Regioselective Asymmetric Alkynylation of N–Alkyl Pyridiniums

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Supporting Information Placeholder

ABSTRACT: Disclosed in this communication is a novel asymmetric addition of alkynyl nucleophiles to N-alkyl pyridinium electrophiles. The coupling is effected under mild and simple reaction conditions, affording dihydropyridine products with complete regiochemical and stereochemical control. In addition to several manipulations of the dihydropyridine products, the utility of this transformation is demonstrated through a concise, dearrative, and asymmetric synthesis of (+)-lupinine, a natural acetylcholine esterase inhibitor.

Alkaloid natural products have a rich history in medicine, thus fortuitously inspiring innovation and invention by synthetic chemists for decades. In turn, the piperidine heterocycle, a common moiety in bioactive alkaloids and rationally designed drugs, has demanded creative new strategies for its construction for many years. Moreover, the asymmetric construction of substituted congeners of this important heterocycle has been of the utmost importance in the synthetic pursuit of complex and medicinally relevant molecular targets.

Notable biologically active targets that contain multiply-substituted piperidines are depicted in Figure 1A. For example, the bisquinolizidine alkaloids related to (-)-neothiobinupharidine (2) have shown unique inhibition of the NF-kB signaling pathway, and tangutorine (3) has displayed noteworthy anticancer activity. Recently, the akuammiline alkaloid 2(S)-cathafoline (4) has shown remarkable inhibition of Ca²⁺-ion channels while (+)-lupinine (1) has been shown to inhibit acetylcholine esterase. Beyond their impressive biological profiles, each of these synthetic targets contains at least one of its stereogenic centers adjacent to the nitrogen atom of the piperidine heterocycle (See starred atoms, Figure 1A). While considering this salient structural feature, we aimed to develop a unified approach that would be general for building various piperidines thus allowing concise and asymmetric access to these privileged bioactive substances.

Recently, our research group became interested in the synthesis of biologically active alkaloids through the dearomatization of heterocyclic aromatic starting materials. For example, (+)-lupinine (1) was envisioned to arise from dihydropyridine 5, which would be derived from the dearomatic coupling of alkyne 7 and pyridinium 6 (Figure 1). One example of this type of asymmetric dearomative alkynylation was demonstrated by Charette in 2001 exploiting a chiral iminoyl group to activate the pyridine 8 electrophile. Regioselective addition of the alkynyl Grignard reagent, as directed by the activating group, provided dihydropyridine 9. While innovative, superstoichiometric amounts of the pyridine heterocycle (8) had to be employed in this protocol, limiting its broader utility.

Years later, Ma and coworkers demonstrated a similar transformation using a chiral Cu catalyst and acylated pyridiniums for enantioselective dearomative alkynylation (see 13). This transformation was also remarkable, but only pyridine (8) was...
used as a substrate, and only ynone and ynoate nucleophiles provided high enantioselctivities. Shortly after this, another single example of an enantioselective alkylation of an acylated pyridine was also demonstrated by Arndtsen.\textsuperscript{26} Furthermore, while much work has been done activating pyridines and other aromatic azacycles\textsuperscript{27,28} via acylation, including these aforementioned examples, selective additions to \textit{N}-alkyl pyridiniums have lagged behind,\textsuperscript{29-32} despite their potential to provide more robust and strategic access to piperidine heterocycles with higher regio- and stereocontrol (\textit{vide infra}).

Prior work in our group established a working model for the innate regioselectivity for the attack of Grignard nucleophiles on substituted pyridiniums.\textsuperscript{15} In one scenario (See Figure 1C), an amide substituent provided for a regioselective nucleophilic addition at C2 that resulted in the conversion of pyridinium 11 to dihydropyridine 13. Seeing that it was likely amide chelation of the organometallic that drove this regioselective preference (see intermediate 12), it was thought that substituting the amide with a simple chiral guide (X\textsuperscript{*}, see 15) would both control the regiochemistry and stereochemistry of the dearomatative addition (see 16). Regioselective preference was likely to be enhanced by using alkynylmagnesium nucleophiles (e.g. 14), which show preferential pyridinium addition at C2 and C6. It should be noted that, while alkynyl nucleophiles have been added into pyridiniums previously, there are no examples of highly regioselective and stereoselective additions to substituted pyridiniums. Overall, overcoming these obstacles in selectivity would add value to employing dearomatative strategies and tactics toward the asymmetric synthesis of polysubstituted piperidine motifs.

Initially, exploration of the desired alkylation reaction was inspired by Koga’s work on addition to unsaturated aldehydes from several decades ago (See Table 1).\textsuperscript{[33]} Following simple condensation between \textit{L-}\textit{t}-leucine \textit{t}-butyl ester and 3-formylpyridine to form 17, chemoselective methylation and the temperature (\textasciitilde\textdegree C or rt) had little effect on the selectivity, but resulted in slightly reduced yields. Changing the solvent from THF to either CH\textsubscript{3}Cl\textsubscript{2} or Et\textsubscript{2}O also resulted in only a minor effect on the yield. In the interest of using a removable nitrogen substituent, activation of the pyridine 17 with either benzyl bromide or allyl bromide (See Supporting Information for experimental details) followed by propynylmagnesium bromide resulted in comparable yields.

With optimized conditions in hand, we embarked on exploring the asymmetric alkylation of various substituted pyridine starting materials (Scheme 1). Following alkylation with either allyl bromide or methyl triflate, pyridinium intermediates with substitution at the 4, 5, and 6 positions were successfully converted to the corresponding alkylated dihydropyridines. Furthermore, chloride (21), bromide (22), aryl (23), and alkyl (20, 24) substitution were all tolerated under the reaction conditions with adducts isolated in 37–70\% yields. A

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**Table 1.** Reaction optimization studies. \textsuperscript{[a]}Determined by \textsuperscript{1}H NMR analysis of the crude mixture with an internal standard. \textsuperscript{[b]}Isolated yield.

<table>
<thead>
<tr>
<th>Deviation from conditions above</th>
<th>Yield of 19</th>
<th>dr of 19</th>
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<tbody>
<tr>
<td>2 equiv of 18</td>
<td>43%\textsuperscript{[a]}</td>
<td>&gt; 20:1</td>
</tr>
<tr>
<td>–40 °C instead of –78 °C</td>
<td>64%\textsuperscript{[a]}</td>
<td>&gt; 20:1</td>
</tr>
<tr>
<td>rt instead of –78 °C</td>
<td>67%\textsuperscript{[a]}</td>
<td>&gt; 20:1</td>
</tr>
<tr>
<td>Et\textsubscript{2}O instead of THF</td>
<td>63%\textsuperscript{[a]}</td>
<td>&gt; 20:1</td>
</tr>
<tr>
<td>CH\textsubscript{3}Cl\textsubscript{2} instead of THF</td>
<td>62%\textsuperscript{[a]}</td>
<td>&gt; 20:1</td>
</tr>
<tr>
<td>BnBr\textsuperscript{[b]} instead of MeOTf</td>
<td>70%</td>
<td>&gt; 20:1</td>
</tr>
<tr>
<td>AllylBr\textsuperscript{[b]} instead of MeOTf</td>
<td>69%</td>
<td>&gt; 20:1</td>
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\textsuperscript{[a]}See Supporting Information for conditions.
dihydroquinoline product (25) was also afforded following activation via methylation with MeOTf and treatment with propynylmagnesium bromide. In addition, dihydropyridine 26, derived from 3-acetylpyridine was also produced with this method demonstrating that aldimines are not the only competent functionality for directing the desired addition. It should be noted that, despite the substitution pattern for any of these substrates, only one stereoisomer and one regioisomer was observed in each case.

Next, the activating group was further evaluated to understand its effect on the alkylation reaction. Having established that methyl (19), benzyl (28,29), and allyl (31) were all competent activating groups in the initial reaction screening, it was also determined that p-methoxybenzyl as well as methoxethyl were also competent N-activating groups. The alkylation of these pyridinium intermediates provided 27 in 65% yield and 30 in 70% yield, each with full regioselectivity and stereoselectivity. Activation of the pyridine nitrogen of 17 via acylation was also briefly explored, but upon addition of either organocopper or organomagnesium acetylides, complex mixtures of regioisomers, diastereomers, and rotamers were observed, in agreement with previous work on alkylation of substituted systems.35–38

Following the investigation of variously substituted pyridines, the scope of alkyne addition was evaluated using pyridine 17 as a model substrate (Scheme 2). Upon treatment of 17 with allyl bromide, the resultant pyridinium was then treated with a variety of alkynyl Grignard reagents to afford variously substituted dihydropyridine adducts in 30 minutes at −78 °C. Alkynyl nucleophiles bearing alkyl (31,37), aryl (33), silyl (34), and alkoxy (35) substituents were all competent in this reaction giving synthetically useful yields with complete regio- and stereochemical control. In addition, ethynylmagnesium bromide was used as a nucleophile to selectively give adduct 32 in 75% yield. Alkynes bearing propargylic ethers (38 and 39) and a homopropargylic chloride (36) also gave useful functionalyzed products under this reactivity manifold, allowing for further synthetic manipulation following the dearomatization (vide infra). Aliphatic and aromatic heterocyclic alkynes also performed well yielding dihydropyridine adducts containing indole (40), pyridine (43), quinoline (44), and pyrroldine (45) rings. It is worth noting that no epimerization was observed in adduct 45, as only one diastereomer was observed in the crude reaction mixture. Adducts with important isosteres39 like a cyclopropane and chemical reporter groups like a diyne40 also were incorporated into products 41 and 42 in 49% and 48% yield, respectively.

A stereochemical model for this asymmetric alkylation reaction is depicted in Scheme 3A. Following activation of a pyridine 17 with allyl bromide to form an intermediate pyridinium bromide, the alkynyl Grignard 46 was introduced at −78 °C. According to reactivity previously described by Koga for the addition to unsaturated imines, chelation of the magnesium center by the imino ester guides the alkynyl nucleophile to the C2 position (See intermediate 47).33 The tert-butyl group passively blocks the top face of the pyridinium allowing for the alkynyl nucleophile to approach from the opposite side, providing for high stereocontrol. In addition to the exquisite stereoccontrol that is observed in this transformation, the absolute regioselectivity can be attributed both to the chelation by the imino ester as well as the preference for alkynes to add at the “harder” 2 position versus the 4 position of the pyridinium.29 Further evidence for this regioselective and stereoselective chemical model was confirmed by X-ray analysis of 34 which shows addition of the alkynyl nucleophile opposite of the bulky tert-buty1 side chain to give the expected C2 stereochemical outcome. At this time, however, solvent effects on the transition state of the addition cannot be completely ruled out, as this has been crucial in the success of other asymmetric dearomatizations.41

The dihydropyridines forged from these asymmetric dearomatizations can be manipulated in various ways, as shown in Scheme 3B. Following cleavage of the chiral guide under treatment with basic alumina (See Supporting Information), aldehyde 48 (R = H or Me) can be converted to various products. Horner–Wadsworth–Emmons olefination at elevated temperature an provided ester 49, while activation of 48 with TIPSOTf in MeCN followed by TBAF treatment provided Mannich product 50 in 50% yield.42 A Cu-catalyzed alkynyl-azole “click” reaction with benzyl azide provided 51 in modest
yield. It is anticipated that each of these modifications both to the periphery and the core of the alkynyl dihydropyridines will be instrumental in future dearomative synthetic endeavors.

Lastly, as the synthesis of alkaloid natural products was the original inspiration and impetus for this work, a concise synthesis of (+)-lupinine was accomplished (Scheme 3C). Starting from pyridine 17, activation with allyl bromide and selective alkynylation provided a mixture of aldehyde 53 and the corresponding imine in a 1:2 ratio and 86% overall yield. This ratio was the result of prolonged exposure of the product mixture to basic alumina during the purification (See Supporting Information for details). Conveniently, imine 36 could by hydrolyzed to aldehyde 53 by further exposure to basic alumina. Then, reduction of dihydropyridine 53 under Luche conditions provided allylic alcohol 54 in 50% yield. This intermediate was then deallylated under Pd catalysis, subjected to hydrogenation with Raney Ni, and then cyclized under basic conditions in one pot to provide (+)-lupinine (1) in a 75% yield and a >10:1 diastereomeric ratio with its epimer epilupinine.

In conclusion, a new asymmetric platform for the regioselective and stereoselective dearomatization of substituted pyridines has been achieved. In addition to having a broad scope and functional group tolerance, this transformation is useful for synthesizing a variety of dihydropyridines that can be broadly useful for the asymmetric synthesis of various azaheterocyclic building blocks. Furthermore, the implementation of this method in total synthesis was demonstrated through the concise asymmetric construction of the bicyclic alkaloid (+)-lupinine (1). The success of this synthesis served as a test for the implementation of this stereoselective and redox-economic dearomative approach to the synthesis of bioactive alkaloids. It is anticipated that further synthetic applications expanding upon this dearomative retrosynthetic analysis will be reported by our research group in due course.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website.

Supporting Information (PDF)

Crystallographic Information (CIF)

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Notes

The authors declare no competing financial interest.

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REFERENCES


