

Enantioselective α -Vinylolation of Aldehydes via the Synergistic Combination of Copper and Amine Catalysis

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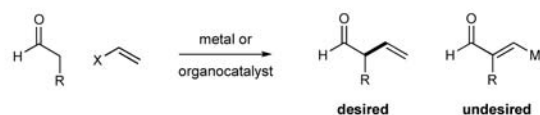
S Supporting Information

ABSTRACT: The enantioselective α -vinylolation of aldehydes using vinyl iodonium triflate salts has been accomplished via the synergistic combination of copper and chiral amine catalysis. These mild catalytic conditions provide a direct route for the enantioselective construction of enolizable α -formyl vinylic stereocenters without racemization or olefin transposition. These high-value coupling adducts are readily converted into a variety of useful olefin synthons.

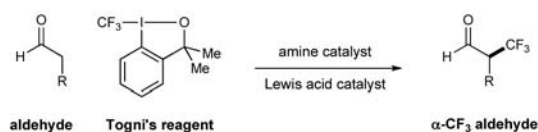
While monocatalysis strategies have successfully delivered vast numbers of new reactions over many decades, mult catalysis concepts that allow access to many elusive or unattainable transformations have only recently begun to evolve. In particular, synergistic catalysis, wherein two catalysts and two catalytic cycles work in concert to create a single new bond, has emerged as a powerful mechanistic approach to asymmetric reaction engineering.^{1–3} In this context, our laboratory has recently described the enantioselective α -trifluoromethylation and α -arylation of aldehydes via the synergistic combination of copper catalysis, organocatalysis, and iodonium salts (eq 1), two protocols that enable the production of structural motifs of high value to the pharmaceutical sciences.^{4–6} Continuing this theme, we questioned whether this mult catalysis strategy might be extended to the enantioselective α -vinylolation of carbonyls, an important yet complex synthetic task given the capacity of such adducts to readily undergo olefin–carbonyl conjugation or product racemization.⁷ Herein we describe the successful execution of these ideals via the combination of two discrete catalytic cycles (chiral amine–enamine, Cu(I)–Cu(III)) that enables the enantioselective construction of a wide variety of α -substituted- β,γ -unsaturated aldehydes without olefin isomerization or stereochemical ablation (eq 2).^{8–11}

As shown in Scheme 1, we postulated that a Cu(I) catalyst **1** (e.g., CuBr), in the presence of a vinyl iodonium triflate, will undergo chemoselective oxidative addition into the weak alkenyl–iodine bond to generate a highly electrophilic Cu(III)–vinyl complex **2**.^{12,13} In a simultaneous organocatalytic cycle, condensation of chiral amine catalyst **3** with an aldehyde should furnish the transient yet nucleophilic enamine **4**. Computational studies predict that the *Si*-face of this enamine will be shielded by the phenyl ring of the imidazolidinone framework, leaving the *Re*-face accessible to electrophilic attack (see DFT-S).¹⁴ On this basis, complexation of the enamine π -system with Cu(III)–vinyl complex **2** is expected to lead to η^1 -

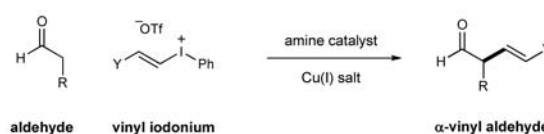
Challenging Transformation: Aldehyde α -Vinylolation w/o Conjugation



Enantioselective Trifluoromethylation: Mild and w/o Racemization (Eq 1)



Synergistic Catalysis: Enantioselective Aldehyde α -Vinylolation (Eq 2)



iminium organocopper species (not shown), which, upon reductive elimination, will forge a carbon–carbon bond and reconstitute the active Cu(I) catalyst (thereby completing the metal cycle). Hydrolysis of the resulting iminium **6** will also release amine **3** to complete the organocatalysis cycle while furnishing the desired enantioenriched α -vinyl aldehyde.

Our investigation began with an examination of the coupling of octanal with styrenyl iodonium triflate **7** in the presence of chiral amine catalyst **3** and various Cu(I) catalysts. As expected, no product formation was observed in the absence of copper salts (Table 1, entry 1). However, upon exposure to a catalytic amount of CuOAc, the desired α -vinyl aldehyde was produced with moderate yield and with excellent levels of asymmetric induction (entry 2, 67% yield, 97% ee). While the use of CuI resulted in diminished yields (entry 4, 23% yield), CuCl was found to be a competent coupling catalyst (entry 3, 82% yield, 97% ee). Optimal efficiency and enantiocontrol was achieved, however, using CuBr to afford the desired α -vinyl aldehyde adduct in 98% yield and 97% ee (entry 5). Importantly, lowering the copper and chiral amine catalyst loadings to 5 and 10 mol %, respectively, had little impact on the overall performance of this mult catalysis protocol (entry 6, 97% yield, 97% ee).

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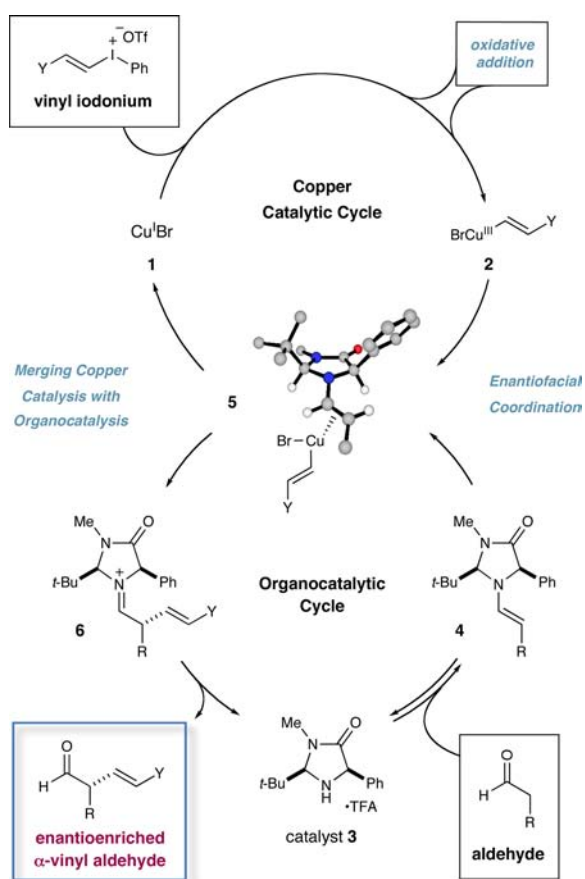
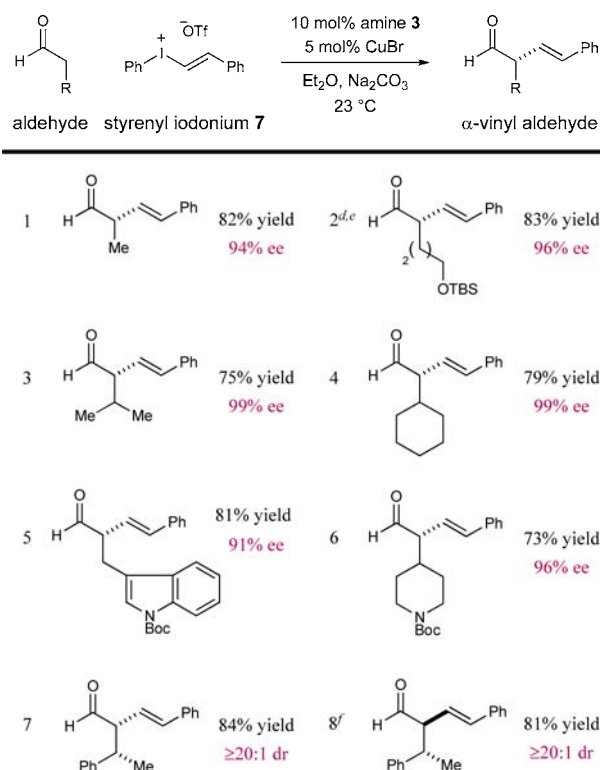
Scheme 1. Proposed Synergistic Cycles for Aldehyde α -Vinylolation

Table 1. Impact of Amine and Copper Catalysts on Vinylolation

entry	catalyst 3 (mol %)	copper catalyst	yield (%) ^a	ee (%) ^b
1	20	none	0	–
2	20	CuOAc (10 mol %)	67	97
3	20	CuCl (10 mol %)	82	97
4	20	CuI (10 mol %)	23	97
5	20	CuBr (10 mol %)	98	97
6	10	CuBr (5 mol %)	97	97

^a¹H NMR yield. ^bDetermined by chiral SFC analysis of the corresponding alcohol; absolute configuration determined by chemical correlation.

With optimal conditions in hand, we next examined the structural diversity of the aldehydic coupling partner (Table 2). Notably, sterically demanding β -branched aldehydes were found to be competent substrates, providing α -vinylated products with excellent levels of optical purity (entries 3, 4, and 6, 73–79% yield, 96–99% ee). Moreover, the presence of pendent heteroatom functionalities does not have a deleterious impact on the reaction performance (entries 2, 5, and 6). It should be noted that the two cycles involved are sufficiently mild to maintain configurational integrity of the pendent Z-

Table 2. Scope of Aldehyde in Enantioselective Vinylolation^{a,b,c}

^aAbsolute configuration assigned by chemical correlation or analogy. ^bIsolated yield of the corresponding alcohol. ^cEnantiomeric excess determined by chiral HPLC analysis of the corresponding alcohol. ^dUsing 20 mol % 3. ^eUsing 20 mol % CuBr. ^fUsing (S,S)-3 catalyst.

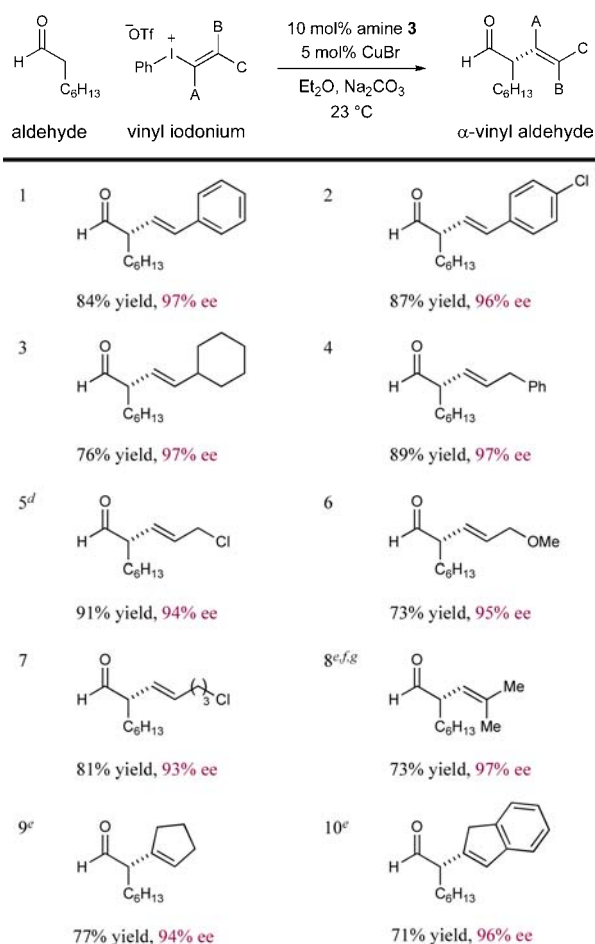
alkene (entry 1, 69% yield, 96% ee). More importantly, aldehydes bearing β -stereogenicity can be routinely employed to enable the highly diastereoselective construction of both *syn*- and *anti*- α,β -disubstituted adducts (entries 7 and 8, 81–84% yield, $\geq 20:1$ dr). As a salient indicator of the dominance of catalyst versus substrate control in these processes, both β -stereocenter-containing antipodes undergo α -olefination with the same sense of induction (as dictated by the amine catalyst).

Next we turned our attention to the scope of the vinyl coupling partner. As demonstrated in Table 3, these merged catalysis conditions are amenable to a broad range of iodonium triflates. Both electron-neutral and electron-poor styrenes (entries 1 and 2) as well as alkylvinyl groups (entries 3, 4, and 7) are transferred with high levels of enantiocontrol ($\geq 93\%$ ee).¹⁵

Traditionally metal-sensitive structural motifs such as allyl chlorides and allyl ethers (entries 5 and 6, 94–95% ee) are also readily incorporated to deliver useful synthons for subsequent elaboration. Moreover, this novel transformation is not limited to 1,2-disubstituted alkenyl group couplings but can also be applied to trisubstituted carbocycles (entries 9 and 10, 71–77% yield, 94–96% ee).¹⁶ The selective transfer of sterically demanding trisubstituted olefins can also be accomplished; however, this system requires the use of the corresponding mesityliodonium salt to avoid a competing aryl group transfer (entry 8).¹⁷

To demonstrate the utility of this new enantioselective vinylolation reaction, we performed a series of subsequent synthetic manipulations to demonstrate that this protocol can

Table 3. Scope of the Vinyl Iodonium Coupling Component^{a,b,c}



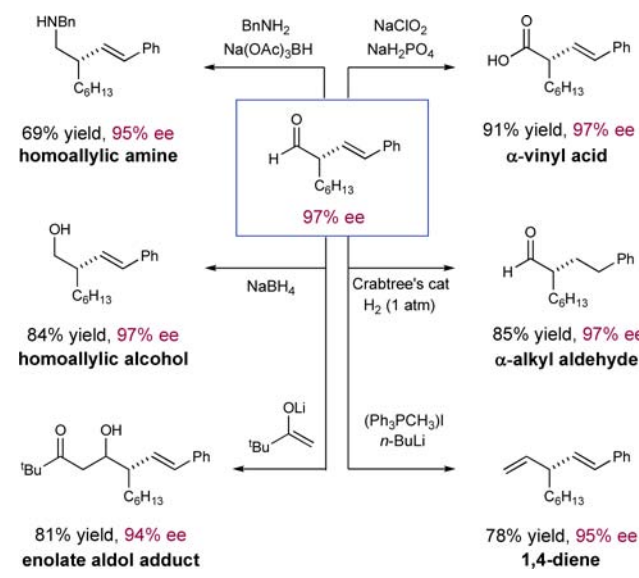
^aAbsolute configuration assigned by chemical correlation or analogy.

^bIsolated yield of the corresponding alcohol. ^cEnantiomeric excess determined by chiral HPLC or SFC analysis of the corresponding alcohol. ^d¹H NMR yield. ^eUsing 20 mol % 3. ^fUsing 30 mol % CuBr. ^gUsing the corresponding mesitylvinylidonium triflate.

lead to a diverse series of synthons (Scheme 2). For example, Pinnick oxidation of the formyl group provides an α -olefin carboxylic acid in high yield with complete preservation of enantiomeric purity. Moreover, hydrogenation of the alkene group using Crabtree's catalyst leads to α -alkyl aldehyde products, again without loss in optical purity. Importantly, the formyl group can be utilized in a reductive amination sequence to afford a homoallylic amine scaffold in 95% ee, a remarkable outcome given that an iminium intermediate is formed in this process. As a related system, homoallylic alcohols can also be readily accessed via NaBH₄ reduction, to generate adducts that should be of value for subsequent diastereoselective reactions such as epoxidation, cyclopropanation etc.¹⁸ The addition of metal enolates to these vinylation adducts can also be accomplished to generate aldol products with high efficiency. Last, the use of traditional Wittig conditions leads to 1,4-diene formation without loss of enantiopurity at the stereogenic 3-position, again demonstrating the versatility of these 1,3-formyl-olefin adducts.

In conclusion, we have developed a highly enantioselective protocol for the α -vinylation of aldehydes with vinyl iodonium triflates using copper(I) and chiral amine catalysis in a

Scheme 2. Access to Diverse Enantioenriched Vinyl Synthons



synergistic coupling mechanism. This operationally simple, room temperature and mild protocol should also be cost-effective given the substrates, catalysts and catalyst loadings employed. We fully expect that the α -formyl olefin products generated in this study will be of high value for practitioners of chemical synthesis and medicinal chemistry.

■ ASSOCIATED CONTENT

Supporting Information

Experimental procedures and spectral data. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

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NOTE ADDED AFTER ASAP PUBLICATION

The Supporting Information published online on May 29, 2012, showed incorrect spectra for Table 2, Entry 3. The corrected version was reposted on June 6, 2012.