

Platinum-Catalyzed Reductive Aldol and Michael Reactions

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For the Pt-catalyzed nucleophilic addition of enones, Pt complexes were employed in the presence of various phosphine ligands and H₂ (or Et₃SiH), affording inter- and intra-molecular coupling products in good to modest yield. Depending on reaction protocols, different phosphine ligands were required to optimize the conditions. In the aldol reaction, the Pt catalyst involving P(2,4,6-(OMe)₃C₆H₂)₃ or P(*p*-OMeC₆H₄)₃ was chosen. Michael reaction proceeds in good yields in the presence of P(*p*-CF₃C₆H₄)₃. Regarding the activity of the reductants, H₂ exhibited superior activity to Et₃SiH, resulting in a shorter reaction time and higher yield in the aldol and Michael reaction. In light of the deuterium labeling studies, the catalytic cycle including the hydrometalation of the enones by the platinum hydride species was proposed.

Key Words: Platinum chloride, Reductive aldol reaction, Reductive Michael reaction

Introduction

Platinum complexes have received a great deal of attention as powerful catalysts for the synthesis of a diverse range of molecules including natural building blocks.¹ Pt-catalyzed coupling protocols can be divided into two categories; 1) the reactions of Pt bound alkenes (alkynes) with nucleophiles and 2) the reactions of Pt bound alkenes (alkynes) with electrophiles (Scheme 1). In the Pt-catalyzed cycloisomerization of enynes and the hydroalkoxylation (hydroamination) of alkenes, Pt complexes pull the electron density from the π -bond of alkenes or alkynes upon coordination. As a result, the Pt bound alkenes or alkynes become electron-deficient, thereby allowing them to react with nucleophiles.² In contrast, Pt complexes in the presence of silanes or hydrogen exhibit different reactivity toward the π -bond.^{3,4} In the Pt-catalyzed hydroformylation of alkenes, Pt complexes react with the H₂ and alkenes, affording Pt-alkyl complexes, which undergo migratory insertion to carbon monoxide.⁵ Considering the remarkable activity of Pt complexes toward electron deficient functional groups during the hydroformylation, Pt complexes with suitable reductants are expected to conduct the reductive coupling of alkenes (alkynes) and electron deficient functional groups.⁶ Hence, we embarked on a study of reductive coupling reactions utilizing PtCl₂ with electronically different phosphine ligands in the presence of reductants.

In the case of Rh, Ir, Pd, and Ni complexes, the reductive couplings of alkenes and alkynes through their hydrometalation are well known.^{7,8} In particular, activated alkenes (enones) undergo hydrometalation at the alkene of the enone in the presence of transition metal catalysts and reductants, affording hydrometalated products (metal enolates). Transition metal

enolates participate in the successive addition to activated alkenes (Michael addition), aldehydes (aldol addition), and imines (Mannich reaction), etc.^{9,11} Among the available reductants, hydrogen and silanes have emerged as promising ones, since reductive coupling reactions using these reagents provide high selectivity of the products and functional group tolerance. In this sense, the reductive couplings of enones in the presence of hydrogen and silanes constitute one of the most active areas of transition metal catalysis.^{10,11}

With the aim of developing platinum catalyzed protocols for the reductive coupling of enones with electrophiles (aldehydes and enones), we explored various platinum complexes with hydrogen and silanes to obtain the corresponding reductive coupling products. Previously, the Pt-catalyzed hydrogenative Michael cyclization of bis-enones and related mechanistic studies were conducted.¹² In this account, we discuss Pt-catalyzed intra- and inter-molecular reductive aldol reactions where both hydrogen and the silane are utilized as a hydride source. Deuterium labeling experiments were conducted to confirm the proposed catalytic cycle involving the hydrometalation of the enones and subsequent trapping by the aldehyde. Moreover, the Pt complexes showing good catalytic activity in the aldol reaction were applied to silane-mediated Michael cyclization of bis-enones.

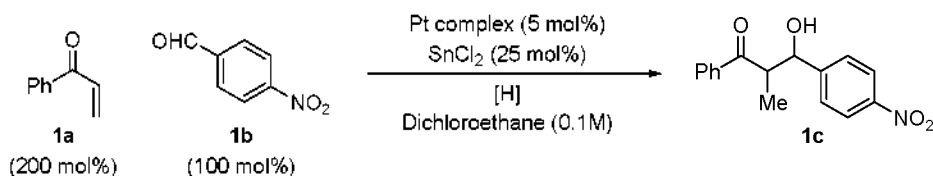
Results and Discussion

In an effort to improve both the diastereoselectivity and reactivity of the catalyst, various phosphine ligands were screened and a low reaction temperature was employed. Table 1 summarizes the results. The reactivity and selectivity of the aldol reaction catalyzed by PtCl₂ with P(*p*-CF₃C₆H₄)₃ and P(*p*-OMeC₆H₄)₃ were previously reported (Table 1, entries 1 and 2).¹² The Pt complex involving the electron-rich phosphine appears to promote the hydrogen-mediated reductive aldol reaction in good yield. P(furyl)₃ and 1,2-bis(diphenylphosphino)ethane (dppf) showed diminished yields with poor selectivity (Table 1, entries 3 and 4). The Pt catalyst involving the electron-rich P(2,4,6-(OMe)₃C₆H₂)₃ exhibited an increased



Scheme 1. Pt-catalyzed cyclization

Table 1. Optimization of the reductive aldol reaction



Entry	Pt complex	Ligand	[H]	Temp	Yield(<i>syn:anti</i>)
1	PtCl ₂ (5 mol%)	P(<i>p</i> -CF ₃ C ₆ H ₄) ₃ (5 mol%)	H ₂ (1 atm)	80 °C	50% (45:55)
2	PtCl ₂ (5 mol%)	P(<i>p</i> -OMeC ₆ H ₄) ₃ (5 mol%)	H ₂ (1 atm)	80 °C	83% (57:43)
3	PtCl ₂ (5 mol%)	P(furyl) ₃ (5 mol%)	H ₂ (1 atm)	80 °C	57% (48:52)
4	PtCl ₂ (5 mol%)	dppe (5 mol%)	H ₂ (1 atm)	80 °C	16% (58:42)
5	PtCl ₂ (5 mol%)	P(2,4,6-(OMe) ₃ C ₆ H ₂) ₃ (5 mol%)	H ₂ (1 atm)	80 °C	91% (62:38)
6	PtCl ₂ (5 mol%)	P(2,4,6-(OMe) ₃ C ₆ H ₂) ₃ (10 mol%)	H ₂ (1 atm)	80 °C	91% (84:16)
7	PtCl ₂ (5 mol%)	P(2,4,6-(OMe) ₃ C ₆ H ₂) ₃ (5 mol%)	H ₂ (1 atm)	50 °C	36% (42:58)
8	PtCl ₂ (5 mol%)	P(<i>p</i> -CF ₃ C ₆ H ₄) ₃ (5 mol%)	Et ₃ SiH (150 mol%)	50 °C	65% (74:26)
9	PtCl ₂ (5 mol%)	P(<i>p</i> -OMeC ₆ H ₄) ₃ (5 mol%)	Et ₃ SiH (150 mol%)	50 °C	66% (50:50)
10	PtCl ₂ (5 mol%)	P(furyl) ₃ (5 mol%)	Et ₃ SiH (150 mol%)	50 °C	64% (73:27)
11	PtCl ₂ (5 mol%)	P(2,4,6-(OMe) ₃ C ₆ H ₂) ₃ (5 mol%)	Et ₃ SiH (150 mol%)	50 °C	21% (79:21)

yield (91%) along with improved diastereoselectivity (Table 1, entry 5). Next, the amount of P(2,4,6-(OMe)₃C₆H₂)₃ was increased to 10 mol% (Table 1, entry 6). Pleasingly, in this case, the ratio of *syn* to *anti* products was increased (*syn:anti* = 84:16), while maintaining a similar chemical yield (Table 1, entry 6). In addition to ligand screening, the temperature was lowered to 50 °C, with the expectation that the selectivity of the *syn* and *anti* products would be improved. As shown in entry 7 of Table 1, neither the chemical yield nor the diastereoselectivity was increased. Throughout the optimization procedure, 10 mol% of P(2,4,6-(OMe)₃C₆H₂)₃ or 10 mol% of P(*p*-OMeC₆H₄)₃ with 5 mol% of PtCl₂ and 25 mol% of SnCl₂ was selected as the catalyst of the hydrogenative aldol reaction.

Based on the results of the hydrogenative aldol reaction, Pt complexes containing P(*p*-CF₃C₆H₄)₃, P(*p*-OMeC₆H₄)₃, P(furyl)₃, and P(2,4,6-(OMe)₃C₆H₂)₃ were utilized in the silane-mediated aldol reaction. Since silanes have been utilized in various reductive coupling reactions, Pt complexes with Et₃SiH would be expected to catalyze the reductive aldol reaction. As

indicated in entries 8-10 of Table 1, the silane-mediated reductive aldol reactions using P(*p*-CF₃C₆H₄)₃, P(*p*-OMeC₆H₄)₃, and P(furyl)₃ provided the products in similar yield (65%, 66%, and 64%), but with different diastereoselectivities. Unlike the hydrogenative aldol reaction, employing P(2,4,6-(OMe)₃C₆H₂)₃ diminished the yield of the silane-mediated aldol reaction (Table 1, entry 11). In addition to phosphine ligand screening, a higher temperature (80 °C) was applied, providing the product in diminished yield. As an optimal catalyst for the silane-mediated reductive aldol reaction, 5 mol% of PtCl₂ and 25 mol% of SnCl₂ were applied in the presence of suitable phosphine ligands. Depending on the substrates, different phosphine ligands were required to obtain a good yield.

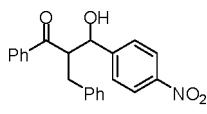
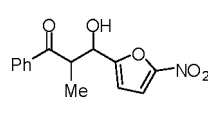
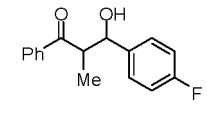
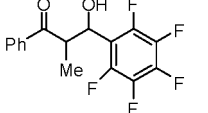
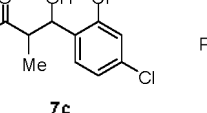
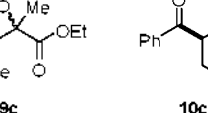
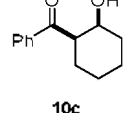
With appropriate sets of reaction conditions, we investigated a range of carbonyl compounds and enones to show the substrate scope of the hydrogen-mediated aldol reactions. The results are shown in Table 2. Initially, β -substituted enones were evaluated in the hydrogen-mediated intermolecular reductive aldol reaction (Table 2, compounds 2c and

Table 2. The substrate scope of the reductive aldol reaction using H₂

46% ^a (<i>syn:anti</i> = 55:45) ^a 51% ^b (<i>syn:anti</i> = 40:60) ^b	31% ^a (<i>syn:anti</i> = 62:40) ^a 75% ^b (<i>syn:anti</i> = 15:85) ^b	68% ^a (<i>syn:anti</i> = 73:27) ^a 99% ^b (<i>syn:anti</i> = 60:40) ^b	46% ^a (<i>syn:anti</i> = 75:25) ^a 71% ^b (<i>syn:anti</i> = 68:32) ^b	24% ^a (<i>syn:anti</i> = 60:40) ^a 75% ^b (<i>syn:anti</i> = 15:85) ^b
60% ^a (<i>syn:anti</i> = 50:50) ^a 79% ^b (<i>syn:anti</i> = 51:49) ^b	20% ^a (<i>syn:anti</i> = 83:17) ^a 94% ^b (<i>syn:anti</i> = 79:21) ^b	trace ^a 53% ^b	67% ^a 58% ^b	

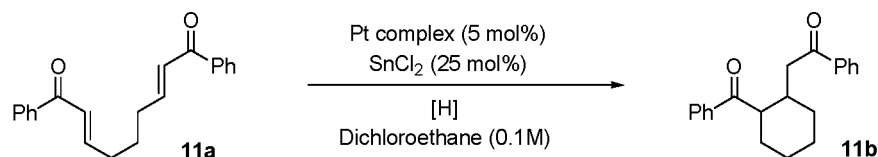
^aP(2,4,6-(OMe)₃C₆H₂)₃ (10 mol%), ^bP(*p*-OMeC₆H₄)₃ (10 mol%). In the case of compound 10c, 5 mol% of phosphine ligand was used.

Table 3. The substrate scope of the reductive aldol reaction using Et₃SiH

		
27% ^a (<i>syn:anti</i> = 70:30) ^a trace ^b	29% ^a (<i>syn:anti</i>) ^a 42% ^b (<i>syn:anti</i> = 64:36) ^a	
		
trace ^a 48% ^b (<i>syn:anti</i> = 58:42) ^b	72% (<i>syn:anti</i> = 5:95) ^a trace ^b	
		
67% ^a (<i>syn:anti</i> = 76:24) ^a 27% ^b (<i>syn:anti</i> = 42:58) ^b	43% ^a 19% ^b	60% ^a 35% ^b

^aP(*p*-OMeC₆H₄)₃; ^bP(*p*-CF₃C₆H₄)₃

3c). Upon the exposure of chalcone **2a** and nitrobenzaldehyde **1b** to the reaction conditions using P(2,4,6-(OMe)₃C₆H₂)₃ and P(*p*-OMeC₆H₄)₃, **2c** was formed in 46% and 51% yields, respectively, indicating that the substituent at the β-position of the enone strongly retarded the hydrometalation of the enone and the following coupling reaction to **1b**. The formation of compound **3c** illustrated that the β-substituted aliphatic enone participated in this reaction to form the desired product in low yield, which might be associated with the steric effect of the octyl group at the β-position as well as the electronic effect of the methyl group proximal to the carbonyl group. In addition to acyclic enones, the reaction of cyclohexenone with *p*-nitrobenzaldehyde **1b** did not provide the coupling product.

Table 4. Optimization of reductive Michael cyclization

Entry	Pt complex	Ligand	[H]	Temp	Yield(<i>syn:anti</i>)
1	PtCl ₂ (5 mol%)	P(<i>p</i> -CF ₃ C ₆ H ₄) ₃ (5 mol%)	H ₂ (1 atm)	80 °C	69% (22:78)
2	PtCl ₂ (5 mol%)	P(<i>p</i> -OMeC ₆ H ₄) ₃ (5 mol%)	H ₂ (1 atm)	80 °C	98% (33:67)
3	PtCl ₂ (5 mol%)	PPh ₃ (5 mol%)	H ₂ (1 atm)	80 °C	76% (26:74)
4	PtCl ₂ (5 mol%)	P(<i>p</i> -OMeC ₆ H ₄) ₃ (5 mol%)	H ₂ (1 atm)	80 °C	66% (28:72)
5	PtCl ₂ (5 mol%)	P(2,4,6-(OMe) ₃ C ₆ H ₂) ₃ (5 mol%)	H ₂ (1 atm)	80 °C	79% (35:65)
6	PtCl ₂ (5 mol%)	P(furyl) ₃ (5 mol%)	H ₂ (1 atm)	80 °C	94% (38:62)
7	PtCl ₂ (5 mol%)	dppe (5 mol%)	H ₂ (1 atm)	80 °C	NR
8	PtCl ₂ (5 mol%)	P(<i>p</i> -CF ₃ C ₆ H ₄) ₃ (5 mol%)	Et ₃ SiH (150 mol%)	50 °C	71% (29:71)
9	PtCl ₂ (5 mol%)	P(<i>p</i> -CF ₃ C ₆ H ₄) ₃ (5 mol%) ^a	Et ₃ SiH (150 mol%)	50 °C	24% (50:50)
10	PtCl ₂ (5 mol%)	P(<i>p</i> -OMeC ₆ H ₄) ₃ (5 mol%)	Et ₃ SiH (150 mol%)	50 °C	52% (25:75)

A variety of aldehydes were evaluated with phenylvinylketone **1a** in the reductive aldol reaction using hydrogen. The heterocyclic aldehyde **4b** participated in this reaction to provide **4c** in 68% yield in the presence of P(2,4,6-(OMe)₃C₆H₂)₃. By replacing the phosphine ligands with P(*p*-OMeC₆H₄)₃, the yield of **4c** was dramatically increased, while the diastereoselectivity was slightly lowered. Aromatic aldehydes possessing electron withdrawing groups exhibit improved yields (**5c–7c**) under the conditions employing P(*p*-OMeC₆H₄)₃ rather than P(2,4,6-(OMe)₃C₆H₂)₃. Especially, compound **6c** showed significant diastereoselectivity changes in the presence of P(*p*-OMeC₆H₄)₃. The aliphatic aldehyde, octanal **8b** was also coupled to phenylvinylketone **1a** to afford **8c** in 94% yield under the conditions using P(*p*-OMeC₆H₄)₃. In addition to the aldehyde, the activated ketone, ethyl pyruvate was tested as a carbonyl partner. P(*p*-OMeC₆H₄)₃ promoted the hydrogenative coupling reaction of ethyl pyruvate and phenylvinylketone to afford **9c** as a single diastereomer in good yields, while the Pt catalyst with P(2,4,6-(OMe)₃C₆H₂)₃ formed trace amounts of the products. In most intermolecular reactions, under the reaction conditions involving P(*p*-OMeC₆H₄)₃ rather than P(2,4,6-(OMe)₃C₆H₂)₃, higher yields were observed. In the case of **2c** and **6c**, remarkable diastereoselectivity changes were detected upon changing the phosphine ligands.

Unlike the intermolecular aldol reaction, the intramolecular aldol reaction proceeds to afford **10c** as a single diastereomer. Presumably, the equilibrium between *E* and *Z* forms of the Pt-enolate and the free rotation of the tethered aldehyde might be restricted, providing high selectivity. Both P(*p*-OMeC₆H₄)₃ and P(2,4,6-(OMe)₃C₆H₂)₃ were employed under 1 atmosphere of hydrogen, affording **10c** in 67% and 58% yields, respectively.

In parallel to the hydrogenative aldol reactions, the silane-mediated aldol reactions of various enones and aldehydes were performed (Table 3). As the catalyst, 5 mol% of PtCl₂ and 25 mol% of SnCl₂ were employed in the presence of 5 mol% of P(2,4,6-(OMe)₃C₆H₂)₃ or P(*p*-CF₃C₆H₄)₃. Upon the exposure of chalcone to the reductive aldol conditions, the

desired product **2c** was formed in 27% yield. In the case of the nitro-substituted furfuraldehyde and 4-fluorobenzaldehyde, the corresponding aldol products (**4c** and **5c**, respectively) were formed in modest yield. In contrast, multihalogen substituted benzaldehydes required an electron-rich phosphine ligand to obtain a higher yield (**6c** and **7c**). In addition to the aldehyde, the activated ketone (ethyl pyruvate) reacts with phenylvinylketone to afford **9c** as a single diastereomer. As illustrated in **10c**, the intramolecular aldol cyclization occurs to provide the desired product in 60% yield. Overall, compared to the hydrogenative aldol reaction, the silane-mediated aldol reaction exhibited a lower chemical yield despite the prolonged reaction time.

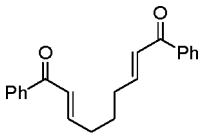
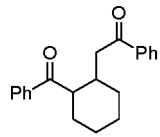
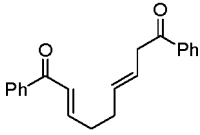
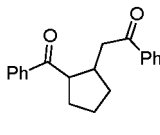
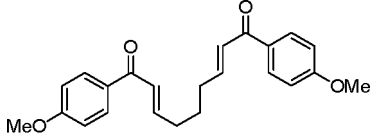
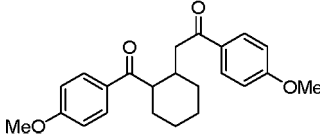
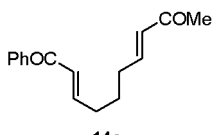
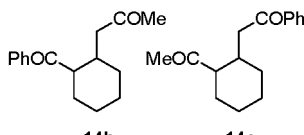
We previously reported the results of the reductive Michael reaction in the presence of electronically different phosphines and the bidentate phosphine (Table 4). Under hydrogenation conditions using 5 mol% of $P(C_6F_5)_3$, $P(p-CF_3C_6H_4)_3$, PPh_3 , and $P(p-OMeC_6H_4)_3$, the reductive cyclizations of (2E,7E)-1,9-diphenylnona-2,7-diene-1,9-dione **11a** were previously presented (Table 4, entries 1-4).¹² Subsequently, electronically and sterically different phosphine ligands were tested in an attempt to improve the diastereoselectivity and chemical yield. The Pt complex with $P(2,4,6-(OMe)_3C_6H_2)_3$ provided the product in 79% yield as a mixture of the *syn* and *anti* products (*syn:anti* = 35:65) (Table 4, entry 5). The Pt catalyst involving $P(furyl)_3$ exhibited good activity, comparable to that of $P(p-CF_3C_6H_4)_3$ but low diastereoselectivity (Table 4, entry 6). The activity of the bidentate phosphine ligand (dppe) was inferior to that of the monodentate phosphines (Table 4, entry 7). Throughout the optimization of the phosphine ligands, reasonable yields were obtained using the Pt catalysts involving $P(p-CF_3C_6H_4)_3$, PPh_3 and $P(furyl)_3$. Subsequently,

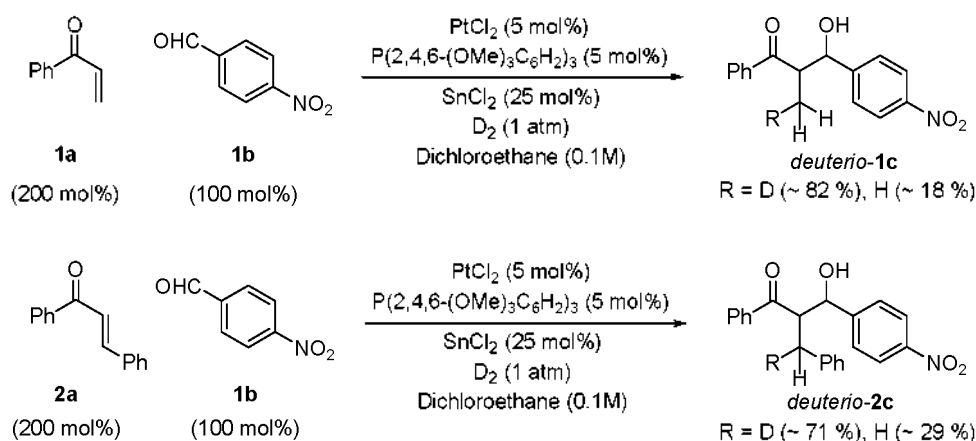
to improve the diastereoselectivity, a range of solvents (dichloromethane, acetone, diethyl ether, CH_3CN , THF, and benzene) were applied. Under the conditions involving those solvents, desired reductive cyclization products were observed in modest to good yields, but the diastereoselectivity was not increased.

Next, reductive Michael reactions were conducted using Et_3SiH as a reductant. The optimal catalyst for the hydrogenative Michael cyclization was applied. Using $PtCl_2$ (5 mol%), $P(p-CF_3C_6H_4)_3$ (5 mol%), $SnCl_2$ (25 mol%), and Et_3SiH (150 mol%), compound **11a** was converted to **11b** in 71% yield as a mixture of the *syn* and *anti* isomers (*syn:anti* = 29:71) (Table 4, entry 8). Similar to the hydrogenative Michael reaction, in the absence of $SnCl_2$, the catalytic activity of the Pt complex significantly decreases to provide the cyclization product in 24% yield (Table 4, entry 9). In the presence of the electron-rich phosphine $P(p-OMeC_6H_4)_3$, product **11b** was formed in 52% yield (Table 4, entry 10). Throughout the screening of the phosphine ligands, the combination of $PtCl_2$, $P(p-CF_3C_6H_4)_3$, and $SnCl_2$ was chosen for the reductive Michael cyclization of the bis-enones.

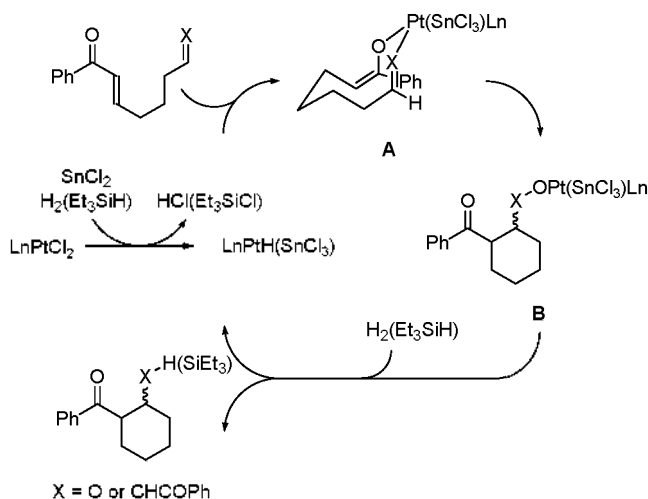
To explore the scope of the substrates, a variety of bis-enones were evaluated under the standard reductive cyclization conditions involving 5 mol% of $PtCl_2$, 25 mol% of $SnCl_2$, 5 mol% of $P(p-CF_3C_6H_4)_3$, and 150 mol% of Et_3SiH (Table 5). Compared to the hydrogenative Michael cyclizations forming **11b-14c**,¹² with the silane as a reductant, the catalytic activity appears to be lower. The reductive cyclization of **12a** provided the 5-membered ring **12b** in 57% yield with a mixture of the *syn* and *anti* products (*syn:anti* = 1:1). The electron-rich aromatic bis-enone **13a** was converted to **13b** in 89% yield with 1:4 ratio of the *syn:anti* products.

Table 5. Silane-mediated reductive Michael cyclization

Entry	Substrate	Product	Yield (<i>syn:anti</i>)
1			71 % (1:2.5)
2			57 % (1:1)
3			89 % (1:4)
4			20 % (1:1) 14b:14c = 4:1



Scheme 2. Deuterium labeling experiments



Scheme 3. A proposed catalytic cycle

However, the unsymmetrical bis-enones **14a** exhibited much lower yields than the symmetric aromatic bis-enones. The symmetric aliphatic bis-enones and bis-enoates were not converted to the desired product.

With the aim of providing a better understanding of the platinum-catalyzed reductive coupling mechanism, deuterium labeling experiments were conducted. In D_2 -mediated intermolecular aldol reaction, the degree of deuterium incorporation at the β -position of the ketone was clearly determined by 1H NMR. Deuterium labeling studies of the aldol reaction were conducted using phenylvinylketone **1a** and chalcone **2a** with *p*-nitrobenzaldehyde **1b**. Upon the exposure of compounds **1a** and **1b** to the hydrogenation conditions using D_2 , 82% of the hydrogen at the methyl group of *deuterio-1c* was exchanged with deuterium. In the case of *deuterio-2c* formed from chalcone **2a**, 71% of deuterium labeling was observed at the methylene. These results imply that the reversible hydrometalation of the platinum hydride to the enone occurs prior to the C-C bond formation, affording a mixture of the deuterated and non-deuterated aldol products.

The catalytic cycle for the Pt-catalyzed reductive Michael reaction was previously proposed.¹² Here, we propose a similar mechanism for the Pt-catalyzed reductive aldol reaction.

The catalytic cycle begins with the generation of $LnPtH(SnCl_3)$ from $PtCl_2$, phosphine, $SnCl_2$, and H_2 (or Et_3SiH). In Pt-catalyzed hydrogenation and hydroformylation, $SnCl_2$ is known to function as a co-catalyst.^{4,5} Generally, the chloride of the Pt complex reacts with $SnCl_2$ to afford $SnCl_3^-$. The role of the strong π -acceptor $SnCl_3^-$ is speculated to promote the desired coupling and to prevent the reduction of Pt(II) complexes during the reaction.^{5c} Upon its exposure to the reaction mixture containing $LnPtH(SnCl_3)$, the enone undergoes hydrometalation to form intermediate **A**, as indicated by the deuterium labeling results. The subsequent addition of the Pt-enolate **A** to the aldehyde forms the intermediated **B**. Finally, the oxidative addition of hydrogen or the silane to the intermediate **B** and reductive elimination afford the product with the concomitant generation of $LnPtH(SnCl_3)$.

Conclusion

We have developed the catalytic nucleophilic addition of Pt-enolates to aldehydes or enones in the presence of H_2 or Et_3SiH . Using $PtCl_2$ with phosphine ligands and reductants, various enones, aldehydes and bis-enones participated in this reaction to afford the corresponding products in good to modest yield. Noticeably, under reductive aldol reaction conditions, in most cases, electron-rich phosphines ($P(p-OMeC_6H_4)_3$ and $P(2,4,6-(OMe)_3C_6H_2)_3$) are favored as ligands. On the other hand, the reductive Michael reaction requires $P(p-CF_3-C_6H_4)_3$ for good conversion. Accordingly, tuning the electronic properties of the Pt and phosphine complexes is vital for the good conversion of the reactions involving different substrates. In terms of the reactivity of the reductants, the hydrogen-mediated aldol and Michael reactions exhibited significantly higher yields than the silane-mediated reactions. To understand the mechanism, deuterium labeling experiments were carried out using enones and aldehydes, supporting the formation of $LnPtH$ and its subsequent hydrometalation to the enone. Our methodology involving Pt catalysts and reductants (H_2 and Et_3SiH) demonstrates the nucleophilic activity of Pt-enolates complexes to the electron-deficient functional groups, which is an exciting extension of the catalytic activity of Pt complexes in organic reactions.

Experimental Section

General. All reactions were run under an atmosphere of argon, unless otherwise indicated. Anhydrous solvents were transferred by an oven-dried syringe. Flasks were flame-dried and cooled under a stream of nitrogen. Dichloroethane was distilled from calcium hydride. Analytical thin-layer chromatography (TLC) was carried out using 0.2-mm commercial silica gel plates (DC-Fertigplatten Kieselgel 60 F₂₅₄). Preparative column chromatography employing silica gel was performed according to the method of Still.¹³ Solvents for chromatography are listed as volume/volume ratios. Proton nuclear magnetic resonance (¹H NMR) spectra were recorded with a Mercury (400 MHz) spectrometer. Chemical Shifts are reported in delta (δ) units, parts per million (ppm) downfield from trimethylsilane. Coupling constants are reported in Hertz (Hz). Carbon-13 nuclear magnetic resonance (¹³C NMR) spectra were recorded with a Mercury 400 (100 MHz) spectrometer. Chemical shifts are reported in delta(δ) units, parts per million (ppm) relative to the center of the triplet at 77.00 ppm for deuteriochloroform. ¹³C NMR spectra were routinely run with broadband decoupling. Product **1c**,¹² **5c**,^{11a} **9c**,¹⁴ **10c**,^{11a} **11b-14c**¹² exhibited spectral properties consistent with previous literature reports. Spectral characterization of new compounds **2c-4c** and **6c-8c** are available in supplementary data.

Representative Experimental Procedure for Hydrogen-Mediated Reductive Aldol Reaction. To the premixed solution of PtCl₂ (5 mol%), P(2,4,6-(OMe)₃C₆H₂)₃ (10 mol%), and SnCl₂ (25 mol%), in dichloroethane (0.1 M) under H₂ (1atm) were added 200 mol% of **1a** and 100 mol% of **1b** at room temperature. The resulting mixture was allowed to run at 80 °C for 2 hours. The solvent was removed with a rotary evaporator to produce a residue which was purified by column chromatography on a silica gel eluting with hexane and ethyl acetate.

Representative Experimental Procedure for Triethylsilane-Mediated Reductive Aldol Reaction. To the premixed solution of PtCl₂ (5 mol%), P(*p*-OMeC₆H₄)₃ (5 mol%), SnCl₂ (25 mol%), and Et₃SiH (150 mol%) in dichloroethane (0.1 M) were added 200 mol% of **1a** and 100 mol% of **1b** at room temperature. The resulting mixture was allowed to run at 50 °C for 24 hours. The solvent was removed with a rotary evaporator to produce a residue which was purified by column chromatography on a silica gel eluting with hexane and ethyl acetate.

Representative Experimental Procedure for Hydrogen-Mediated Reductive Michael Cyclization. To the premixed solution of PtCl₂ (5 mol%), P(*p*-CF₃C₆H₄)₃ (5 mol%), and SnCl₂ (25 mol%), in dichloroethane (0.1 M) under H₂ (1atm) were added 100 mol% of **11a** at room temperature. The resulting mixture was allowed to run at 80 °C for 2 hours. The solvent was removed with a rotary evaporator to produce a residue which was purified by column chromatography on a silica gel eluting with hexane and ethyl acetate.

Representative Experimental Procedure for Triethylsilane-Mediated Reductive Aldol Reaction. To the premixed solution of PtCl₂ (5 mol%), P(*p*-CF₃C₆H₄)₃ (5 mol%), SnCl₂ (25 mol%),

and Et₃SiH (150 mol%) in dichloroethane (0.1 M) were added 100 mol% of **11a** at room temperature. The resulting mixture was allowed to run at 50 °C for 24 hours. The solvent was removed with a rotary evaporator to produce a residue which was purified by column chromatography on a silica gel eluting with hexane and ethyl acetate.

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Supporting Information. Experimental procedures and spectral characterization of compounds **2c-4c** and **6c-8c** are available at the bkcs website (<http://www.kcsnet.or.kr/bkcs>).

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