

# Construction of Highly Functionalized Diazoacetoacetates via Catalytic Mukaiyama–Michael Reactions

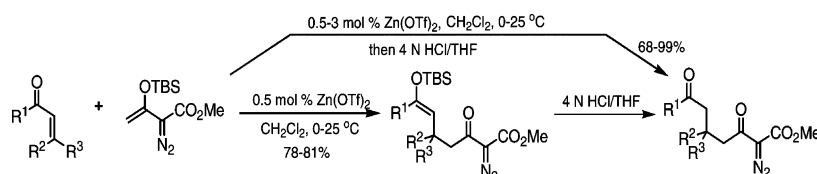
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## ABSTRACT



Functionalized diazo acetoacetates are prepared by an efficient Mukaiyama–Michael reaction between methyl 3-(trialkylsiloxy)-2-diazo-3-butenate and  $\alpha,\beta$ -unsaturated enones. Vinyl ether and ketone derivatives are both accessible in good to excellent yield through this methodology. The mild Lewis acid zinc(II) triflate is the optimal catalyst, and its loading can be as low as 0.1 mol %. In addition, zinc triflate was also found to be a superior catalyst for the related Mukaiyama–aldol reaction.

Metal-catalyzed diazo decompositions of diazoacetoacetate derivatives are among the most well-known reactions for the construction of cyclopropane, cycloalkanone, and  $\beta$ -lactam moieties.<sup>1,2</sup> The advantages of diazoacetoacetate compounds compared to diazoacetates include their higher thermal and acid stability and greater reaction control during diazo decomposition.<sup>1</sup> Traditionally, these diazo compounds were prepared by multistep synthesis involving diazo transfer as the final step.<sup>3,4</sup> Alternative methods were developed which involve the base-mediated condensation of diazo compounds with various electrophiles.<sup>5,6</sup> However these reactions generally require stoichiometric amounts of the reagents under

relatively harsh reaction conditions. Although diazo compounds are known to lose their diazo functionality in the presence of Lewis acids,<sup>7</sup> Calter described Lewis acid promoted electrophilic additions of carbonyl compounds to metal enolates of diazoacetoacetates,<sup>8a,b</sup> and Wang reported that these enolates also undergo Michael addition.<sup>8c</sup> Despite these advances, efficient syntheses of structurally complex diazo compounds under mild conditions remains a challenging problem.

We recently published an efficient Mukaiyama–aldol reaction between methyl 3-(trialkylsiloxy)-2-diazo-3-butenate **1**<sup>9</sup> and both aromatic and aliphatic aldehydes **2** (Scheme 1) catalyzed by scandium(III) triflate.<sup>10</sup> These

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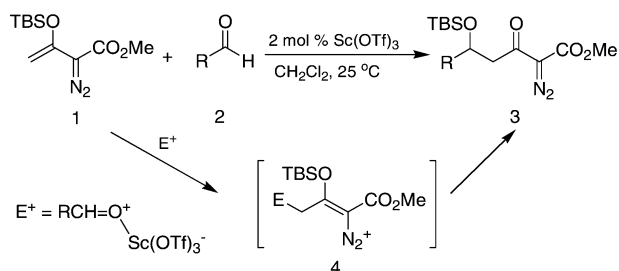
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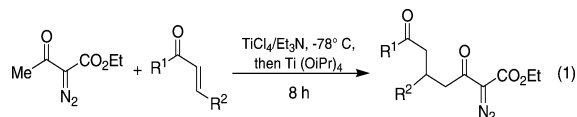
Scheme 1



reactions proceeded to form **3** in high yield without decomposition of the diazo group. Their success is due, at least in part, to the stability of the vinyl diazonium ion intermediate (**4**) that is formed by electrophilic addition to **1**. However, the low Lewis acidity of scandium(III) triflate required relatively long reaction times and limited application to aldehydes with strong electron-withdrawing groups.

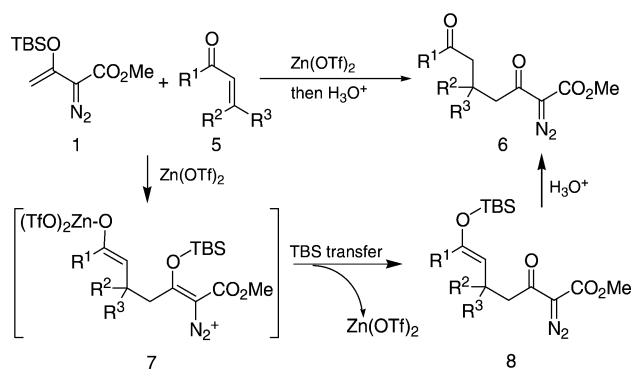
The Mukaiyama-Michael reaction, the condensation between enolsilanes and  $\alpha,\beta$ -unsaturated carbonyl compounds,<sup>11</sup> is a suitable extension of the Mukaiyama-aldol reaction. Reactions mediated by stoichiometric amounts of strong Lewis acid catalysts have been reported,<sup>12</sup> and the use of various lanthanides and other Lewis acids as catalysts has been extensively studied.<sup>13,14</sup> In the only report of a Mukaiyama-Michael reaction to form diazoacetates,<sup>8c</sup>  $\text{TiCl}_4$  is used in stoichiometric amounts to form the conden-

sation product (eq 1); reactions occurred at  $-78^\circ\text{C}$ , and 1,2-addition was competitive with 1,4-addition. In view of this,



a Mukaiyama-Michael reaction between **1** and  $\alpha,\beta$ -unsaturated carbonyl compounds using catalytic amounts of Lewis acid under milder conditions would be desirable. Herein, we report that highly functionalized diazo compounds (**6** and **8**) can be prepared by an efficient Lewis acid catalyzed Mukaiyama-Michael reaction between **1** and enones **5** (Scheme 2) under mild conditions with low catalyst loading

Scheme 2



(8) (a) Calter, M. A.; Sugathapala, P. M.; Zhu, C. *Tetrahedron Lett.* **1997**, 38, 3837–3840. (b) Calter, M. A.; Zhu, C. *J. Org. Chem.* **1999**, 64, 1415–1419. (c) Deng, G.; Tian, X.; Qu, Z.; Wang, J. *Angew. Chem., Int. Ed.* **2002**, 41, 2773–2776.

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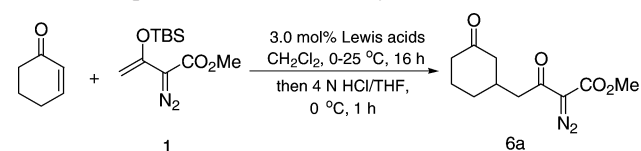
(14) For representative examples of asymmetric Mukaiyama-Michael reactions using Lewis acid catalysts, see: (a) Evans, D. A.; Rovis, T.; Kozlowski, M.; Dowey, C. W.; Tedrow, J. S. *J. Am. Chem. Soc.* **2000**, 122, 9134–9142. (b) Evans, D. A.; Scheidt, K. A.; Johnston, J. N.; Willis, M. C. *J. Am. Chem. Soc.* **2001**, 123, 4480–4491. (c) Desimoni, G.; Faita, G.; Filippone, S.; Mella, M.; Zampori, M.; Zema, M. *Tetrahedron* **2001**, 57, 10203–10212. (d) Sibi, M. P.; Chen, J. *Org. Lett.* **2002**, 4, 2933–2936. (e) Wang, X.; Adachi, S.; Iwai, H.; Takatsuki, H.; Fujita, K.; Kubo, M.; Oku, A.; Harada, T. *J. Org. Chem.* **2003**, 68, 10046–10057. (f) Suga, H.; Kitamura, T.; Kakehi, A.; Baba, T. *Chem. Commun.* **2004**, 1414–1415. (g) Harada, T.; Adachi, S.; Wang, X. *Org. Lett.* **2004**, 6, 4877–4879. (j) Ishihara, K.; Fushimi, M. *Org. Lett.* **2006**, 8, 1921–1924. (h) Takenaka, N.; Abell, J. P.; Yamamoto, H. *J. Am. Chem. Soc.* **2007**, 129, 742–743.

of inexpensive zinc triflate. As in previous examples of Mukaiyama-Michael reactions with enolsilanes, silyl group transfer across seven atoms was considered to be a major challenge.

In catalyst screening with cyclohexenone, we found that scandium(III) triflate, the previously preferred catalyst for Mukaiyama-aldol reactions of **1**,<sup>10</sup> was relatively ineffective for the Michael reaction with the same nucleophile. Thus, reaction of **1** with 1.1 equiv of cyclohexenone in the presence of 3.0 mol %  $\text{Sc}(\text{OTf})_3$  gave the Mukaiyama-Michael reaction product (**6a**) in only 43% isolated yield after subsequent hydrolysis (entry 1, Table 1). The low yield was determined to be caused by slow reaction as unreacted cyclohexenone and hydrolyzed silyl enol ether from **1** were also isolated. Lanthanum(III) triflate increased the yield slightly to 50%; however, the reaction was not significantly faster (entry 2). The use of boron trifluoride-etherate<sup>8b</sup> under the same conditions resulted in decomposition of **6a** (entry 3).

In related studies of the Cu(II)-catalyzed Mukaiyama-aldol reactions, silylation of the metal aldolate (silyl transfer) had been proposed to be the rate-limiting step.<sup>15a</sup> Furthermore, Ellis and Bosnich concluded that a Lewis acid having

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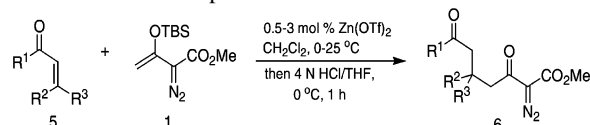
**Table 1.** Optimization of the Mukaiyama–Michael Reaction<sup>a</sup>

entry	Lewis acid	yield of <b>6a</b> <sup>b</sup> (%)
1	Sc(OTf) <sub>3</sub>	43
2	La(OTf) <sub>3</sub>	50
3	BF <sub>3</sub> ·OEt <sub>2</sub> (100 mol %)	0
4	Cu(OTf) <sub>2</sub>	29
5	AgOTf	44
6	Sn(OTf) <sub>2</sub>	48
7	Zn(OTf) <sub>2</sub>	79
8 <sup>c</sup>	Zn(OTf) <sub>2</sub>	96

<sup>a</sup> Vinyl diazoacetate **1** (0.5 mmol) was added to a solution of enone **5a** (0.55 mmol) and metal triflate (0.015 mol) in CH<sub>2</sub>Cl<sub>2</sub> (2 mL). The solution was stirred at 0 °C and then allowed to warm to room temperature. After 16 h, the crude reaction mixture was concentrated and hydrolyzed with 1:5 4 N HCl/THF at 0 °C. See the Supporting Information for details. <sup>b</sup> Yield of isolated **6a** following column chromatography. <sup>c</sup> 1.5 equiv of **1** was used under the same conditions.

a strong alkoxide–Lewis acid bond will inhibit silyl transfer.<sup>15b</sup> Thus, we focused our attention on the use of less oxophilic Lewis acids. A screening of selected mild Lewis acids (Table 1, entries 4–7) revealed that the inexpensive zinc(II) triflate gave a significantly improved yield (79%) of Mukaiyama–Michael product **6a** in the reaction between **1** and 1.1 equiv of cyclohexenone (entry 7). The yield of **6a** could be further improved to 96% with the use of 1.5 equiv **1** (entry 8).<sup>16,17</sup> Other Lewis acids were found to be much less effective than Zn(OTf)<sub>2</sub>; with Cu(II) and Ag(I) triflate diazo decomposition occurs, resulting in a complex reaction mixture and low yield (entries 4 and 5).<sup>1</sup> Having established optimum conditions with Zn(OTf)<sub>2</sub>, efforts were undertaken to reduce the required amount of catalyst: a 96% yield of **6a** was obtained with 0.5 mol % catalyst, and with only 0.1 mol % of Zn(OTf)<sub>2</sub> the yield of **6a** was 94%.

Using 3 mol % of Zn(OTf)<sub>2</sub> and the optimal conditions for production of **6**, reactions of **1** with various  $\alpha,\beta$ -unsaturated ketones were examined (Table 2). Cyclic enones **5a–d** and several acyclic enones **5h–j** all gave excellent yields of the desired product.<sup>18</sup>  $\beta,\beta$ -Disubstituted enones such as **5e** and **5g** are sterically hindered and were previously reported to react sluggishly<sup>19</sup> or give low yields of Mukaiyama–Michael product.<sup>14a</sup> However, although we found that its reaction with **1** was notably slower than with **5a**, the Michael adduct **6e** was still isolated in good yield with 3

**Table 2.** Zinc Triflate-Catalyzed Mukaiyama–Michael Reactions of **1** with Representative Enones<sup>a</sup>

entry	enone <b>5</b>	product <b>6</b>	yield <sup>b</sup> , %
1	<b>a</b>		96
2	<b>b</b>		84
3	<b>c</b>		94
4 <sup>c</sup>	<b>d</b>		99
5 <sup>d</sup>	<b>e</b>		68
6 <sup>e</sup>	<b>f</b>		74
7 <sup>d,f</sup>	<b>g</b>		71
8	<b>h</b>		88
9	<b>i</b>		92
10	<b>j</b>		99

<sup>a</sup> Vinyl diazoacetate **1** (0.75 mmol) was added to a solution of enone **5** (0.5 mmol) and zinc triflate (0.015 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (2 mL). The solution was stirred at 0 °C and then allowed to warm to room temperature. After 16 h, the crude reaction mixture was concentrated and then hydrolyzed with 1:5 4 N HCl/THF at 0 °C. See the Supporting Information for details.

<sup>b</sup> Yield of isolated **6** following column chromatography. <sup>c</sup> The product was isolated as two diastereomers in 3:1 ratio. <sup>d</sup> Reaction time was 36 h. <sup>e</sup> Two equivalents of **1** was used, 62% yield with 1.5 equiv of **1**; 71% yield with 0.5 mol % of catalyst. <sup>f</sup> Reaction was performed in 1 mL of CH<sub>2</sub>Cl<sub>2</sub>, and the concentration of enone **5g** was 0.5 M; 78% yield with 0.5 mol % of catalyst.

(16) The use of other common solvents such as chloroform, toluene, THF, and ether gave diminished yields of **6a**.

(17) In contrast to our report of Mukaiyama reaction (ref 10), the OTMS silyl enol ether analogue of **1** gave a lower yield of **6** than did **1**. The additional advantage of the OTBS silyl enol ether is its greater stability.

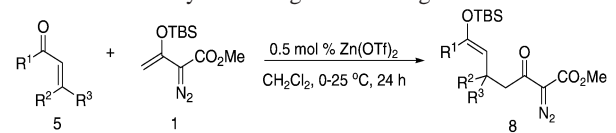
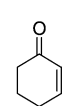
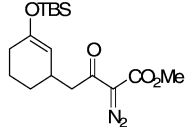
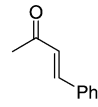
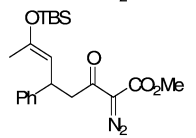
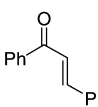
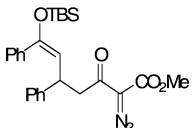
(18) Other common Michael acceptors such as  $\alpha,\beta$ -unsaturated esters, unsaturated nitriles, and unsaturated nitro compounds were found to be unreactive under the standard reaction condition.

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mol % catalyst after 36 h. Decreased reaction rate was also observed with **5g**, but a good yield of **6g** was obtained by increasing the concentration of reactants in the reaction solution. The reaction of methyl vinyl ketone **5f** was facile; however, oligomerization was found to be the major side-reaction. The yield of **5f** could be improved to 74% by using 2 equiv of **1**. Several of these reactions were performed in the presence of only 0.5 mol % zinc triflate catalyst and afforded comparable product yield (entries 1, 6, 7). In all cases, only 1,4-addition products were observed and isolated from the reaction mixture.

In performing these reactions, we anticipated that silyl enol ether **8** would be the major product prior to hydrolysis, and indeed it was. However, the product mixture after 16 h in a reaction apparatus that was open to the atmosphere contained variable amounts of **6** and **8** that were marginally reproducible. However, when the reaction apparatus was closed to the atmosphere, only **8** was obtained. Zinc triflate is sufficiently hygroscopic to draw water into the reaction solution and cause the hydrolysis of **8**. Reducing the amount of catalyst and performing the reaction in a closed apparatus resulted in good yields of **8**, representative examples for which are given in Table 3.

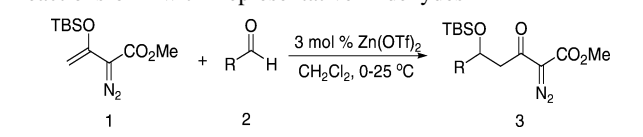
**Table 3.** Catalytic Mukaiyama–Michael Reactions of **1** with 0.5 mol % of Catalyst Loading and Leading to Product **8**<sup>a</sup>

			
entry	enone <b>5</b>	product <b>8</b>	% yield <sup>b</sup>
1	<b>a</b> 		78
2 <sup>c</sup>	<b>i</b> 		81
3	<b>j</b> 		83

<sup>a</sup> Vinyl diazoacetate **1** (0.5 mmol) was added to a solution of enone **5** (0.6 mmol) and metal triflate (0.0025 mol) in CH<sub>2</sub>Cl<sub>2</sub> (2 mL). The solution was stirred at 0 °C and then allowed to warm to room temperature and react for 24 h. <sup>b</sup> Yield of isolated **5** following column chromatography. <sup>c</sup> The product was isolated as two diastereomers in 3:1 ratio favoring the *E*-isomer.

Recognizing the effectiveness of zinc triflate for the Mukaiyama–Michael reaction, we returned to the Mukaiyama–aldol reaction to examine its effectiveness for that transformation (Table 4). The isolated yields of both **3a** and

**Table 4.** Zinc(II) Triflate Catalyzed Mukaiyama–Aldol Reactions of **1** with Representative Aldehydes<sup>a</sup>

				
entry	aldehyde	R	% yield of <b>3</b> (ref 10) <sup>b</sup>	% yield of <b>3</b> (this work) <sup>c</sup>
1 <sup>d</sup>	<b>2a</b>	C <sub>6</sub> H <sub>5</sub> -	93	97
2	<b>2b</b>	<i>trans</i> -β-styryl	89	91
3	<b>2c</b>	4-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	0	80
4	<b>2d</b>	2-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -	0	61

<sup>a</sup> Same procedure as in Table 2. <sup>b</sup> Yield of isolated **3** from Sc(OTf)<sub>3</sub> catalyzed aldol reaction; see ref 10 for details. <sup>c</sup> Yield of isolated **3** following column chromatography. <sup>d</sup> Use of 0.2 mol % of Zn(OTf)<sub>2</sub> gave a 97% yield of **3a**; use of 0.1 mol % Zn(OTf)<sub>2</sub> gave 92% yield of **3a**.

**3b** were slightly higher with zinc triflate than with scandium triflate, and Zn(OTf)<sub>2</sub> loading could be as low as 0.1 mol %. Cinnamaldehyde **2b** undergoes exclusive 1,2-addition with **1** in the presence of zinc(II) triflate; no 1,4-addition product was observed (entry 2). Furthermore, zinc(II) triflate was found to catalyze the reactions of 4-nitrobenzaldehyde **2c** and 2-nitrobenzaldehyde **2d**, which had previously been found to be unreactive using scandium triflate (entries 3, 4).

In conclusion, an efficient Mukaiyama–Michael methodology involving reactions between a diazo-containing silyl enol ether and various α,β-unsaturated ketones has been developed using inexpensive, commercially available, zinc triflate. Reaction conditions are mild, product yields are high under low catalyst loading, and the catalyst is also effective in the corresponding Mukaiyama–aldol reactions. This methodology can be used for the efficient construction of highly substituted diazo-containing compounds and allow late-stage installation and manipulation of the diazo functionality.

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**Supporting Information Available:** Experimental procedures and spectroscopic data for all new compounds. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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