$HCo(CO)_4 \xrightarrow{-CO} [HCo(CO)_3]$$
(b)

$$\operatorname{RCH=CH}_{2} + [\operatorname{HCo}(\operatorname{CO})_{3}] \xrightarrow{\operatorname{RCH}_{2}\operatorname{CH}_{2}\operatorname{Co}(\operatorname{CO})_{3} \xrightarrow{\operatorname{CO}} \operatorname{RCH}_{2}\operatorname{CH}_{2}\operatorname{Co}(\operatorname{CO})_{4} (c)$$

$$\operatorname{RCH}_{2}\operatorname{CH}_{2}\operatorname{Co}(\operatorname{CO})_{4} \rightleftharpoons \operatorname{RCH}_{2}\operatorname{CH}_{2}\operatorname{CCo}(\operatorname{CO})_{3} \rightleftharpoons \operatorname{RCH}_{2}\operatorname{CH}_{2}\operatorname{CCo}(\operatorname{CO})_{4} \qquad (d)$$

$$\operatorname{RCH}_{2}\operatorname{CH}_{2}^{\circ}\operatorname{CCO}(\operatorname{CO})_{3} + \operatorname{HSi} \xrightarrow{} \operatorname{RCH}_{2}\operatorname{CH}_{2}^{\circ}\operatorname{CCO}(\operatorname{HSi})(\operatorname{CO})_{3} \quad (e)$$

$$\operatorname{RCH}_{2}\operatorname{CH}_{2}^{\circ}\operatorname{CCO}(\operatorname{HSi})(\operatorname{CO})_{3} \longrightarrow \operatorname{RCH}_{2}\operatorname{CH}_{2}\operatorname{CHO} + \operatorname{SiCO}(\operatorname{CO})_{3} \quad (f)$$

$$sico(co)_3 \xrightarrow{co} sico(co)_4$$
 (g)

(i)

.

$$\operatorname{RCH}_{2}\operatorname{CH}_{2}\operatorname{CHO} + \operatorname{SiCo}(\operatorname{CO})_{4} \longrightarrow \operatorname{RCH}_{2}\operatorname{CH}_{2}\operatorname{CCO}(\operatorname{CO})_{4} \qquad (h)$$

$$\operatorname{RCH}_{2}\operatorname{CH}_{2}_{H}^{\operatorname{CCO}(\operatorname{CO})}_{H} 4 \xrightarrow{-\operatorname{CO}}_{\operatorname{RCH}_{2}} \operatorname{RCH}_{2}\operatorname{CH}_{2}_{H}^{\operatorname{CCO}(\operatorname{CO})}_{H}_{H}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{RCH}_{2}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{RCH}_{2}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{RCH}_{2}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{RCH}_{2}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{OSi}}_{\operatorname{OSi}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{OSi}}_{\operatorname{OSi}}_{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{\operatorname{H}}^{\operatorname{OSi}}_{$$



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Ojima and co-workers have already reported that hydrosilylation of acetophenone catalyzed by $RhCl(PPh_3)_3$ proceeded under more drastic conditions than that of acetone and the reactivity of triethylsilane and dimethylphenylsilane in hydrosilylation of benzaldehyde catalyzed by $RhCl(PPh_3)_3$ were almost similar.¹² From these results, the ratelimiting step in hydrosilylation of carbonyl compounds catalyzed by $RhCl(PPh_3)_3$ may be the stage of the addition of the silyl-rhodium complex to the carbonyl of the aldehyde or the ketones, since the steric hindrance of the carbonyl compounds effects on the reaction rate and that of hydrosilanes does not effect on the rate.

It was described in this chapter that the reaction of cyclchexene was catalyzed by Co₂(CO)₈ and not by RhCl(PPh₃)₃ and Ru3(CO)12. The hydroformylation reaction of cyclohexene catalyzed by Ru₃(CO)₁₂ has already been reported.⁸ These results suggest that in the reaction of an olefin with a hydrosilane and carbon monoxide catalyzed by RhCl(PPh3) and Ru3(CO)12, the reaction of the aldehyde with silyl-metal complexes is the rate-limiting step. This is the reason why RhCl(PPh3)3 and Ru3(CO)12 are not effective catalysts in the reaction of cyclohexene. In the reaction catalyzed by Co₂(CO)₈, the oxidative addition of a hydrosilane to the acylcobalt complex(eq. e) may be the rate-limiting step from the results of the reaction with various hydrosilanes. In the case of 1-methylcyclohexene, 2-diethylmethylsiloxyethenyl-1cyclohexane was not obtained. It has been reported that hydroformylation reaction of 1-methylcyclohexene gave

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cyclohexaneacetaldehyde.¹³ The result suggests that the oxidative addition of a hydrosilane to acylcobalt complexes is faster than that of hydrogen.

The relative reactivity of the cyclic olefins for hydroformylation has already been reported.¹⁴ The order was shown below. The order is similar to hydroboration with

 $A > O \simeq O > O > O$

disiamylborane¹⁵, the platinum(II) -catalyzed heterogeneous hydrogenation reaction¹⁶, and the palladium(II) -catalyzed olefin carbonylation reaction¹⁷. The fact suggests that the formation of the π -complex will be important in determining the overall yield of product, because the order is the same as that of ring strain.¹⁷ While the relative reactivity of cycloalkenes in the reaction of cycloalkenes with diethylmethylsilane and carbon monoxide catalyzed by Co₂(CO)₈ showed the alternative order as shown below. Therefore, the ratelimiting step is not the reaction of an olefin with HCo(CO)₄.

 $> \bigcirc > \bigcirc > \bigcirc > \bigcirc > \bigcirc >$

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1-3 Experimental

General comments

All temperature are uncorrected. IR spectra were recorded i di Null'Anglanta (su). Nationalità di Statu with a Shimazu IR-400. The H-NMR spectra were measured on a Japan Electron Optics JNM-PS-100 spectrometer with Me, Si as an internal standard. The position of Me, Si was recognized by adding it after recording spectrum without the standard. Otherwise the signal of the standard may be confused with that of a organosilicon compound. Mass spectra were recorded on a Hitachi mass spectrometer Model RMU-6E. High resolution mass spectra were recorded on a Hitachi RMU-7M. Analytical gas chromatography(GLC) were carried out on a Shimazu GC-3BF and a Shimazu GC-6A equipped with flame ionization detecters. The GLC columns used were as follows: (A) 6m X 3mm stainless steel containing 5% Silicon OV-1 supported on 60-80 mesh Uniport KS; (B) 6m X 3mm stainless steel containing 15% DEGS supported on 60-80 mesh cromosorb W. Preparative GLC was carried out using a Varian Aerograph Model 90-P gas chromatograph. Olefins were commercially available and were purified by distillation. Benzene was dried over sodium-lead alloy and stored over 5A Co₂(CO)₈ were prepared according to the molecular sieves. literature^{18|}. Ru₃(CO)₁₂ and RhCl(PPh₃)₃ were commercially available. Hydrosilanes were prepared by standard methods¹⁹. Authentic samples of disiloxanes were prepared by RhCl(PPh₂)₃ catalyzed silylation of water with appropriate hydrosilanes (benzene as solvent, reflux 48 h, quantitative yields).

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General procedure for the reaction of cycloalkenes with hydrosilanes and carbon monoxide in the presence of Co₂(CO)₈

A 100 ml stainless steel autoclave equipped with a : 300 kg/cm² pressure gauge and a magnetic stirring bar was charged with 30 mmol of cycloalkenes, 10 mmol of the hydrosilane, 0.2 mmol of Co₂(CO)_o and 20 ml of benzene. The autoclave was flushed $N_2(80 \text{ kg/cm}^2)$ and CO(50 kg/cm²), then charged with CO to SO kg/cm^2 . The reaction vessel was then heated with magnetic stirring in an oil bath. After 1 h, the pressure reached 105-108 kg/cm² at 140°C. The temperature was maintained at 140°C for 20 h. The vessel was then cooled to room temperature. The pressure decreased to 76-79 kg/cm². The gas pressure was slowly released and the autoclave was opened. Analysis of the reaction mixture by GLC (column A) with an internal standard (n-tridecane, n-tetradecane, n-pentadecane, or n-eicosane) showed it to contain the products (I and IV-VIII) and (R3Si)20. RRT(relative retention time) of the products to an internal standard is shown below for each case. RRT (relative retention time)=retention time of the compounds/ retention time of the standard. Distillation of the reaction mixture and/or preparative GLC afforded the analytical samples.

Diethylmethylsiloxymethylenecyclohexane(Ic)

Analysis by GLC(column A, 130°C, n-tetradecane as an internal standard(RRT 1)); Ic(71%, RRT 0.79) and (Et₂MeSi)₂O (8%, RRT 0.29). Ic; bp 130-138°C/28 torr); IR(neat) 2924, 1680, 1450, 1380, 1260, 1240, 1220, 1160, 1090, 1070, 1010, 970, 910,

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820, 800, 760 cm⁻¹; NMR(CCl₄) <u>6</u> 0.08(s 3H), 0.58(m 4H), 0.96(m 6H), 1.36-1.60(c 6H), 1.88(m 2H), 2.10(m 2H), 5.88 (s lH); mass spectrum m/e(rel. intensity) 212(33), 183(40), 101(9), 89(100).

Anal. Calcd for C₁₂H₂₄OSi: P⁺, 212.1595; C, 67.85; H, 1138. Found: P⁺, 212.1561; C, 67.80; H, 11.40.

Diethylmethylsiloxymethylenecyclopentane(IV)

Analysis by GLC(column A, 120°C, n-tridecane as an internal standard(RRT 1)); IV(48%, RRT 0.80) and $(\text{Et}_2\text{MeSi})_2^0$ (8% RRT 0.46). IV; bp 98-100°C/15 torr; IR(neat) 2950, 2875, 1685, 1460, 1415, 1250, 1170, 1120, 1005, 960, 950, 840-820, 800, 760 cm⁻¹; NMR(CCl₄) δ 0.08(s 3H), 0.60(m 4H), 0.96(m 6H), 1.58(m 4H), 2.12(m 4H), 6.04(s 1H); mass spectrum m/e(rel. intensity) 198(30), 169(39), 103(9), 89(100).

Anal. Calcd for C₁₁H₂₂OSi: P⁺, 198.1438; C, 66.59; H, 11.18. Found: P⁺, 198.1409; C, 66.35; H, 10.94.

Diethylmethylsiloxymethylenecycloheptane(V)

Analysis by GLC(column A, 140°C, n-tetradecane as an internal standard(RRT 1)); V(74%, RRT 1.29) and $(\text{Et}_2\text{MeSi})_2\text{O}$ (9%). V; bp 155-165°C/50 torr; IR(neat) 2950, 1660, 1450, 1260, 1190, 1160, 1130, 1000, 960, 870, 810, 760 cm⁻¹; NMR (CCl₄) δ 0.08(s 3H), 0.58(m 4H), 0.96(m 6H), 1.40-1.70(c 8H), 1.99(m 2H), 2.20(m 2H), 5.92(s 1H); mass spectrum m/e(rel. intensity) 226(13), 197(35), 101(11), 89(100).

Anal. Calcd for C₁₃H₂₆OSi: P⁺, 226.1750; C, 68.95; H, 11.57. Found: P⁺, 226.1731; C, 68.97; H, 11.70.

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Diethylmethylsiloxymethylenecyclooctane(VI)

Analysis by GLC(column A, 150°C, <u>n</u>-tetradecane as an internal standard(RRT 1)); VI(69%, RRT 1.96) and $(\text{Et}_2\text{MeSi})_2^O$ (11%). VI; bp 126-135°C(25 mm); IR(neat) 2920, 1660, 1480, 1250, 1235, 1200, 1160, 1090, 1060, 1000, 960, 860, 800, 770 cm⁻¹; NMR(CCl₄) δ 0.03(s 3H), 0.58(m 4H), 0.97(m 6H), 1.36-1.72(c 10H), 1.98(m 2H), 2.14(m 2H), 5.94(s 1H); mass spectrum m/e(rel. intensity) 240(13), 211(33), 121(21), 101(16), 89(100).

Anal. Calcd for C₁₄H₂₈OSi: P⁺, 240.1908; C, 69.92; H, 11.73. Found: P⁺, 240.1914; C, 69.88; H, 11.64.

<u>l-Diethylmethylsiloxymethylene-2-methylcyclohexane(VII)</u>

Analysis by GLC(column A, 130°C, <u>n</u>-tridecane as an internal standard(RRT 1)); VII (22%, RRT 1.64), unidentified compound(5%, RRT 1.52), and $(\text{Et}_2\text{MeSi})_2$ O (15%). The structure of the unidentified compound is not clear, but NMR spectrum of an 1:1 mixture(by GLC) of VII and it is not consistent with 2-diethylmethylsiloxyethenyl-1-cyclohexane. VII; bp 120-125°C(17 mm); IR(neat) 2950, 2900, 1680, 1460, 1260, 1150, 1100; 1010, 960, 880, 820, 800, 770 cm⁻¹; NMR(CCl₄) <u>&</u> 0.08 (s 3H), 0.60(m 4H), 0.88-1.20(m 6H)overlapped with 0.96(s 3H), 1.20-1.86(c 6H), 1.86-2.12(c 2H), 2.60-2.80(c 1H), 5.88(s 1H); mass spectrum m/e (rel. intensity) 226(11), 211(5), 197(11), 101(11), 89(100).

Anal. Calcd for C₁₃H₂₆OSi: P⁺, 226.1751; C, 68.95; H, 11.57. Found: P⁺, 226.1731; C, 68.59; H, 11.75.

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2-Diethylmethylsiloxymethylenenorbornene(VIII)

Analysis by GLC(column A, 150°C, <u>n</u>-tetradecane as an internal standard (RRT 1)); VIII(45%, RRT 1.71) and $(Et_2M=Si)_2O$ (7%). VIII; bp 130-140°C(10 mm); IR(neat) 2940, 2860, 1690, 1250, 1160, 1140, 1100, 1000, 850, 810, 750 cm⁻¹; NMR(CCl₄) δ 0.08(s 3H), 0.57(m 4H), 0.90(= 5H), 1.12-1.32(c 4H), 1.47(m 1H), 1.61(m 1H), 1.86(= 1H), 2.04(m 1H), 2.31(m 1H), 2.55(m 1H), 6.02(m 1H); In addition, small absorptions at 3.06 and 5.79 were also observed. The product may contain two stereo isomers. mass spectrum m/e (rel. intensity) 224(39), 195(100), 101 (44), 89(39).

Anal. Calcd for C₁₃H₂₄OSi: C, 69.57; H, 10.77. Found: C, 69.41; H, 10.91.

Dimethylethylsiloxymethylenecyclohexane(Ib)

Analysis by GLC(column A, 110°C, <u>n</u>-tridecane as an internal standard(RRT 1)); Ib(57%, RRT 0.70) and $(\text{EtMe}_2\text{Si})_2^0$ (5%, RRT 0.13)...Ib; bp 82-88°C(13 mm); IR(neat) 2910, 2850, 1680, 1450, 1380, 1250, 1150, 1080, 1060, 1000, 960, 900, 840, 780 cm⁻¹; NMR(CC1₄) δ 0.11(s 6H), 0.57(m 2H); 0.96 (m 3H), 1.35-1.59(c 6H), 1.88(m 2H), 2.10(m 2H), 5.84(s 1H); mass spectrum m/e (rel. intensity) 198(25), 169(22), 89(14), 75(100).

Anal. Calcd for C₁₂H₂₂OSi: P⁺, 198.1438; C, 66.59; H, 11.18. Found: P⁺, 198.1428; C, 66.87; H, 11.18.

Triethylsiloxymethylenecyclohexane(Id)

Analysis by GLC(column A, 135°C, <u>n</u>-pentadecane as an internal standard (RRT 1)); Id(74%, RRT 0.77) and $(\text{Et}_3\text{Si})_2$ O (23%, RRT 0.44). Id; bp 135-139°C(20 mm); IR(neat) 2920, 2895, 1675, 1445, 1260, 1240, 1220, 1150, 1090, 1010, 910, 820, 740 cm⁻¹; NMR(CCl₄) δ 0.70(m 6H), 1.09(m 9H), 1.44-1.74(c 6H), 2.01(m 2H), 2.22(m 2H), 6.02(s 1H); mass spectrum m/e (rel. intensity) 226(24), 197(41), 169(11), 103(100), 87(17), 75(55).

Anal. Calcd for C₁₃H₂₆OSi: C, 68.95; H, 11.57. Found: C, 69.05; H, 11.62.

Dimethylphenylsiloxymethylenecyclohexane(Te)

Analysis by GLC(column A, 160°C, <u>n</u>-pentadecane as an internal standard (RRT 1)); Ie(51%, RRT 1.97) and (PhMe₂Si)₂O (18%, RRT 2.19). Ie; bp 150-151°C(16 mm); IR(neat) 3050, 2910, 2850, 1680, 1590, 1430, 1250, 1200, 1150, 1120, 970, 830, 780, 720, 700 cm⁻¹; NMR(CCl₄) δ 0.39(s 6H), 1.35-1.59(c 6H), 1.86(m 2H), 2.16(m 2H), 5.88(s 1H), 7.26 (m 3H), 7.44(m 2H); mass spectrum m/e (rel. intensity) 246 (42), 231(4), 168(10), 153(19), 137(78), 135(100):

Anal. Calcd for C₁₅H₂₂OSi: P⁺, 246.1439. Found: P⁺, 246.1462.

Diphenylmethylsiloxymethylenecyclohexane(If)

Analysis by GLC(column A, 220°C, <u>n</u>-eicosane as an internal standard (RRT 1)); If(26%, RRT 1.65) and (Ph₂MeSi)₂O (6%, RRT 6.31). If; bp 150-156°C(0.55 mm); IR(neat); 3050,

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2910, 2850, 1680, 1590, 1430, 1250, 1140, 1120, 1080, 1060, 900, 830, 790, 720, 700 cm⁻¹; NMR(CCl₄) <u>δ</u> 0.63(s 3H), 1.53 (c 6H), 1.91(m 2H), 2.25(m 2H), 6.01(s 1H), 7.34(m 6H), 7.55 (m 4H); mass spectrum m/e (rel. intensity) 308(38), 215(33), 199(63), 197(100), 137(91).

Anal. Calcd for C₂₀H₂₄OSi: C, 77.87; H, 7.84. Found: C, 77.58; H, 7.68.

Trimethylsiloxymethylenecyclohexane(Ta)

A solution of cyclohexene(30 mmol) and $\text{Co}_2(\text{CO})_8$ (0.2 mmol) in 20 ml of benzene was placed in a 100 ml stainless steel autoclave. The autoclave was flushed with N₂(80 kg/cm²) and CO(50 kg/cm²) and cooled to -70°C. Trimethylsilane (10 mmol) was introduced to the autoclave by trap-to-trap distillation under vacume(10 mm) and then the autoclave was charged with CO to 80 kg/cm² at 25°C and heated at 140°C. The GLC retention time of the product was identical to that of trimethylsiloxymethylenecyclohexane which was independently prepared from cyclohexanecarboxaldehyde and trimethylchlorosilane $\frac{20}{20}$

Analysis by GLC(column A, 100°C, <u>n</u>-dodecane as an internal standard (RRT 1)); Ia(4%, RRT 0.65). Ia; bp 91°C (23 mm); IR(neat) 2950, 2850, 1675,1445, 1250, 1210, 1150, 1080, 900, 840, 740 cm⁻¹; NMR(CCl₄) $\underline{\delta}$ 0.14(s 9H), 1.44(c 6H), 1.84(m 2H), 2.04(m 2H), 5.80(s 1H); mass spectrum m/e (rel. intensity) 184(44), 169(42), 155(21), 75(100) 73(98).

Anal. Calcd for C₁₀H₂₀OSi: p⁺, 184.1282. Found: p⁺, 184.1279.

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l-(Diethylmethylsiloxymethyl)-cyclobex-l-ene(XI)

The experimental procedure was the same as the general procedure except that the reaction was carried out at 200°C.

Analysis by GLC(column A, 100°C, n-tetradecane as an internal standard (RRT 1)); XI(17%, RRT 0.81), Ic(67%, RRT 0.79), and $(\text{Et}_2\text{MeSi})_2$ O (15%, RRT 0.29). XI; IR(neat); 2920, 2870, 1675, 1460, 1420, 1255, 1160, 1090, 1070, 1010, 965, 840, 800, 760 cm⁻¹; NMR(CCl₄) δ 0.01(s 3H), 0.53(m 4H), 0.93 (m 6H), 1.60(m 4H), 1.80-2.12(c 4H), 3.85(s 2H), 5.50(m 1H); mass spectrum m/e (rel. intensity) 212(14), 183(31), 103(7), 89(100).

Anal. Calcd for C₁₂H₂₄OSi: P⁺, 212.1594; C, 67.85; H, 11.38. Found: P⁺, 212.1560; C, 67.83; H, 11.44.

General procedure for the reaction of 1-hexene with diethylmethylsilane and carbon monoxide in the presence of a metal complex

A solution of 3.7 ml(30 mmol) of 1-hexene, 1.02g(10 mmol)of diethylmethylsilane, and 0.136g(0.4 mmol) of $Co_2(CO)_8$ in 20 ml of benzene was placed in a 100 ml stainless steel autoclave. The autoclave was flushed with $N_2(50 \text{ kg/cm}^2)$ and $CO(50 \text{ kg/cm}^2)$, then charged with CO to 50 kg/cm² and heated at 140°C with stirring. After 20 h, the autoclave was cooled and depressurized. Analysis of the reaction mixture by GLC(Silicon OV-1, 5%, on Uniport KS, 6m, 115°C) with an internal standard(n-tetradecane, RRT=1) showed it to contain enol silyl ethers((Z)-(IXa); RRT 0.36, (E)-(IXa); RRT 0.39, (Z)-(Xa); RRT 0.45, (E)-(Xa); RRT 0.60) and (Et₂MeSi)₂O in 57%(total) and 5% yields, respectively.

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Analytical samples were obtained by fractional distillation (bp 105-115°C/12 torr) followed by repeting preparative GLC. The results under various reaction conditions were shown in Table VIII and IX.

The spectral data are: (Z)-(IXa), IR(heat) r=2950, 1670, 1460, 1260, 1170, 1090, 1010, 970, 750-800 cm⁻¹; NMR(CCl₄) <u> δ </u> 0.08(s 3H), 0.62(m 4H), 0.70-1.14(m 9H), 1.14-1.50(c 4H) overlaped with 1.48(d 3H), 2.00(m 2H) 5.90(s 1H); mass spectrum m/e (rel. intensity) 214(35), 199(6), 185(28), 171(50), 101(100); high resolution MS, parent ion m/e 214.1747 (calcd for C₁₂H₂₆OSi, 214.1751).

(E)-(IXa), IR(neat); 2920, 1670, 1460, 1260, 1170, 1100, 1010, 970, 740-800 cm⁻¹; NMR(CCl₄) δ 0.08(s 3H), 0.60(m 4H), 0.84-1.12(m 9H), 1.12-1.50(m 4H) overlaped with 1.50(d 3H), 1.83(m 2H), 5.92(m 1H); mass spectrum m/e (rel. intensity) (Z)-(Xa), IR(neat); 3050, 2950, 1660, 1460, 1400, 1260, 1170, 1140, 1090, 1010, 750-820 cm⁻¹; NMR(CCl₄) δ 0.10(s 3H), 0.60(m 4H), 0.80-1.15(m 9H), 1.15-1.40(c 6H), 1.96(m 2H), 4.30(dt, J=6 Hz and 8 Hz, 1H), 6:02(d, J=6 Hz, 1H); mass spectrum m/e (rel. intensity) 214(4), 185(76), 157(38), 101 (85), 89(100); high resolution MS, parent ion m/e 214.1728 $C_{12}H_{26}OSi$ Calcd 214.1751.

(E)-(Xa), IR(neat); 2920, 1660, 1460, 1420, 1260, 1160, 1100, 1010, 960, 910 cm⁻¹; NMR(CCl₄) δ 0.10(s 3H), 0.60(m 4H), 0.80-1.14(m 9H), 1.14-1.48(c 6H) 1.80(m 2H), 4.80(dt, J=12 Hz and 6Hz, 1H), 6.08(d, J=12 Hz, 1H); mass spectrum m/e (rel. intensity) 214(3), 185(54), 157(36), 101(66), 89(100); high

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•	ot a) is								8) E-Xa)	45	21 2	54	33	45	.58
0) 8 a	n (%)) (E-X	25	24	25	23	14		0)12	tion (-Xa) (44	35	31	48	37	28
co ₂ (C	butio (Z-Xa	43	44	43	48	32		Ru ₃ (C	tribu Xa) (Z						*. ·
e of	istri -IXa)	16	17	17	16	27		e of	t dis) (E-I	6	7	1	5	9	ω
cesenc	luct d Xa) (E							resenc	roduc z-IXa	4	2	ω	10	5	9
che Pr	Prod (Z-I	16	E F	Ч	Н	27		the P1	otal H ield (40	16	30	22	33	59
0 in t	tal eld(%)	57	52	38	21	22		h ti L ui	л. т	20 4	20	50	20	50	44
and C(ти лт то' Уі́€	00	00	44	50	50		and C(remp. (°C)	140	140	140	200	140	140
SiH :	emp. h	40	40	L40 /	L40	500		SiH S	cm ²)						
Et2M€	e Te			• •				Et2Me	CO (kg/c	50	80	80	80	80	30
with	essur fin.	49	70	79	80	78		with	Izene	0	0	0	0	0	0
xene	CO Pr inít.	50	80	80	80	80		xene	2 Ber (n			4	7		
1-не	zene 1)	0	0	0	0	0		1-не	(CO) ₁ mol)	.4	.05	-	4.		
on of	Ben (m	8	4 7	4	4	4	nmo1)	on of	Ru ₃ (m	0	0	0			
leactio	SiH (mmol)	10	20	20	20	, 30	(0.4 1	keactic	SiH (mmol)	10	10	20	30	20	15
VIII F	lefin mol)	30	60	20	20	20	² (co) ^ε	IX	lefin mol)	30	30	20	205	20	10
Table	Run O. (r	1	7	Ω.	4.	ហ	a) Co	Table	Run O.		2	ო	4	ы	6

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resolution MS, parent ion - Et m/e 185.1346 C₁₀H₂₁OSi Calcd 185.1359.

Reaction of methyl acrylate with diethylmethylsilane and carbon monoxide-

A solution of 2.7 ml(30 mmol) of methyl acrylate, 1.02g (10 mmol) of Et_2MeSiH , and 0.068g(0.2 mmol) of $Co_2(CO)_8$ in 20 ml of benzene was placed in a 100 ml stainless steel autoclave. The autoclave was flushed two times with $CO(50 \text{ kg/cm}^2)$, then charged with CO to 80 kg/cm² and heated at 140°C with stirring. After 20 h, the autoclave was cooled and depressurized. Analysis of the reaction mixture by GLC (Silicon OV-1, 5%, on Uniport KS, 3m, 120°C) with an internal standard(n-tetradecane, RRT=1) showed it to contain (Z) - and (E) -1-diethylmethylsiloxy-2carbomethoxyprop-l-ene(IXb) and (Z)- and (E)-L-diethylmethylsiloxy-3-carbomethoxyprop-1-ene(Xb)((Z)-(IXb); RRT 0.633, (E)-(IXb); RRT 0.744, (Z)-(Xb); RRT 0.625, (E)-(Xb); RRT 0.706) and (Et₂MeSi)₂O in 49%(total) and 7% yields, respectively. Analytical samples were obtained by fractional distillation (130-160°C/17 torr) followed by preparative GLC. The results under various reaction conditions were shown in Table III and Table V.

The spectral data are described below.

(Z)-(IXb), IR(neat); 2950, 2860, 1725, 1700, 1640, 1300, 1260, 1190, 1130, 820, 770 cm⁻¹; NMR(CCl₄) δ 0.17(s 3H), 0.68(m 4H), 1.00(m 6H), 1.69(s 3H), 3.65(s 3H), 6.53(s 1H); mass spectrum (rel. intensity) 216(2), 201(6), 187(100), 159(23), 103(90), 91(35), 89(35), 75(52); high resolution MS parent ion - Et m/e 187.0804 C₈H₁₅O₃Si calcd 187.0789.

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(E)-(IXb), IR(neat); 2920, 1720, 1640, 1300, 1220, 1120, 820, 760 cm⁻¹; NMR(CCl₄) δ 0.22(s 3H), 0.74(m 4H), 1.00(m 6H), 1.70 (d, J=2 Hz, 3H), 3.64(s 3H), 7.35(q, J=2 Hz, 1H); mass spectrum (rel. intensity) 216(2), 201(7), 187(100), 103(95), 75(87); high resolution MS parent - Et ion m/e 187.0821 C₈H₁₅O₃Si calcd 187.0790.

(Z) - (XD), IR(neat); 2950, 2870, 1740, 1660, 1330, 1260, 1160, 1120, 1040, 1010, 840, 750 cm⁻¹; NMR(CCl₄) δ 0.15(s 3H), 0.66 (m 4H), 1.00(m 6H), 3.03(d, J=8 Hz, 2H), 3.60(s 3H), 4.59(dt, J=8 Hz and 8 Hz, 1H), 6.15(d, J=8 Hz, 1H); mass spectrum (rel. intensity) 216(3), 187(86), 157(20), 103(100), 101(43), 75(48), 73(41); high resolution MS parent -Et ion m/e 187.0802 C₈H₁₅O₃Si calcd 187.0790.

(E)-(Xb), The NMR prectrum of (E)-(Xb) was determined from the mixture of (E)-(Xb) and (E)-(IXb). NMR(CCl₄) δ 0.14(s), 0.51-0.81(m), 0.90-1.11(m), 2.79(d, J=8 Hz), 3.58(s), 4.89(dt, J=12 Hz and 8 Hz), 6.15(d, J=12 Hz).

Reaction of cyclohexene with diethylmethyldeuteriosilane and carbon monoxide

A solution of 10.3 ml(100mmol) of cyclohexene, 1.03g (10 mmol) of Et_2MeSiD , and 0.068g(0.2 mmol) of $\text{Co}_2(\text{CO})_8$ in 20 ml of benzene was placed in a 100 ml stainless steel autoclave. The autoclave was flushed with N₂(50 kg/cm²) and CO(50 kg/cm²), then charged with CO to 50 kg/cm² and heated at 140°C with stirring. After 1 h, the pressure reached 105 kg/cm² at 140°C. The temperature was maintained at 140°C for 20 h. The vessel was then cooled to room temperature. The

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pressure decreased to 77 kg/cm². The gas pressure was slowly released and the autoclave was opened. Analysis of the reaction mixture by GLC(Silicon OV-1, 3m, 130°C) with an internal standard(n-tetradecane) showed it to contain diethylmethylsiloxymethylenecyclohexane(Ic) and (Et₂MeSi)₂O in 87% and 12% yields, respectively. Analytical samples were obtained by fractional distillation(124-129°C/20 torr) followed by preparative GLC.

The amount of deuterium incorporation was determined by comparing the area in the vinyl and the methylene of cyclohexane part with the methyl group of siloxy part. These areas were measured five times at least with the largest amptitude and averaged. The result was shown in Table X.

	MeSi	CH ₂ C=COSi C	CH ₂ C=C ^{OS}	C=C NH	CH2 ^{CH2} CH2 ^{CH2}
ratio of areas	3	1.88	1.82	0.088	5.80
deuterium content(%)	-	7.5	7.5	91	3

Table X Comparison of Areas in NMR Spectrum

From the Table X, the amount of deuterium incorporation was determined to be 91% at the vinyl position, 15% at the adjacent methylene group to carbon-carbon double bond, and 3% at the another methylene group of the cyclohexane part.

Mass spectrometric analysis showed it to contain 12% d_0 , 75% d_1 , 11% d_2 , 2% d_3 . The second trial was carried out under the same conditions described above except that 30 mmol of cyclohexene and 80 kg/cm² of carbon monoxide were used. The result was shown in Table XI.

	MeSi CH	OSi C=C	CH ₂ C=C	$C=C_{H}^{OSi}$ $CH_{2}^{CH_{2}}$
ratio of areas	3	1.93	1.73	0.18 -
deuterium content(%)	-	3.5	13.5	82 –

Table XI Comparison of Areas in NMR Spectrum

Mass spectrometric analysis showed it to contain 18% d_0 , 61% d_1 , 18% d_2 , 4% d_3 .

Reaction of n-heptanal with diethylmethylsilane in the presence of carbon monoxide and Co₂(CO)₈

A solution of n-heptanal(10 mmol), diethylmethylsilane (10 mmol), and $\text{Co}_2(\text{CO})_8(0.4 \text{ mmol})$ in benzene(20 ml) was placed in a 100 ml stainless steel autoclave. The autoclave was flushed two times with N₂(50 kg/cm²), then charged with CO (50 kg/cm²) and heated at 140°C with stirring. After 20 h, the autoclave was cooled and depressurized. Analysis of the reaction mixture by GLC(Silicon OV-1, 5%, 6m, 110°C) with an internal standard(n-tetradecane) showed it to contain X and XVI in 32.8% and 7.2% yields, respectively. The IR(neat) and NMR(CCl₄) data of XVI are: IR, 2930, 1470, 1420, 1255, 1100, 1010, 800, 760; NMR δ 0.00(s 3H), 0.32(m 4H), 0.92(m 9H), 1.28(c 10H), 3.50(t, J=6 Hz, 2H). Reaction of n-heptanal with diethylmethylsilane in the presence of $Co_2(CO)_8$

A solution of diethylmethylsilane(4 mmol) and $Co_2(CO)_8$ (0.04 mmol) in xylene(4 ml) was heated at 135°C. To the solution was added a solution of n-heptanal(2 mmol) in xylene. Gas evolution occurred immediately. Analysis of the reaction mixture by GLC(Silicon OV-1, 5%, 6m, 140°C) with an internal standard(n-tetradecane) showed it to contain (Z)-(Xa),(E)-(Xa), and XVI in 49%, 30%, and 21% yields, respectively. The results at various temperature were shown in Table XII.

Table XII Yields and Distribution at Various temperature^a

Temp.(°C)	Yield(Xa + XVI)	Xa :	XVI
10	94	4:	96
28	100	6:	94
50	100	18:	82
80	76 ^C	53:	47
110 ^b	93	64:	36
135 ^b	100	79:	21

a) In benzene unless otherwise noted.

b) In xylene. c) Result after 1 h.

Reaction of styrene with diethylmethylsilane and carbon monoxide in the presence of $Co_2(CO)_8$

A solution of styrene(30 mmol), diethylmethylsilane(10 mmol), and $\text{Co}_2(\text{CO})_8(0.2 \text{ mmol})$ in benzene(20 ml) was placed in a 100 ml stainless steel autoclave. The autoclave was flushed with N_2 (80 kg/cm²) and CO(50 kg/cm²), then charged with CO(80 kg/cm²) and heated at 140°C with stirring. After 20 h, the autoclave

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was cooled and depressurized. Analysis of the reaction mixture by GLC(Silicon OV-1, 5%, 3m, 140°C) with 0.3 g of n-tridecane as an internal standard showed it to contain eight compounds (XVIIa;RRT 1.86, XVIIb;RRT 2.21, XVIIc;RRT 2.54, XVIId;RRT 3.00, XVIIe; RRT 3.44, XVIIf; RRT 5.00, XVIIg; RRT 7.47, and (Et₂MeSi)₂0). Detector response calibrations are not determined except for XVIIa. The ratios of the peak areas of the compounds and the standard were obtained as follows: XVIIa; 0.32(yield 6%), XVIIb; 0.20, XVIIc; 0.13, XVIId; 0.17, XVILe; 0.45, XVIIf; 0.10, XVIIg; 0.27. Analytical samples were obtained by fractional distillation(110-120°C/17 torr) followed by preparative GLC. The product(XVIIa) was identified with (2-phenylethyl) diethylmethylsilane; NMR(CCl_A) δ -0.04(s 3H), 0.51(m 4H), 1.04(m 8H), 2.55(m 2H), 7.05(s 5H). The product (XVIIb) was identified with β -diethylmethylsilylstyrene; IR (neat), 2940, 2860, 1600, 1570, 1490, 1250, 1000, 980, 790, 730 cm⁻¹; NMR(CCl_A) δ 0.10(s 3H), 0.63(m 4H), 0.99(m 6H), 6.29 (d, J=20 Hz, 1H), 6.81(d, J=20 Hz, 1H), 7.05-7.35(c 5H). An exact mass determination gave m/e 204.1303(calcd for C13H20Si, 204.1333).

Reaction of styrene in the presence of $Ru_3(CO)_{12}$

The experimental procedure was the same as that described above except that styrene(10 mmol), diethylmethylsilane(20 mmol), Ru₃(CO)₁₂(0.05 mmol), and n-tridecane(0.2g) as an internal standard were used. The ratios of the peak areas of the compounds and the standard were obtained as follows: XVIIa; 0.11 (1.4%), XVIIb; 0.29, XVIIc; 0.07, XVIId; 0.02, XVIIe;0.14, XVIIf; 0.04, XVIIg; 0.03.

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Reaction of acrylonitrile with diethylmethylsilane and carbon monoxide in the presence of $Ru_3(CO)_{12}$

A solution of acrylonitrile(10 mmol), diethylmethylsilane (30 mmol), and Ru₃(CO)₁₂(0.01 mmol) in benzene(20 ml) was placed in a 100 ml stainless steellautoclave. The autoclave was flushed with N₂(80 kg/cm²) and CO(50 kg/cm²), then charged with \mathbb{C} (80 kg/cm²) and heated at 140°C with stirring. After 20 h, the autoclave was cooled and depressurized. Analysis of the reaction mixture by GLC(DEGS, 15%, on Cromosorb W, 6m, 130°C) with an internal standard(n-pentadecane) showed it to contain 1-cyanoethyl(diethylmethyl)silane in 56% yield. The IR(neat) and NMR(CCl₄) data are: 2950, 2860, 2220, 1460, 1260, 1010, 790, 750 cm⁻¹; δ 0.14(s 3H), 0.69(m 4H), 1.02(m 6H), 1.26(d, J=8 Hz, 3H), 1.65(q, J=8 Hz, 1H). An exact mass determination gave m/e 155.1125(calcd for C₈H₁₇NSi, 155.1128).

Competitive reaction of cycloalkenes

In a typical procedure, a solution of 0.62g(7.5 mmol) of cyclohexene, 0.83g(7.5 mmol) of cyclooctene, 0.75 ml(5 mmol) of diethylmethylsilane, and 0.1 mmol of $Co_2(CO)_8$ in 10 ml of benzene was placed in a 100 ml stainless steel autoclave. The autoclave was flushed two times with $CO(50 \text{ kg/cm}^2)$, then charged with $CO(50 \text{ kg/cm}^2)$ and heated at 140 °C with stirring. After 2 h, the autoclave was cooled and depressurized. Analysis of the reaction mixture by GLC(Silicon OV-1, 5% on Uniport KS, 6m, 150°C) showed it to contain (Ic) and (VI) in 5.5% and 4.3% yields, respectively. The results were shown in Table XIII.

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Table	XIII	Compet.	ıtıve	Reaction	or Cy	cloalkenes	•
Run	hr	Y	ield o	f enol s	ilyl e	ther	Yield of
				\bigcirc	\bigcirc	A	(Et ₂ MeSi) ₂ O
1	20	3.6	6.9	1 			99
2	20		31.	47			18
3	20	•	31	•	23	•	22
4	20	6.1			3.6		29
5 ^b	2		6	12.4	•		8.2
6 ^b	, 2		5.5		4.3	.	5
7^{b}	3	34			28		9
8 ^b	2	4				24	5

a) Reaction conditions; Co₂(CO)₈(0.2 mmol), olefin
 (30 mmol 1:1), Et₂MeSiH(10 mmol), CO(50 kg/cm²), 140°C,
 20 h, benzene(20 ml) unless otherwise noted.

b) Reactions were carried out in a half scale and olefins were weighted in gram unit.

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Experimentally, to a solution of LiAlD₄(18 mmol) in 15 ml of diethyl ether was added dropwise a solution of Et₂MeSiCl (74 mmol) in 80 ml of diethyl ether for 20 min at room temperature and then the solution was heated to reflux for l hr. Diethylmethyldeuteriosilane was obtained in 59% yield.

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Chapter 2 A Catalyzed Reaction of Cyclic Ethers with a

Hydrosilane and Carbon Monoxide

2-1 Introduction

In chapter 1, the new catalyzed reaction of olefins with hydrosilanes and carbon monoxide was described. To know the scope and limitation to the substrate is very important in order to explore the utility of the reaction. From this point of view, the catalyzed reaction of cyclic ethers with a hydrosilane and carbon monoxide was examined. It was found that when tetrahydrofuran (THF) was used as a solvent in the reaction of olefins, the product obtained was not the expected one, enol silyl ether, but a new compound. This is the reason why cyclic ethers were used as a substrate.

2-2 Results and discussion

The reaction of cyclic ethers (tetrahydrofuran, oxetane, and cyclohexeneoxide) with diethylmethylsilane and carbon monoxide was catalyzed by $\text{Co}_2(\text{CO})_8$ to give diethylmethylsiloxyaldehyde (I), (II), (III), respectively, in reasonable yields; (I) 53%, (II) 40%, and (III) 51% as shown in eq. 1, 2, and 3.

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 $Et_2^{MeSiH} + CO - \frac{Co_2(CO)_8}{CO_2(CO)_8}$ Et,MeSiO

(I)

(1)



The reaction of cyclic ethers except for oxetane was: carried out using 50 mmol of cyclic ethers, 10 mmol of diethylmethylsilane, and 60 kg/cm² of carbon monoxide at 140°C for 20 h. The reaction conditions in the case of oxetane were the same as that described above except that 20 mmol of oxetane were used at 70°C.

Hydroformylation of cyclic ethers is known to give hydroxy aldehydes as the initial products¹. The reaction, however, frequently suffers from undesirable side reactions or further reactions, since the hydroxy aldehydes are very labile compounds. For example, hydroformylation² of tetrahydrofuran did not afford the expected 4-hydroxybutanal but yielded various products arising from further carbonylation. as shown in eq. 4 and hydroformylation³ of cyclohexeneoxide gave only condensation product of the expected 2-hydroxycyclohexanecarboxaldehyde as shown in eq. 5. While the reactions using a hydrosilane show a direct approach to the synthesis of pre-protected hydroxy aldehydes. Synthetically, functionalyzed aldehydes are very useful in elongation of carbon skeltone.

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It is interesting to note that Chalk has reported the polimerization of tetrahydrofuran catalyzed by Co2(CO)8 and triethylsilane(1:2) and proposed ring opening of tetrahydrofuran by a silylcobaltcarbonyl for the initiation step⁴. Corriu and co-workers have reported that a silylcobaltcarbonyl containing an optically active silicon reacted with methanol and triethylamine and the cleavage of the Si-Co bond took place with retention². It is reasonable that the silicon group of the silylcobaltcarbonyl electrophilically attacked the oxygen of methanol. These results suggest that the ring opening of tetrahydrofuran by the silylcobaltcarbonyl is the initial step. This means that one can make a carbon cobalt bond utilyzing the high affinity of silicon to oxygen as a driving force. This is the new method to make bonds of carbon and transition metals. The reasonable reaction path was shown in scheme 1.

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2-3 Experimental

Reaction of tetrahydrofuran with diethylmethylsilane and: carbon monoxide in the presence of $Co_2(CO)_8$

A solution of 4 ml(50 mmol) of tetrahydrofuran, 1.02g (10 mmol) of diethylmethylsilane, and 0.068g(0.2 mmol) of Co₂(CO)₈ in 20 ml of benzene was placed in a 100 ml stainless steel autoclave. The autoclave was flushed with $N_2(60 \text{ kg/cm}^2)$ and CO(60 kg/cm²), then charged with CO(60 kg/cm²) and heated at 140°C with stirring. After 20 h, the autoclave was cooled and depressurized. Analysis of the reaction mixture by GLC (Silicon OV-1, 5%, on Uniport KS, 3m, 135°C) with n-tetradecane as an internal standard showed it to contain 5-(diethylmethylsiloxy)-n-pentanal (I) in 53% yield. An analytical sample of (I) was obtained by fractional distillation(bp 120-130°C/ 30 torr) followed by preparative GLC(Silicon OV-1, 5%, on Uniport KS, 3m X 8mm, 140°C). The spectral data are shown below. (I), IR(neat), 2950, 2870, 1720, 1250, 1090, 1010, 800, 760 cm^{-1} ; NMR(CCl₄) δ 0.02(s 3H), 0.56(m 4H), 0.96(m 6H), 1.58 (m 4H), 2.38(m 2H), 3.56(t, J=6 Hz, 2H), 9.60(m 1H):

Anal. Calcd for C₁₀H₂₂O₂Si: C, 59.34; H, 10.96. Found: C, 59.20; H, 11.10.

Reaction of oxetane with diethylmethylsilane and carbon monoxide

The reaction conditions were the same as that described above except that 20 mmol of oxetane were used and the temperature was 70°C. The spectral data are shown below.

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Analysis of the reaction mixture by-GLC(Silicon OV-1, 5%, on Uniport KS, 3m, 110°C) with n-tridecane as an internal standard showed it to contain 4-(diethylmethylsiloxy)-n-butanal(II) in 40% yield.

(II), IR(neat), 2920, 2860, 1725, 1480, 1460, 1250, 1080, 1000, 790, 750 cm⁻¹; NMR(CCl₄) δ 0.03(s), 0.51(m), 0.93(m), 1.79(m), 2.43(=), 3.57(t, J=6), 9.64(m).

Anal. Calcd for C₈H₂₀O₂Si: C, 57.39; H, 10.70. Found: C, 57.35; H, 10.79.

Reaction of cyclohexeneoxide with diethylmethylsilane and carbon monoxide

The experimental procedure was the same as that in the case of tetrahydrofuran. Analysis of the reaction mixture by GLC(Silicon OV-1, 5%, on Uniport KS, 3m, 160°C) with n-tetradecane as an internal standard showed it to contain 2-(diethylmethylsiloxy)cyclohexanecarboxaldehyde(III) and an identified compound.

(III), RRT; 1.30, bp 165-166/30 torr, IR(neat), 2950, 2870, 1725, 1460, 1250, 1155, 1110-1060, 1010, 960, 800, 760 cm⁻¹; NMR(CCl₄) <u>&</u> 0.05(s 3H), 0.56(m 4H), 0.95(m 6H), 1.11-1.50 (c 4H), 1.50-2.04(c 4H), 2.04-2.40(c 1H), 3.76(td, J=10 Hz and 4 Hz, 1H), 9.66(d, J=2 Hz, 1H); mass spectrum(rel. intensity) 199(18), 110(24), 89(100), 81(31), 61(88).

Anal. Calcd for C₁₂H₂₄O₂Si: C, 63.10; H, 10.59. Found: C, 62.88; H, 10.75.

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2-4 References and Footnotes

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Chapter 3 A Catalyzed Reaction of Aldehydes with a Hydrosilane and Carbon Monoxide

3-1 Introduction

It was shown that the new catalyzed reaction using hydrosilane and carbon monoxide combination converts olefins $(C=C) \implies$ siloxy olefins in chapter 1 and cyclic ethers(C-O-C) to siloxy aldehydes in chapter 2. It is interesting question whether aldehydes(C=O) can be used as a substrate in this new reaction. In chapter 2, the new method making a carbon cobalt bond utilyzing the high affinity of silicon to oxygen as a driving force was also described. From these points of view, the catalyzed reaction of aldehydes with a hydrosilane and carbon monoxide was examined.

3-2 Results and discussion

The reaction of n-heptanal with diethylmethylsilane(three fold excess to the aldehyde) and carbon monoxide in the presence of $Co_2(CO)_8$ and PPh₃ gave a mixture of (Z) - and (E)-1,2-bis-(diethylmethylsiloxy)-1-octene(Id) and unidentified compounds(II) in 66% and 14% yields, respectively.

 $Co_2(CO)_8$ -PPh₃ OSiEt,Me Et, MeSiQ 2Et₂MeSiH RCH со (1)R' (I)

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Aldehyde	Yield of	(I) Yield of (II)	(Et2 ^{MeSi)} 20
propanal	(Ia)25	8	3
isobutanal	(Ib)21	1	2
butanal	(Ic)67	12	
n-heptanal	(Id)66	14	
cyclohexane- carboxaldehyd	le ^{(Ie)37}	5	2
n-hexanal	(If)54	12	_

Table I Reaction of Aldehydes with Et₂MeSiH and CO^a

a) Reaction conditions; aldehyde(10 mmol), Et₂MeSiH(30 mmol) Co₂(CO)₈(0.2 mmol), PPh₃(0.2 mmol), CO(50 kg/cm²), 140°C, 20 h, benzene(20 ml).

The use of PPh_3 as a co-catalyst is necessary to prevent undesired hydrosilylation of the aldehyde.¹ In the case of cyclohexanecarboxaldehyde, the yield of (Ie) was low because of the formation of diethylmethylsiloxymethylenecyclohexane(61%) by dehydrogenative silylation. Interestingly, the similar reaction(i. e., using five fold excess of the hydrosilane, but in this case without PPh₃) of tetrahydrofuran was found to give the same type of compound(Ig) in 89% yield. The intermediate may be 5-(diethylmethylsiloxy)-n-pentanal which was the product when the reaction was carried out using five fold excess of tetrahydrofuran over the silane and without PPh₃ as shown in eq. 2.

OSiEt₂Me Et₂MeSiO Et,MeSiO (2) OSiEt,Me

(Ig)

The fact that the reaction of tetrahydrofuran with five fold excess of the hydrosilane gave the corresponding 1,2bissiloxyalkene in high yield suggests that $HCo(CO)_4$ is not the active catalyst in the reaction of cyclic ethers with a hydrosilane and carbon monoxide catalyzed by $Co_2(CO)_8$ in chapter 2.

Hydroformylation (oxo reaction) of aldehydes has been known to give formates as shown in eq. 3.² Thus the carbon atom of carbon monoxide is connected with the oxygen atom of aldehyde carbonyl. Contrastly, the carbon atom of carbon monoxide is linked with the carbon atom of aldehyde carbonyl in the reaction using a hydrosilane as shown in eq. 4.³



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The observed dramatic change may be attributed to the difference of reactivities of $HCO(CO)_4$ and $R_3SiCO(CO)_4$. Consequently, a carbon cobalt bond was formed utilyzing the high affinity of silicon to oxygen as a driving force.

3-3 Experimental

Reaction of n-heptanal with diethylmethylsilane and carbon monoxide in the presence of $Co_2(CO)_8$ and PPh₃

A solution of 1.14 g(10 mmol) of n-heptanal, 4.4 ml (30 mmol) of diethylmethylsilane, 0.068 g(0.2 mmol) of Co₂(CO)₈, and 0.052 g(0.2 mmol) of PPh₃ in 20 ml of benzene was placed in a 100 ml stainless steel autoclave. The autoclave was flushed with $N_2(50 \text{ kg/cm}^2)$ and CO(50 kg/cm²), then charged with CO(50 kg/cm²) and heated at 140° C with stirring. After 20 h, the autoclave was cooled and depressurized. Analysis of the reaction mixture by GLC (Silicon OV-1, 5% on Uniport KS, 3m, 170°C) with n-cetane as an internal standard showed it to contain (Id) in 66% yield. In addition to Id there were at least two unidentified products (IId) (total ca. 14%) at longer retention time. An analytical sample of (Id) (a mixture of (Z) and (E) isomers) was obtained by fractional distillation(bp 120-123°C/0.5 torr) followed by preparative GLC (Silicon OV-1, 5%, on Uniport KS, 3m × 8mm, 190°C).

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1,2-Bis(diethylmethylsiloxy)-l-octene(Id)

Analysis by GLC(Silicon OV-1, 170°C, n-cetane as an internal standard (RRT 1)); Id(RRT 2.04), bp 120-123°C/ 0.5 torr, IR(neat) 2950, 2870, 1685, 1460, 1250, 1160, 1090, 1000, 800, 760 cm⁻¹; NMR(CCl₄) <u>6</u> 0.08(s 3H), 0.10(s 3H), 0.63(m 8H), 0.80-1.11(m 15H), 1.29(c 8H), 1.80(m 2H), 5.44(s 0.83H), 5.90(s 0.17H):

Anal. Calcd for C₁₈H₄₀O₂Si₂: C, 62.72; H, 11.67. Found: C, 62.60; H, 11.71.

1,2-Bis(diethylmethylsiloxy)-l-pentene(Ic)

Analysis by GLC(Silicon OV-1, 140°C, n-tetradecane as an internal standard (RRT 1)); Ic(RRT 2.0), bp 130-150°C/25 torr, IR(neat) 2940, 2860, 1685, 1460, 1360, 1250, 1150, 1000, 850, 790, 750 cm⁻¹; NMR(CCl₄) <u>6</u> 0.08(s 3H), 0.10(s 3H), 0.32-0.80 (m 8H), 0.84-1.08(m 15H), 1.26-1.60(m 2H), 1.78(t, J=7 Hz, 2H), 5.44(s 0.92H), 5.92(s 0.08H); mass spectrum(rel. intensity) 302(43),273(25), 189(43), 161(54), 101(100), 73(78), 45(29).

Anal. Calcd for C₁₅H₃₄O₂Si₂: C, 59.54; H, **11**.32. Found: C, 59.21; H, 11.55.

1,2-Bis(diethylmethylsiloxy)-l-cyclohexylethylene(Ie)

Analysis by GLC(Silicon OV-1, 170°C, n-heptadecane as an internal standard (RRT 1)), Ie(RRT 1.9), bp 140-150°C/0.8 torr, IR(neat) 2920, 1685, 1250, 1210, 1160, 1120, 1010, 970, 860, 840, 760-800 cm⁻¹; NMR(CCl₄) <u>§</u> 0.09(s 3H), 0.12(s 3H), 0.48-0.75(m 8H), 0.87-1.08(m 12H), 1.11-1.41(c 5H), 1.41-1.86(c 6H), 5.48(s 0.91H), 5.78(s 0.09H); mass spectrum(rel. intensity)

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342(100), 313(38), 189(54), 161(73), 101(88), 73(87), 45(33). Anal. Calcd for C₁₈H₃₈O₂Si₂: C, 63.09; H, 11.17. Found: C, 62.99; H, 11.21.

1,2-Bis(diethylmethylsiloxy)-l-heptene(If)

Analysis by GLC(Silicon OV-1, 6m, 180°C, n-pentadecane as an internal standard (RRT 1)); If, bp 117-119°C/1.0 torr, IR (neat) 2950, 1690, 1470, 1420, 1260, 1210, 1160, 1100, 1000, 840, 800, 770 cm⁻¹; NMR(CCl₄) δ 0.08(s 3H), 0.09(s 3H), 0.62 (m 8H), 0.87-1.08(m 15H), 1.17-1.56(c 6H), 1.80(m 2H), 5.42 (s 0.87H), 5.88(s 0.13H); mass spectrum(rel. intensity) 330(50), 301(17), 189(40), 161(53), 101(100), 73(90), 45(30).

Anal. Calcd for C₁₇H₃₈O₂Si₂: C, 61.75; H, 11.58. Found: C, 61.48; H, 11.69.

1,2-Bis(diethylmethylsiloxy)-3-methyl-1-butene(Ib)

Ib, bp 150-160°C/30 torr, IR(neat) 2950, 2870, 1685, 1460, 1250, 1180, 1000, 970, 790, 760 cm⁻¹; NMR(CCl₄) δ 0.08(s), 0.10(s), 0.60(m), 0.80-1.08(m), 1.98(m), 5.44(s), 5.70(s).

Anal. Calcd for C₁₅H₃₄O₂Si₂: C, 59.54; H, **11**.32. Found: C, 59.59; H, 11.46.

1,2-Bis(diethylmethylsiloxy)-l-butene(Ia)

Ia, IR(neat) 2950, 2870, 1685, 1460, 1250, 1160, 1005, 795, 760 cm⁻¹; NMR(CCl₄) δ 0.09(s 3H), 0.12(s 3H), 0.63(m 8.2H), 0.92(m 16H), 1.83(q, J=10 Hz, 1.5H), 5.42(s 0.75H); mass spectrum (rel. intensity) 288(41), 259(23), 189(35), 161(46), 101(100), 73(79), 45(26).

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Anal. Calcd for C₁₄H₃₂O₂Si₂: C, 58.26; H, 11.17. Found: C, 58.27; H, 11.26.

1,2-Bis(diethylmethylsiloxy)-4-methyl-l-pentene(IIc)

(IIc), NMR(CCl₄) <u>δ</u> 0.08(s 3H), 0.10(s 3H), 0.66(m 8H), 0.93-1.14(m 18H), 1.65-1.74(m 2H), 1.74-2.07(c 1H), 5.48(s 1H); mass spectrum(rel. intensity) 316(31), 287(14), 273(28), 189(31), 161(44), 101(100), 89(38), 73(79), 61(32), 45(31).

Anal. Calcd for C₁₆H₃₆O₂Si₂: C, 60.69; H, 11.46. Found: C, 60.43; H, 11.55.

3-4 References and Footnotes

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- 3 When the reaction was carried out using excess aldehyde, 1,2-bissiloxyalkene was not obtained. The products have not been identified yet, but one of them may be α -siloxy aldehyde from IR and mass spectra. This result suggests that in the reaction of olefins described in chapter 1, the key intermediate is not a siloxycarbene complex but an aldehyde.

The results obtained from the present work may be summarised as follows;

(1) Hydrogenation is related to hydrosilylation in a sence that hydrogen in the former is replaced by a hydrosilane in the latter. However, hydroformylation has no such a counterpart. In the present work, a new catalyzed reaction using hydrosilanes and carbon monoxide related to hydroformylation(oxo reaction) of an olefin has been found.

H-H	
hydrogenation	hydrosilylation
$C=C + H_2 \xrightarrow{\longrightarrow} C \xrightarrow{\longrightarrow} C \xrightarrow{\longrightarrow} L \xrightarrow{\longrightarrow} H \xrightarrow{\longrightarrow} H$	C=C + HSi→C—C H Si
hydroformylation	present work
$C=C + CO + H_2 \rightarrow C - C$ H $C=O$ H	$C=C + CO + HSi \rightarrow C-C=C-OSi$

(2) It is suggested that the initial stage of the new reaction is similar to that of hydroformylation (oxo reaction) and the key intermediate is an aldehyde.

(3) The range of the applicability of the new reaction has been extended to cyclic ethers and aldehydes, cyclic ethers give siloxyaldehydes and aldehydes yield bis-siloxyolefins. These reactions were a series of new catalyzed ones using hydrosilanes and carbon monoxide. In the case of olefins, $HCo(CO)_4$ may be the active catalyst, while in the cases.of cyclic ethers and aldehydes, $R_3SiCo(CO)_4$ may play an important

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role as the active catalyst. Interestingly, either of $HCo(CO)_4$ or $R_3SiCo(CO)_4$ formed by the reaction of R_3SiH with $Co_2(CO)_8$ becomes the active catalyst according to the nature of the substrates.

$$C=C + R_{3}SiH + CO \xrightarrow{HCo(CO)_{4}} C-C=C-OSiR_{3}$$

$$H$$

$$C=C + R_{3}SiH + CO \xrightarrow{R_{3}SiCo(CO)_{4}} R_{3}SiO-C-C-C-H$$

 $\begin{array}{c} 0 \\ H \\ R^{T}CH + R_{3}SIH + CO \end{array} \xrightarrow{R_{3}SICO(CO)_{4}} R_{3}SIO_{R'}C=C \\ H \end{array}$

These products (siloxyalkenes, siloxyaldehydes, and 1,2-bissiloxyalkenes) are important intermediate of organic syntheses. It is to be hoped that a series of new catalyzed reactions comparable with hydroformylation (oxo reaction) will be established on the occasion when the scope and limitation of the new catalyzed reactions using hydrosilanes and carbon monoxide are completely revealed.

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