L-\textit{tert}-Leucine-Derived AmidPhos–Silver(I) Chiral Complexes for the Asymmetric [3+2] Cycloaddition of Azomethine Ylides

Zhipeng Zhou
Xiaojun Zheng
Jialin Liu
Jinlei Li
Pushan Wen
Haifei Wang*

College of Life Science and Chemistry, Hunan University of Technology, Zhuzhou 412007, Hunan Province, P. R. of China
whf2107@hotmail.com

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Abstract The L-\textit{tert}-leucine-derived AmidPhos/silver(I) catalytic system has been developed for the asymmetric [3+2] cycloaddition of azomethine ylides with electronic-deficient alkenes with or without Et\textsubscript{3}N. Under optimal conditions, highly functionalized \textit{endo}-4-pyrroli- dendines were obtained with modest to high yields (up to 99% yield) and enantioselectivities (up to 98% ee).

Key words L-\textit{tert}-leucine, amidophosphane, silver(I), [3+2] cycloaddition, azomethine ylide

Five-membered nitrogenous heterocycles, in particular, the highly substituted pyrrolidines are useful building blocks for biologically active molecules,\textsuperscript{1} as the structural motifs are widely present in many natural alkaloids and pharmaceutically useful agents. In recent years, synthesis of this five-membered heterocyclic compounds have been the focus of attention.\textsuperscript{2} The 1,3-dipolar cycloaddition reaction of azomethine ylides to electron-deficient alkenes is one of the most useful tool for constructing highly substituted pyrrolidines. Since the first catalytic asymmetric 1,3-dipolar cycloaddition reported by Zhang employing the AgOAc/xyllyl-FAP/i-Pr\textsubscript{2}NEt system,\textsuperscript{3} several examples of the formation of optically pure pyrrolidines based on a combination of chiral metal Lewis acids and organic or inorganic bases have thus far been reported to catalyze the process with high \textit{endo}/\textit{exo} diastereo- and enantioselectivities.\textsuperscript{4-6} Despite these impressive advances, there are still some problems that need to be explored for the reaction. First, the effect of the extra bases on the substrates adaptability has hardly been systematically studied.\textsuperscript{7} Second, synthesis of pyrrolidine derivatives containing aliphatic, heterocyclic substituents and 2-quaternary stereocenter with high enantioselectivities with small amounts of catalysts loading are still limited.\textsuperscript{8}

In previous papers, the most accepted mechanism for the 1,3-dipolar cycloaddition reaction of azomethine ylides to electron-deficient alkenes has been proposed.\textsuperscript{9} Coordination of the iminoester to a chiral metal catalyst, followed by deprotonation with base to form the reactive metal-bound azomethine ylide dipole, which reacts with dipolarophiles, was followed by elimination of cycloadduct to regenerate the chiral catalyst (Scheme 1). Thus, for the catalytic system, an excess amount of base such as a tertiary amine or an inorganic base was involved. However, a few researchers reported that extra bases are not necessary for their catalytic systems, because the metal salt bearing a moderately charged with a basic ligand anion would facilitate deprotonation of the iminoester to generate the azomethine ylide.\textsuperscript{4c,9b,10} Whether such a catalytic system require an extra base or not, we believe that the deprotonation of the iminoester can be accelerated by an suitable base, which is advantageous to improve the reaction rate and enantioselectivity.
tivities, especially for those slower reaction substrates containing aliphatic, heterocyclic, and α-substituted iminoesters. Here, we examined the substrate adaptability of 1,3-dipolar cycloaddition reaction of azomethine ylides to dipolarophiles catalyzed by the chiral l-tert-leucine-derived AmidPhos/Ag2CO3 catalytic system by small amounts of catalyst loading with or without base.

Recently, our group reported a new Ag2CO3/CA-AA-AmidPhos catalytic system which was applied to asymmetric 1,3-dipolar cycloaddition of azomethine ylides.1,2 Through the studies of the reactivity of precatalyst in the cycloaddition, we found the Ag2CO3/L-valine-derived amidoephosphine 1a system can efficiently catalyze the cycloaddition of iminoesters 3a with diethyl maleate in toluene at room temperature with high enantioselective (84% ee) in the absence of base (Table 1, entry 1), only the endo isomer 4a was detected by 1H NMR analysis of reaction mixtures.

Encouraged by these results, the effect of ligands derived from amino acids on the conversions and the enantioselectivities was investigated in toluene (Table 1).1,2 Phenylalanine- and phenylglycine-derived ligands 1b,c were not very effective by comparison with ligand 1a with slightly lower enantioselectivities (Table 1, entries 2 and 3). Delightfully, high enantioselectivity (88% ee) was achieved with the l-tert-leucine-derived ligand 1d (Table 1, entry 4). Next, the influence of the size and chirality of substituent on the terminal amide group was also studied (Table 1, entries 5–8). Four ligands were synthesized by replacing the benzyl group in 1d to 1-((2-naphthyl)methyl) (1e), methyl (1f), (S)-1-phenylethyl (1g), and (R)-1-phenylethyl (1h), respectively. We were pleased to find that the precatalyst 1g, with a (S)-1-phenylethyl moiety incorporated, afforded the desired adduct with >99% yield and 94% ee (Table 1, entry 7). However, when two hydrogen atoms on the terminal amide group are replaced by dibenzyl and dimethyl groups, respectively, the enantioselectivities were sharply decreased (Table 1, entries 9 and 10).

To further optimize the process, different bases were also studied. When the reaction was run with K2CO3, the enantioselectivity was maintained, and the yield was slightly decreased (Table 1, entry 11). Extra Et3N had no particular effect on the yield and ee value (Table 1, entry 12). Other organic bases were also used under the same conditions, lower enantioselectivities (87–93% ee) were obtained (Table 1, entries 13–16). Thus, the optimal conditions for the asymmetric cycloaddition of azomethine ylides are Ag2CO3/1g/toluene with or without Et3N at room temperature.

1,3-Cycloaddition reaction of various iminoesters 3 and diethyl maleate (2a) in the presence of ligand 1g was investigated under the optimized experimental conditions with or without Et3N. Usually the increase in reaction rate will bring a lower selectivity, but the chiral silver AmidPhos catalytic system can efficiently catalyze the cycloaddition of azomethine ylides to dipolarophiles at room temperature with high enantioselective (84% ee) in the absence of base (Table 1, entry 1), only the endo isomer 4a was detected by 1H NMR analysis of reaction mixtures.

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In addition, when R^4^ was 2-naphthyl, 3,4-Cl\(_2^\)C\(_6^\)H\(_3^\), and 2-CIC\(_6^\)H\(_4^\), the endo-4k–m adducts were also successfully obtained with high enantioselectivities (87–96% ee) and yields (65–88%) by using Ag\(_2^\)CO\(_3^\)/1g catalytic system without Et\(_3^N^\). We found the Et\(_3^N^\) had significantly positive influences on the reaction rate, yields, and enantioselectivities. Especially, when R\(^5\) was benzyl or 3-indolylmethyl group, the reaction time was sharply shortened to 24 hours with higher enantioselectivities (94%, 87% ee; Table 3, entries 4 and 5). Moreover, when R\(^5\) was Pheny group, the enantioselectivity was also increased from 62% to 82% with extra Et\(_3^N^\), albeit the reaction did not go to completion (Table 3, entry 6).

Furthermore, we also examined the iminoesters derived from phenylalanine (Table 3, entry 4), tryptophan (Table 3, entry 5), and phenylglycine (Table 3, entry 6) with or without Et\(_3^N^\). We found the Et\(_3^N^\) had significantly positive influences on the reaction rate, yields, and enantioselectivities. Especially, when R\(^5\) was benzyl or 3-indolylmethyl group, the reaction time was sharply shortened to 24 hours with higher enantioselectivities (94%, 87% ee; Table 3, entries 4 and 5). Moreover, when R\(^5\) was Pheny group, the enantioselectivity was also increased from 62% to 82% with extra Et\(_3^N^\), albeit the reaction did not go to completion (Table 3, entry 6).

We also probed other four dipolarophiles in the cycloaddition with 3a as outlined in Figure 1. Only the endo adducts were isolated in all cases. The iminoester 3a reacted perfectly with dimethyl maleate in 94% ee. For dimethyl fumarate and methyl acrylate, much lower enantioselectivities were observed with 30% and 46% ee, respectively. N-Methylmaleimide as a popular dipolarophile in the reported literature was also used to react with α-iminoester 3a with 99% yield and 97% ee.\(^\text{70}\)
In conclusion, we have developed the 1-tert-leucine-derived AmidPhos-silver(I) catalytic system for the asymmetric [3+2] cycloaddition of azomethine ylides with diethyl maleate in high yields and excellent levels of enantioselectivities by using a combination of 2 mol% of ligand 1g with or without Et3N. The study showed the addition of extra Et3N greatly accelerated the reaction rate, increase the yields and the enantioselectivities as well, especially for heterocyclic, aliphatic, and 2-substituted azomethine ylides. In addition, dimethyl maleate, dimethyl fumarate, N-methylmaleimide, and methyl acrylate were also used to react with α-iminoester 3a with high yield and modest to high enantioselectivities further investigation of the reaction scope and detailed mechanism study are under way.

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Supporting Information

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References and Notes


