Organocatalytic synthesis of enantioenriched 1,2,4-triazolines containing a chiral quaternary carbon center†

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An efficient asymmetric cyclization reaction of amino-acid-derived isothiocyanates with azodicarboxylates is described. The chiral 1,2,4-triazolines are prepared in good yield (up to 87%) and enantioselectivities (up to 95% ee).

Triazoles are one of the most important classes of heterocycles because of their broad biological activities such as antiviral, antiepileptic, anticancer, and anti-HIV activities. Many triazoles are used in pharmaceuticals, agrochemicals, corrosion inhibitors, photographic materials, dyes, etc. Therefore, it is important to develop novel methods for the synthesis of structurally diverse triazoles. 1,2,4-Triazolines, one type of important triazole related compounds bearing a potential chiral carbon center at the C-3 position, exhibit good antiviral, anticancer, anti-inflammatory, and anticonvulsant properties. The racemic synthesis of 1,2,4-triazolines has been reported extensively; however, the asymmetric synthetic methods are very limited. In 2010, Jørgensen and co-workers reported preliminary results for the asymmetric synthesis of 1,2,4-triazolines by phase transfer catalysis. Then, Feng, Huang and Shi reported the synthesis of chiral 1,2,4-triazolines independently. In these studies, α-isocyno esters (I) or azlactones (II) reacted with azodicarboxylates to generate the target chiral triazolines (Scheme 1). In 2013, Shi and co-workers reported an efficient [3 + 2] reaction of 3-aminooxindole derived isothiocyanates (III) with azodicarboxylates, producing spirooxindoles in high yields and enantioselectivities. α-Isothiocyanato esters (IV), which can be readily prepared from commercially available amino acids, are useful reactants in the catalytic asymmetric addition reactions with aldehydes, ketones, or imines, affording chiral cyclic thio carbamates in good experimental results. Inspired by Shi’s work, we assumed that the amino acid derived α-isothiocyanato esters would be good reactants in the catalytic asymmetric addition to azodicarboxylates. Such a reaction will produce chiral polystabilised 1,2,4-triazolines bearing a carboxylic functional group at the C-3 position and sulfur substitution at the C-5 carbon (Scheme 1b), and provide an alternative pathway leading to chiral 1,2,4-triazolines. Also, the corresponding products can be converted into structural diversity chiral nitrogen containing heterocycles easily. Herein, we report this chiral tertiary amine catalyzed [3 + 2] reaction of α-isothiocyanato esters with azodicarboxylates, leading to polysubstituted 1,2,4-triazolines in good yields and enantioselectivities.

In the initial study, the phenyl glycine derived isothiocyanato ester 1a was used as reactant to react with di-tert-buty1 azodicarboxylate 2a in the promotion of cinchonine 3a. As expected, the desired product 4a was obtained in a high yield (96%), albeit with a low enantioselectivity (19% ee). We tried to improve the stereocontrol ability of the catalyst by modifying the 9-OH group of cinchonine and cinchonidine. As shown in Table 1, the modified catalysts 3b–d gave 4a in low yields and enantioselectivities (Table 1, entries 2–4). Quinidine 3e and its derivatives 3f–3j were then used as catalysts (Table 1, entries 5–10). Among them, 3f has the best enantioselective control ability (68% ee) (Table 1, entry 6). A comparison of the results gave by catalysts 3d and 3f showed that the methoxy group of 3f benefited the enantioselectivity of 4a significantly; therefore, modified catalysts 3k and 3l were synthesized and used in this reaction.

Scheme 1  Strategy for the synthesis of polysubstituted chiral 1,2,4-triazolines.

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However, no better result was obtained (Table 1, entries 11 and 12). Bifunctional catalysts 3m–3r have been extensively used as chiral organocatalysts, but they were not suitable promoters for this transformation (Table 1, entries 13–18). Therefore, catalyst 3f was selected as the optimal catalyst for further investigating the reaction conditions.

With the optimal catalyst in hand, other reaction conditions were investigated for further improving the yield and enantioselectivity. First, solvents were screened. As shown in Table 2, m-xylene was the best solvent for this transformation. Not only the yield increased to 70%, but the enantioselectivity excess was enhanced to 78% (Table 2, entry 7). Then, this reaction was carried out at a low temperature (−20 °C). The enantioselectivity was further improved to 86% ee (Table 2, entry 8). Additives also affected the reaction outcomes. The yield and enantioselectivity of 4a increased to 77% and 89% ee when Na2SO4 was added to the reaction (Table 2, entry 9).

The substrate scope of the reaction was investigated under the optimal reaction conditions. First, the substituent effects of isothiocyanate esters 1 were studied. The results are shown in Table 3. Electron-withdrawing group substituted isothiocyanates were good reaction partners in this reaction, leading to the chiral 1,2,4-triazolines 4b–4e in good yields and enantioselectivities (Table 3, entries 2–5). The low yield of product 4f (15%) was partially caused by the steric influence of the o-substituent (Table 3, entry 6). We found the electron-donating groups of compounds 1 could give the products 4h–4j in high enantioselectivities, albeit the yield decreased slightly (Table 3, entries 8–10). The 2,5-dimethyl-phenyl-substituted isothiocyanate did not participate in this reaction (Table 3, entry 11). This was probably caused by the steric effect of the bulky o-methyl group. The alkyl groups of compounds 2 and 1 were then investigated. We found the yields increased greatly when the tert-butyl group of 2 was replaced by iso-propyl and ethyl groups, albeit the enantioselectivities decreased (Table 3, entries 12 and 13). These results indicated this transformation would be speeded if an azodicarboxylate bearing small alkyl group was introduced as acceptor. So, compounds 1 having o-substituted phenyls, which could not react with di-tert-butyl azodicarboxylate efficiently (Table 3, entries 6 and 11), were used as donors to react with di-iso-propyl and ethyl azodicarboxylates, respectively (Table 3, entries 17–20). As expected, these reactions took place smoothly, giving the products in moderate to good yields and high to excellent enantioselectivities. Particularly, the highest enantioselectivity (95% ee) in this work was obtained in the reaction of 2-Cl phenyl substituted isothiocyanate with diisopropyl azodicarboxylate (Table 3, entry 17). The damage of enantioselectivities of products 4l and 4m

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**Table 1** Screening catalysts

| Entry | Solvent | T (°C) | Time (h) | Yielda (%) | ee (%) |
|-------|---------|--------|----------|------------|-------|
| 1     | 3a      | 10     | 16       | 96         | 19    |
| 2     | 3b      | 10     | 120      | 10         | <5    |
| 3     | 3c      | 10     | 96       | 11         | <5    |
| 4     | 3d      | 10     | 144      | 25         | −40   |
| 5     | 3e      | 10     | 21       | 64         | <5    |
| 6     | 3f      | 20     | 92       | 22         | −68   |
| 7     | 3g      | 20     | 36       | 14         | −62   |
| 8     | 3h      | 20     | 76       | 9          | −58   |
| 9     | 3i      | 20     | 53       | 16         | 46    |
| 10    | 3j      | 20     | 70       | 29         | <5    |
| 11    | 3k      | 20     | 104      | 19         | 58    |
| 12    | 3l      | 20     | 104      | 14         | 69    |
| 13    | 3m      | 10     | 66       | 29         | <5    |
| 14    | 3n      | 10     | 66       | 29         | <5    |
| 15    | 3o      | 10     | 53       | 29         | <5    |
| 16    | 3p      | 10     | 120      | 20         | 29    |
| 17    | 3q      | 10     | 65       | 10         | 19    |
| 18    | 3r      | 20     | 144      | 20         | <5    |

a Isolated yield. b Determined by chiral HPLC.

**Table 2** Further optimization of reaction conditions

| Entry | Solvent | Time (h) | Yielda (%) | ee (%) |
|-------|---------|----------|------------|-------|
| 1     | Et3O   |          | 89         | 11    |
| 2     | MTBE   |          | 89         | 21    |
| 3     | "BuO   |          | 71         | 52    |
| 4     | DCM    |          | 73         | 55    |
| 5     | CHCl3  |          | 92         | 22    |
| 6     | Toluene|          | 73         | 48    |
| 7     | m-Xylene|         | 64         | 56    |
| 8     | m-Xylene|         | 70         | 57    |
| 9     | m-Xylene|         | 96         | 77    |

a Isolated yield. b Determined by chiral HPLC.
could be complemented through increasing the bulk of alkyl group of isothiocyanate 1. For example, the enantioselectivities increased greatly when we used ethyl and isopropyl isothiocyanate esters as donors for the first time. Triazoline products were obtained in good yields and good to excellent enantioselectivities, and could be converted into other compounds without the loss of enantioselectivities.

### Conclusion

In conclusion, an efficient organocatalytic method was developed for the synthesis of chiral 1,2,4-triazolines, in which, the amino-acid-derived isothiocyanato esters were used as the donors for the first time. Triazoline products were obtained in good yields and good to excellent enantioselectivities, and could be converted into other compounds without the loss of enantioselectivities.

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