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Selective One-Pot Synthesis of Allenyl and Alkynyl Esters from β -Ketoesters

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Abstract

A convenient method is described for the dehydration of β -ketoesters to generate conjugated and deconjugated alkynyl esters and conjugated allenyl esters. This sequential one-pot method involves the formation of a vinyl triflate monoanion intermediate that leads to the selective formation of alkynes or allenes depending on additives and conditions used. Product outcomes appear to be a function of unique mono- and dianion mechanisms which are described.

Introduction

Highly unsaturated carbonyl compounds such as allenyl¹ and alkynyl esters² have proven increasingly useful in organic synthesis. Conjugated allenyl carbonyls are particularly promising as synthetic intermediates due to their chiral axis and electronically distinctive cumulated double bonds that can be chemoselectively manipulated. Despite their synthetic promise,³ numerous challenges remain in the preparation and application of allenyl and alkynyl carbonyls. We have previously reported procedures involving amide bases to convert conjugated alkynyl esters to the corresponding allenyl esters with alkylation at the α -position.⁴ However, this method was limited to α -silylation, stannylation, and methylation with other alkylating agents leading to the deconjugated alkynyl ester product depending on reaction conditions.⁵ We have also described methods for the isomerization of α -alkynyl esters to their corresponding allenes involving manganese coordination chemistry.⁶ However, all of our previously disclosed methods to prepare β -alkynyl and α -allenyl esters require conjugated alkynyl esters starting materials which are not always readily available. As an alternative, we envisioned a one-pot transformation whereby a ketone carbonyl in substrates containing α -electron withdrawing groups can be dehydrated to generate doubly unsaturated products. Indeed, simple acyclic and cyclic ketones have been converted to their alkynyl and allenyl dehydration products via vinyl triflates⁷ and enol phosphonates.⁸ Hydrocarbon alkynes have also been prepared through a decarboxylative elimination of vinyl triflates.⁹ However, to our knowledge, no dehydration strategy has ever been reported for the preparation of allenyl esters. In this paper we describe our development of a dehydration approach starting from cheap and readily available β -keto esters to form three different allene-alkyne isomers depending on reaction conditions.

Results and Discussion

Our initial experiments involved the treatment of β -keto esters **1** with two equivalents LiHMDS at -78°C to form dienolates followed by trapping with TiF_2O .¹⁰ Subsequent warming to room temperature leads to conjugated alkynyl esters **2** in good yields for a variety of readily available β -keto ester substrates excepting those with heteroatom substitution at the γ -position (Table

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1). The choice of a metal HMDS base (i.e. lithium) was critical to the formation of **2** (and in transformations discussed below) whereas the use of LDA failed to give alkyne products. Generally 2-alkynoates are prepared through formation of bonds on either side of the triple bond either by the carboxylation of terminal alkynyl anions¹¹ or by the less favorable alkylation of propiolates that usually require activated electrophiles.¹² Both methods suffer drawbacks from either lack of availability of terminal alkynes or poor nucleophilicity of propiolate anions. Thus the present dehydration approach serves as an important alternative to existing 2-alkynoate synthesis methods.

It is interesting to note that these dehydration reactions did not produce any of the allene isomer. Even substrates containing α -substituents (entries 8 and 9), which were expected to eliminate the triflate *via* an E1_{cb} mechanism, failed to give allene products. Instead, vinyl triflates **4** and **5** were obtained after aqueous quenching. To further study this reaction, triflate intermediate **3a** was obtained by a low temperature aqueous quench of substrate **1a** (R = Me, R₁ = H) after the addition of base and triflic anhydride.¹³ Refluxing **3a** in THF in the presence of Et₃N gave near quantitative yield of alkynyl ester **2a** (Scheme 1). However, vinyl triflates **4** and **5** did not give allene products under these conditions even with extended reaction times. The treatment of **4** and **5** with a much stronger base (excess LiHMDS) did give allene products.

Our experiments with vinyl triflates **3a**, **4**, and **5** reveal a strong mechanistic preference for triflate elimination across the double bond even though the γ -hydrogens present in these substrates can also assume an anti-coplanar orientation. Similar observations have been made with β -bromo- α,β -unsaturated ketones.¹⁴ This strong regioselective preference for triflate elimination is helpful in deciding among mechanistic possibilities for the one-pot process leading to products **2**. The likely reactive intermediate in this reaction is triflate monoanion **6** (Figure 1). Following an E1_{cb} pathway, this intermediate would be expected to give allene product, which is not observed even with low temperature quenching. It is possible that the observed conjugated alkyne product is formed by an intramolecular abstraction of the α -proton *via* a 4-membered ring transition state.¹⁵ However, α -deuterio-analogs of intermediate **6** failed to yield product **7** with the anticipated γ -deuterium (Figure 1). Thus it appears that a more complex mechanism is operative here.

Conjugated alkynyl esters with substitution at the γ -position can also be prepared from acetoacetates (Scheme 2). In a one-pot procedure, the dianion of ethyl acetoacetate is treated with benzyl bromide followed by the addition of an additional equivalent of LiHMDS. The addition of Tf₂O at -78°C followed by warming to room temperature led to a reasonable yield of alkynoate **8**. Given the general availability of arylmethyl halides,¹⁶ this method will likely prove useful especially since the synthesis of **8** and related products by alkylation of 2-butyneates has not proved feasible.¹⁷

Our experience with the reactivity of substrates **4** and **5** prompted us to use an additional equivalent of strong base after the formation of intermediate vinyl triflate anions in an attempt to produce allenes in a one-pot procedure. Thus a series of β -keto esters were sequentially treated with LiHMDS (2 equiv), triflic anhydride (1 equiv), and a final equivalent of LiHMDS followed by quenching at low temperatures (Table 2). For substrates containing no α -substitution (entries 1–4), deconjugated alkynyl esters **9** were obtained in reasonable yields especially with the addition of HMPA prior to quenching. In a previous work we observed a similar preference for deconjugated products using HMPA.⁵ Presumably this agent acts to destabilize lithium complexes of the monoanion and HMDS, which have a tendency to quench the anion through intracomplex protonation.¹⁸ Deconjugated alkynoate **9a** (entry 1) was also produced in good yields despite its well-known propensity for isomerization to allene form **10a**.

It is important to note that this method (Table 2) did not lead to any conjugated alkynyl ester products. This product outcome can be rationalized in terms of the preference for the amide base to deprotonate intermediate **6** (formed after step 2) exclusively at the γ -position (Scheme 3). Overwhelming spectroscopic, crystallographic, and computational investigations establish that lithium cations generally interact with enolates at the oxygen center pointing to the predominance of oxyanion resonance forms.¹⁹ Therefore, as a first approximation, a comparison of the two oxyanion resonance forms resulting from selective deprotonation can be made to the exclusion of other resonance contributors. Deprotonation of monoanion **6** at the γ -position gives oxyanion **11** in which the alkene and alkyne bonds are in conjugation (alkynenolate). Whereas deprotonation at the α -position produces oxyanion **12** containing a cumulated system (cumulenolate) with diminished conjugation of the π -bonds. Although conjugated alkynyl ester products are more thermodynamically stable, their formation appears to require the intermediacy of the less favorable anion **12**. Indeed, in related ketone systems, a semiempirical computational study revealed that alkynenolates were over 5 kcal/mol more stable than their cumulenolate isomers.²⁰

Dehydration of α -substituted β -ketoesters tends to favor the formation of allenes **10** especially when ZnCl_2 is added prior to quenching (Table 2, entries 11–13). This was true whether the reactions were quenched at -78°C or at warmer temperatures. Although these allene products are thermodynamically favored, their formation can be better rationalized using a kinetic argument. Thus anionic intermediate **13** is expected to prefer a geometry in which the ester alkoxy group (OR^2) is *trans* to the larger α -substituent (Figure 2) based on our previous work involving trapping experiments with intermediates similar to **13**⁴ and on other reports.²¹ Protonation of anion **13** at the α -position leading to transition state **TS14** (Figure 2) would augment destabilizing steric interactions between the ester alkoxy and ethynyl groups as the α -carbon undergoes angle compression due to sp^3 hybridization. However, γ -protonation maintains sp^2 geometry at the α -carbon. This rationale appears to explain the product ratios observed for substrates where $\text{R} = \text{H}$ (entries 5 – 8). As the size of the α -substituent (R_1) increases, it is more likely that the ester alkoxy group will prefer the *trans* to this group in intermediate **13** thus increasing the propensity for γ -protonation. Indeed, a bulky *tert*-butyl ester group gave a significantly higher ratio of allene to alkyne relative to the ethyl ester (compare entries 6 and 8). It is unclear how zinc chloride acts to enhance allene formation in this reaction. In general, zinc enolates are thought to be present in solution as mainly the C-metalated species.²² Perhaps the transmetalation of intermediates such as **13** is governed by similar energetic considerations as just described for protonation.

Since large α -groups appear to favor allene products in our dehydration of β -keto esters, we sought to introduce an α -trialkyl silyl group as a control element. Thus two equivalents of LiHMDS were added to vinyl triflate intermediate **6** (Table 3) most likely generating an enolate dianion **15** ($\text{R} = \text{Et}$). We have previously observed that these unusually stable dianions can be readily prepared from conjugated alkynyl esters.⁴ Subsequent addition of TMSCl to **15** presumably gave bis-silyl intermediate **17**, which upon aqueous quench led to a 2:1 mixture of allenylsilane **18a** and propargyl product **9c** most likely the result of silyl group hydrolysis during aqueous workup (entry 1). Very recently, intermediates similar to **17** derived from allenyl esters have been converted to carbinol allenoates with excellent selectivity *via* Mukaiyama aldol-type reactions.³

The same reaction sequence with **15** ($\text{R} = \text{Et}$) using TIPSCl led to the exclusive formation of allenyl ester **18b**. This finding with the much larger TIPS group in the α -position parallels the trend of substrates with α -substitution from the monoanion reactions (Table 2 and Figure 2). The synthesis of product **18b** from **6** is also significant since no other substrate with γ -substitution discussed thus far has been converted exclusively to an allenyl ester product.

Deprotonation of **6** where (R = H) led to intermediate **16** which upon silylation gave deconjugated product **19**.

Conclusion

Conditions have been developed to selectively convert a series of β -ketoesters to conjugated allenyl, alkynyl, or β -deconjugated alkynyl esters. Following a neutral, monoanion, or dianion mechanistic rationale, our method involves a one-pot stepwise addition of LiHMDS, triflic anhydride, and quenching under varying conditions to bring about dehydration. Our investigation has also uncovered an efficient synthesis of ketene acetal intermediate **17** that preferentially adds proton at the γ -position to form allene product. We are currently exploring the reactivity of these types of intermediates with other electrophiles for the potential development of an asymmetric allene synthesis method.

Experimental Section

General procedure for synthesis of conjugate alkynes **2**

To a round bottom flask (50 mL) under argon, dry THF (7 mL) and LiHMDS (4 mL, 1.0 M in THF) were added. The solution was then cooled to $-78\text{ }^{\circ}\text{C}$ (acetone-dry ice bath) followed by the addition of β -keto ester (2 mmol). After stirring for 45 min, triflic anhydride (2 mmol) was added slowly over 15 min. The reaction was then stirred overnight slowly warming to room temperature and quenched with saturated NH_4Cl solution. The organic layer was separated, the aqueous fraction was extracted with ether, and the organic layers were combined and dried with anhydrous Na_2SO_4 . The product was purified by silica gel chromatography in the usual manner using 1–2% EtOAc in hexanes.

Ethyl hex-2-ynoate (**2d**)

^1H NMR (400 MHz, CDCl_3) δ 4.22 (q, $J = 7.14$ Hz, 2H), 2.31 (t, $J = 7.1$ Hz, 2H), 1.66–1.57 (m, 2H), 1.31 (t, $J = 7.14$ Hz, 3H), 1.02 (t, $J = 7.38$ Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) δ 153.8, 89.2, 73.2, 61.7, 21.0, 20.5, 14.0, 13.4; HRMS calc. for $\text{C}_8\text{H}_{12}\text{O}_2$ $[\text{M}+\text{H}]^+$: 141.0910. Found: 141.0899.

Procedure for synthesis of alkynyl ester **8**

To a round bottom flask (50 mL) under argon, dry THF (7 mL) and LiHMDS (4 mL, 1.0 M in THF) were added. The solution was then cooled to $-78\text{ }^{\circ}\text{C}$ (acetone-dry ice bath) followed by the addition of β -keto ester (2 mmol). After stirring for 45 min, BnBr (2 mmol) was added dropwise followed by warming slowly to room temperature over the course of 6 h. The reaction mixture was then cooled to $-78\text{ }^{\circ}\text{C}$ and charged with LiHMDS (2 mL, 1.0 M in THF). After for 45 min, TiF_2O (2 mmol) was added slowly over 15 min and the reaction was stirred overnight slowly warming to room temperature and quenched with saturated NH_4Cl solution. The organic layer was separated, the aqueous fraction was extracted with ether, and the organic layers were combined and dried with anhydrous Na_2SO_4 . The product was purified by silica gel chromatography in the usual manner using 1–2% EtOAc in hexanes.

Ethyl 5-phenyl-pent-2-ynoate (**8**)

^1H NMR (400 MHz, CDCl_3) δ 7.40–7.22 (m, 5H), 4.21 (q, $J = 7.2$ Hz, 2H), 2.89 (t, $J = 7.6$ Hz, 2H), 2.61 (t, $J = 7.6$ Hz, 2H), 1.30 (t, $J = 7.2$ Hz, 3H); ^{13}C NMR (100 MHz, CDCl_3) δ 154.0, 139.9, 128.8, 128.6, 126.9, 88.5, 73.9, 62.1, 34.1, 21.1, 14.3; HRMS calc. for $\text{C}_{13}\text{H}_{14}\text{O}_2$ $[\text{M}+\text{H}]^+$: 203.1067. Found: 203.1059.

General procedure for deconjugated alkyne/conjugated allene isomers via monoanionic enyne enolate with ZnCl₂ additive

To a round bottom flask (50 mL) under argon, dry THF (7 mL) and LiHMDS (4 mL, 1.0 M in THF) were added. The solution was then cooled to $-78\text{ }^{\circ}\text{C}$ (acetone-dry ice bath) followed by the addition of β -keto ester (2 mmol). After stirring for 45 min, triflic anhydride (2 mmol) was added slowly over 15 min. The reaction was maintained at $-78\text{ }^{\circ}\text{C}$ with stirring for an additional 1 h followed by the addition of LiHMDS (2 mmol). The mixture was stirred at $-78\text{ }^{\circ}\text{C}$ for an additional 30 min followed by the addition of HMPA (7 mmol). After stirring for stirring at $-78\text{ }^{\circ}\text{C}$ for an additional 30 min, ZnCl₂ (2.4 mmol) was added as an ether solution. The reaction was quenched after an additional 30 min by pouring into an ice-cold stirring biphasic mixture of ether and saturated aqueous NH₄Cl. NMR spectra were recorded after workup for each reaction confirming the formation of only conjugated allene which were compared with literature references (except for **10h**).

tert-Butyl-2-*n*-butylallenylester (**10h**)

¹H NMR (400 MHz, CDCl₃) δ 5.05 (t, $J = 3.03$ Hz, 2H), 2.18 (m, 2H), 1.47 (s, 9H), 1.45-1.30 (m, 4H), 0.91 (t, $J = 7.19$ Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) δ 213.6, 166.6, 101.7, 80.7, 30.2, 28.0, 27.7, 22.3, 13.9; HRMS calc. for C₁₂H₂₀O₂ [M+H]⁺: 197.1542. Found: 197.1543.

General procedures for silylated deconjugated alkyne/conjugated allene isomers via dianionic enolate

To a round bottom flask (50 mL) under argon, dry THF (7 mL) and LiHMDS (4 mL, 1.0 M in THF) were added. The solution was then cooled to $-95\text{ }^{\circ}\text{C}$ (methanol-liquid N₂ bath) followed by addition of β -keto ester (2 mmol). After stirring for 1 h, triflic anhydride (2 mmol) was added slowly over 15 min. Since Tf₂O is a solid at this reaction temperature, the slow addition should be accompanied by vigorous stirring. Alternatively, Tf₂O can be added as ether solution (not in THF since this solvent reacts to form polymer at room temperature). The reaction mixture was stirred for an additional 1 h at $-95\text{ }^{\circ}\text{C}$ followed by the addition of LiHMDS (4 mL, 1.0 M in THF). After an additional 1 h at $-95\text{ }^{\circ}\text{C}$, R₃SiCl (6 mmol) was added dropwise followed by warming to room temperature over 1 hour and quenching with saturated NH₄Cl solution. The organic layer was separated, the aqueous fraction was extracted with ether, and the organic layers were combined and dried with anhydrous Na₂SO₄. The product was purified by silica gel chromatography in the usual manner using 1–2% EtOAc in hexanes.

Ethyl-2-triisopropylsilylhexa-2,3-dienoate (**18b**)

¹H NMR (400 MHz, CDCl₃) δ 5.24 (t, $J = 6.61$ Hz, 1H), 4.16 (q, $J = 7.05$ Hz, 3H), 2.20-1.05 (m, 2H), 1.34-1.18 (m, 9H), 1.07 (d, $J = 7.40$ Hz, 18H); ¹³C NMR (100 MHz, CDCl₃) δ 216.2, 168.6, 91.5, 89.6, 60.7, 20.9, 18.6, 17.7, 14.2, 12.2; HRMS calc. for C₁₇H₃₂O₂Si [M+H]⁺: 297.2244. Found: 297.2236.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgment

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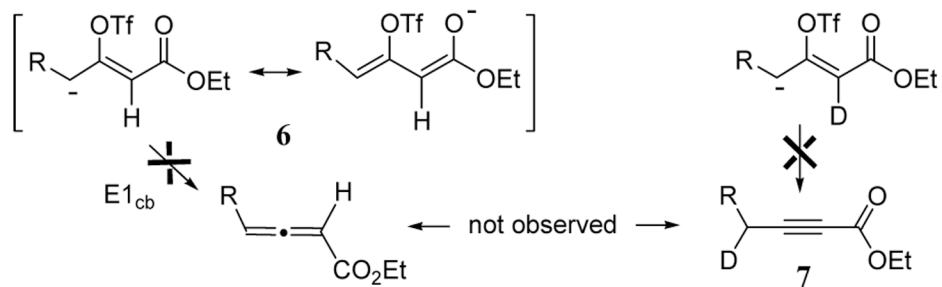


Figure 1.
Bimolecular proton transfer mechanism

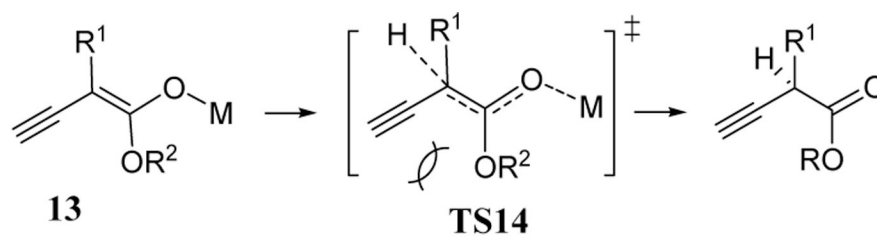
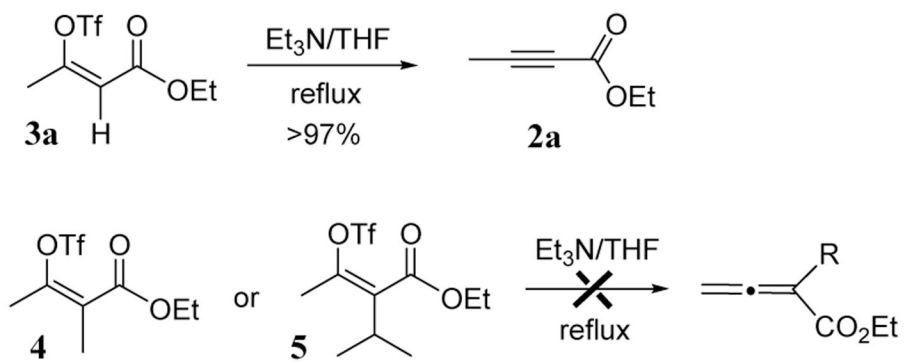
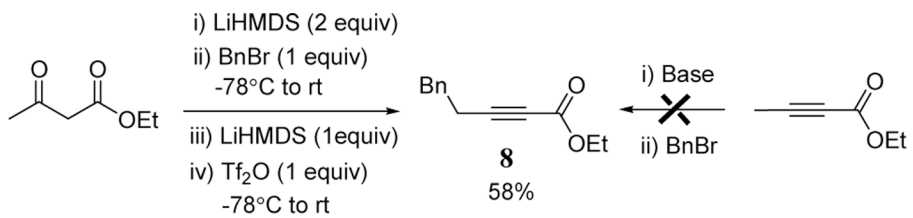
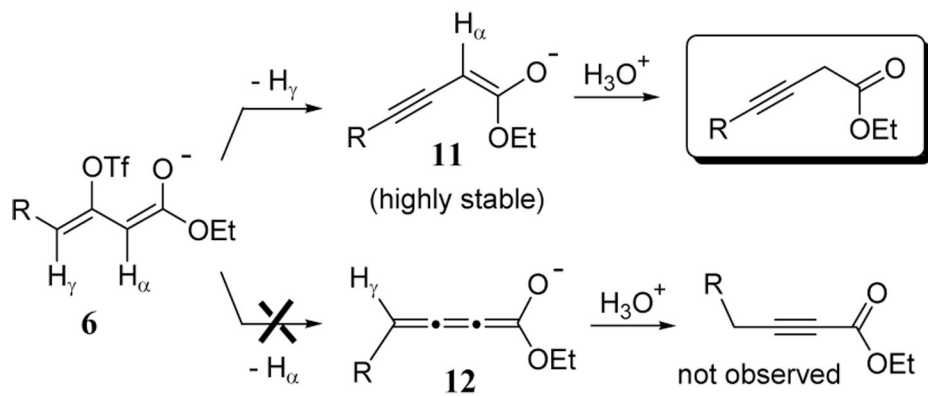


Figure 2.
Deconjugated alkynyl ester disfavored in substrates containing α -substitution

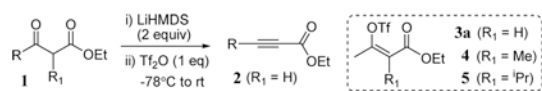


Scheme 1.
Preference for vinylic E2 elimination

**Scheme 2.**Synthesis of γ -substituted alkynyl ester from unsubstituted acetoacetate



Scheme 3.
Rationale for disfavored formation of 2-alkynyl ester

Table 1Synthesis of conjugated alkynyl esters from β -ketoesters

entry	R	R ₁	product	% yield ^a
1	Me	H	2a	84 ^b
2	Et	H	2b	90
3	ⁱ Pr	H	2c	72
4	ⁿ Pr	H	2d	82
5	ⁱ Bu	H	2e	77
6	ClCH ₂	H	2f	0
7	PhS	H	2g	0
8	Me	Me	4	87
9	Me	ⁱ Pr	5	83

^a Isolated yields.^b Quenching at -78°C gives **3a**.

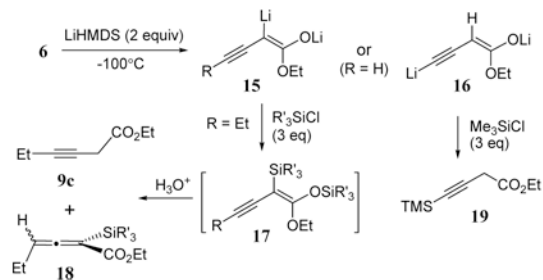
Table 2

Synthesis of 3-alkynoates **9** and allenyl esters **10**.

entry	R	R ¹	R ²	additive	Product (% yield) ^a
1	H	H	Et	None	9a (65)
2	Me	H	Et	None	9b (82)
3	Et	H	Et	None	9c (69)
4	ⁿ Pr	H	Et	None	9d (76)
5	H	Me	Et	None	9e (53)
6	H	ⁿ Bu	Et	None	9f (36)
7	H	ⁱ Pr	Et	None	9g (20)
8	H	ⁿ Bu	^t Bu	None	9h (20)
9	H	H	Et	ZnCl ₂	9a (0)
10	Et	H	Et	ZnCl ₂	9c (72)
11	H	Me	Et	ZnCl ₂	9e (0)
12	H	ⁿ Bu	Et	ZnCl ₂	9f (0)
13	H	ⁱ Pr	Et	ZnCl ₂	9g (0)
14	H	H	Ph	ZnCl ₂	9i (0)
15	H	H	^t Bu	ZnCl ₂	9j (0)
16	H	ⁿ Bu	^t Bu	ZnCl ₂	9h (0)

^a All yields are based on isolated products. For entries 5–8, isomers were isolated as inseparable mixtures and product ratios estimated by NMR.^b Care must be taken to use neutral silica gel during flash chromatographic isolation to avoid isomerization to allene.^c Numerous decomposition byproducts most likely due to ketene formation from phenoxide elimination.

Table 3
Synthesis of 3-alkynyl and allenyl esters via enyne dianions



entry	R	$\text{R}'_3\text{Si}$	Product (% yield) ^a
1	Et	TMS	9c (25) 18a (55)
2	Et	TIPS	- 18b (78)
3	H	TMS	19 (87) -

^a Isolated yields.