

## Synthesis of Oxazolidinone and Tosyl Enamines by Tertiary Amine Catalysis

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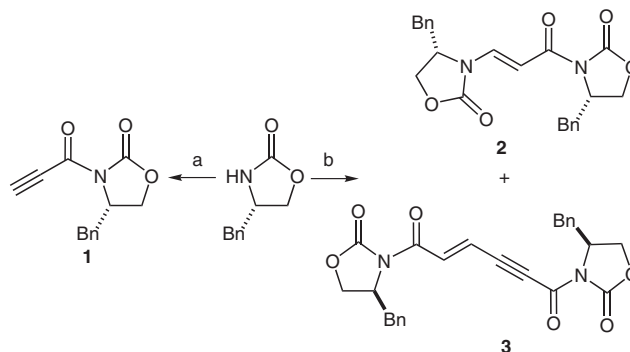
**Abstract:** A procedure for the synthesis of oxazolidinone and tosyl enamines is reported. Alkynoyl oxazolidinones and tosyl imides undergo reaction to form enamines in the presence of catalytic amounts of tertiary amines. The data suggest that an amide anion is formed during the reaction, which undergoes conjugate addition to form the final product.

**Key words:** enamine, oxazolidinone, tosyl imide, catalysis

Enamines have become widely used intermediates in organic synthesis since the development of facile methods for their use in the alkylation and acylation of carbonyl compounds by Stork.<sup>1</sup> These substrates are able to form carbon–carbon bonds easily, and the use of chiral amines provides an entry into asymmetric syntheses.<sup>2</sup> Over the past decade, there has been an expansion of the reactions that may undergo enamine organocatalysis, such as  $\alpha$ -oxidations<sup>3</sup> and alkylations,<sup>4</sup> aldol condensation,<sup>5</sup> Michael additions,<sup>6</sup> and enantioselective reductions.<sup>7</sup> These developments have been showcased in the recent syntheses of (–)-anisomycin<sup>8</sup> and (+)-conicol.<sup>9</sup> The present report describes a method for preparing enamine products from *N*-propynoyl oxazolidinones and tosyl imides through tertiary amine catalysis.

In the process of synthesizing *N*-propynoyl-(4*S*)-4-benzyl-1,3-oxazolidin-2-one (**1**) by Evans' conditions<sup>10</sup> to prepare analogues of locostatin, an *N*-crotonyl oxazolidinone that inhibits cell migration and disrupts specific protein–protein interactions involved in the regulation of cell signaling,<sup>11</sup> we unexpectedly discovered that the reaction underwent a secondary transformation to yield the bis-oxazolidinone enamine **2** (Scheme 1). The structure was confirmed by one- and two-dimensional NMR and ESI-MS (Table S1 in Supporting Information).<sup>12</sup> In addition, compound **2** was hydrolyzed under acidic conditions and found to react sluggishly, suggesting that the presence of the electron-withdrawing group adversely affects the ability of the enamine to undergo hydrolysis.

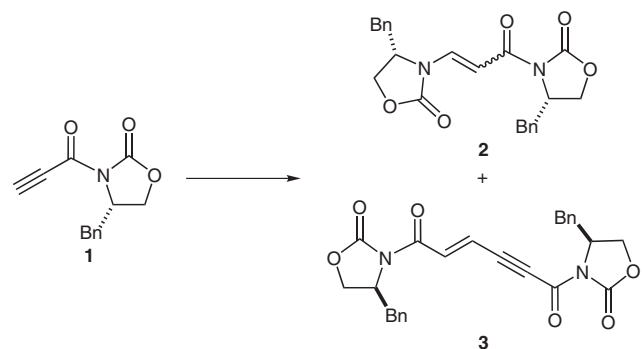
To explain how **2** arose from the reaction, we first explored generation of **1**. Replacing Et<sub>3</sub>N with K<sub>2</sub>CO<sub>3</sub> as the base allowed for generation of the propynoyl mixed anhydride, thus facilitating synthesis of **1** in 42% yield without formation of **2** (Scheme 1). Exposure of **1** to not only stoichiometric but also substoichiometric amounts of



**Scheme 1** Synthesis of *N*-propynoyl-(4*S*)-4-benzyl-1,3-oxazolidin-2-one (**1**), its enamine derivative **2** and its ene-yne dimer derivative **3**. *Reagents and conditions:* (a) 1. *n*-BuLi, THF, –78 °C; 2. propiolic acid, pivaloyl chloride, K<sub>2</sub>CO<sub>3</sub>, THF, r.t.; (b) 1. *n*-BuLi, THF, –78 °C; 2. propiolic acid, pivaloyl chloride, Et<sub>3</sub>N, –78 °C to 0 °C.

Et<sub>3</sub>N gave **2**, along with the ene-yne dimer **3** (Table 1, entries 1–7, and Table S2 in Supporting Information), implying a catalytic reaction. Alternative amines were used at 15 mol% catalyst loading in different solvents, and we found that DABCO also catalyzed the reaction at room temperature (Table 1, entries 8–14). No reaction was observed with DIPEA or imidazole (Table 1, entries 16 and 17). With (–)-sparteine (Table 1, entry 15), mild heating was required for the reaction to reach completion. Other solvents with 15 mol% Et<sub>3</sub>N and DABCO were screened (Table 1, entries 1–14). The catalyst–solvent system of Et<sub>3</sub>N in toluene gave the best results with an 86% yield of **2** (Table 1, entry 7). From these data, we concluded that the reaction requires a nucleophilic tertiary amine with diminished steric encumbrance, since imidazole and DIPEA failed to react and (–)-sparteine required mild heating.

Other systems (*N*-propynoyl oxazolidinones and tosyl imides) also underwent a similar reaction with catalytic Et<sub>3</sub>N (Table 2) in toluene. While the original oxazolidinone substrate **1** (Table 2, entry 1) furnished the enamine **2** with a yield of 86%, the other oxazolidinone substrates reacted to form enamine products with reduced yields of 26–38% (Table 2, entries 2–4). The tosyl imides, however, gave enamine products with 55–72% yields (Table 2, entries 7–10). Interestingly, the reaction appeared to necessitate the presence of an electron-withdrawing group, since tertiary and secondary alkyl ynamides failed to react under the same conditions (Table 2, entries 5 and 6). This result suggests that the oxazolidinone or tosyl moieties presumably act to facilitate the displacement of the nitrogen anion from the alkynoyl imide, as well as to decrease the hard-

**Table 1** Solvents and Amines Tested for Formation of **2** and **3**<sup>a</sup>

Entry	Amine	Solvent	Yield of <b>2</b> (%) and <i>E:Z</i> ratio <sup>b</sup>	Yield of <b>3</b> (%) <sup>b</sup>
1	Et <sub>3</sub> N	MeCN	36 (12:1)	trace <sup>c</sup>
2	Et <sub>3</sub> N	benzene	19 (99:1)	2
3	Et <sub>3</sub> N	CHCl <sub>3</sub>	20 (1:1)	25
4	Et <sub>3</sub> N	Et <sub>2</sub> O	49 (3:2)	trace <sup>c</sup>
5	Et <sub>3</sub> N	EtOAc	65 (12:1)	trace <sup>c</sup>
6	Et <sub>3</sub> N	THF	20 (99:1)	8
7	Et <sub>3</sub> N	toluene	86 (14:1)	trace <sup>c</sup>
8	DABCO	MeCN	37 (99:1)	31
9	DABCO	benzene	65 (99:1)	14
10	DABCO	CHCl <sub>3</sub>	56 (99:1)	5
11	DABCO	Et <sub>2</sub> O	43 (4:1)	5
12	DABCO	EtOAc	54 (8:1)	28
13	DABCO	THF	16 (99:1)	51
14	DABCO	toluene	60 (20:1)	25
15	(-)-sparteine <sup>d</sup>	THF	24 (99:1)	7
16	DIPEA	THF	– <sup>e</sup>	– <sup>e</sup>
17	imidazole	THF	– <sup>f</sup>	– <sup>f</sup>

<sup>a</sup> Reactions were performed at 0.22 M and 15 mol% amine at 24 °C.

<sup>b</sup> Isolated yield and *E:Z* ratio in parentheses.

<sup>c</sup> Trace signifies less than 1% product.

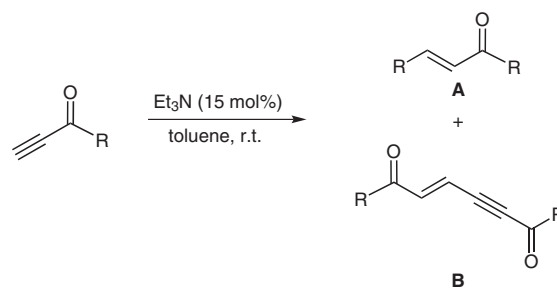
<sup>d</sup> Reaction was performed at 0.22 M, 15 mol% amine at 50 °C.

<sup>e</sup> No reaction, with 62% recovered starting material.

<sup>f</sup> No reaction, with 33% recovered starting material.

ness of the nitrogen anion to allow for conjugate addition onto a second molecule of the alkynoyl imide.

The mechanism of the reaction presumably begins with conjugate addition of the tertiary amine onto the ynamide to form an allenolate intermediate.<sup>13</sup> The allenolate undergoes elimination of the amide anion to generate the nucleophile for subsequent addition onto a separate ynamide moiety, forming the enamine. Alternatively, the allenolate mediates deprotonation of the acetylenic hydrogen of another equivalent of substrate to ultimately form the ene-

**Table 2** Enamine and Ene-Yne Dimer Products from Et<sub>3</sub>N-Catalyzed Reaction of *N*-Propynoyl Oxazolidinones and Tosyl Imides<sup>a</sup>

Entry	R	Yield of <b>A</b> (%) <sup>b</sup>	Yield of <b>B</b> (%) <sup>b</sup>
1		86	trace <sup>c</sup>
2		26	8
3		38	6
4		38	9
5		– <sup>c</sup>	– <sup>c</sup>
6		– <sup>c</sup>	– <sup>c</sup>
7		58	trace <sup>d</sup>
8		68	trace <sup>d</sup>
9		72	trace <sup>d</sup>
10		55	trace <sup>d</sup>

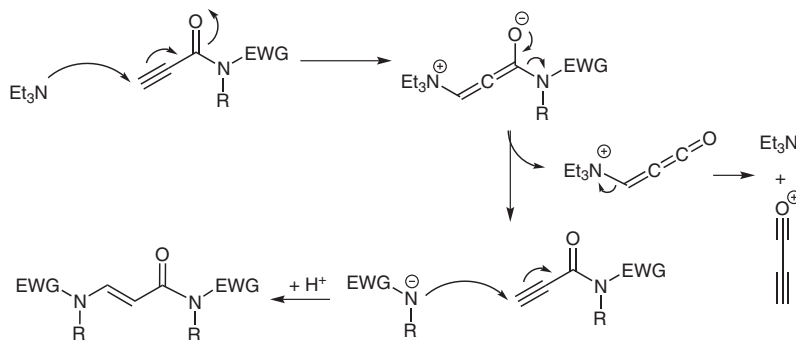
<sup>a</sup> Reactions were performed at 0.22 M and 15 mol% Et<sub>3</sub>N.

<sup>b</sup> Isolated yield.

<sup>c</sup> No observable reaction.

<sup>d</sup> Trace denotes <1% product.

yne dimer (Scheme 2).<sup>14</sup> Treatment of **1** with NaH and 20% aqueous DCl in D<sub>2</sub>O gave deuterated **1**, which was subjected to the catalytic conditions with DABCO and Et<sub>3</sub>N. We found that in both cases, the β position of the enamine product retained high levels of the deuterium la-



**Scheme 2** Proposed mechanism of enamine formation; EWG = electron-withdrawing group

bel (70% for DABCO and 58% for  $\text{Et}_3\text{N}$ ), while the  $\alpha$ -position contained lower levels (26% for DABCO and 39% for  $\text{Et}_3\text{N}$ ). The data suggest that it is more likely that deprotonation occurs through residual water than by direct deprotonation of the acetylenic hydrogen. Quenching the reaction mixture with TMSCl or 20% aqueous DCl in  $\text{D}_2\text{O}$  resulted in no incorporation of the trapping species in the final enamine product. Thus, the vinyl anion may not be neutralized during the aqueous workup, and the proton must be incorporated during the reaction.

We have found that *N*-propynoyl oxazolidinones and tosyl imides will form enamine products when exposed to catalytic amounts of  $\text{Et}_3\text{N}$  and DABCO.<sup>15</sup> The reaction involves conjugate addition of the tertiary amine onto the alkyne imide  $\beta$  carbon to furnish a putative allenolate intermediate. The allenolate then undergoes elimination of an amide (or carbamate) anion, generating the final nucleophile for product formation. This reaction could be used to furnish alternative enamines to yield asymmetric aldol products.

**Supporting Information** for this article is available online at <http://www.thieme-connect.com/ejournals/toc/synlett>.

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- (12) Compound **2** appeared as a single band from chromatographic techniques; additionally, compound **2** underwent palladium hydrogenation, and its spectra were in accordance with the reduced enamine (see Supporting Information).
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- (14) The freed allenone yielded a black tar, possibly the result of decomposition or polymerization.
- (15) For experimental procedures and compound characterization data, see Supporting Information.