Asymmetric Michael/Cyclization Cascade Reaction of 3-Isothiocyanato Oxindoles and 3-Nitroindoles with Amino-Thiocarbamate Catalysts: Enantioselective Synthesis of Polycyclic Spirooxindoles

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ABSTRACT: An unprecedented organocatalytic asymmetric Michael/cyclization cascade reaction of 3-isothiocyanato oxindoles and 3-nitroindoles has been disclosed. A wide range of enantioenriched polycyclic spirooxindoles, containing three contiguous chiral centers with two of them having quaternary stereocenters, could be smoothly obtained with satisfactory results (up to 99% yield, >99:1 dr, and 96% ee). This method is very promising because the reaction is scalable, and the versatile transformations of the products into other spirocyclic oxindoles are also feasible.

Exploring efficient strategies for the construction heterocyclic compounds with rich structural diversity and complexity is a continuing challenge in synthetic chemistry. The spirooxindole skeletons in particular have captured tremendous attention among synthetic and medicinal chemists, due to their prevalence in a broad range of natural and unnatural biologically active products, as well as pharmaceutically important compounds. Over the past several years, we have witnessed rapid progress in the development of creative methodologies for the generation of diverse spirooxindole molecules. Specifically, numerous multifunctional polycyclic spirooxindoles, featuring structural complexity and well-defined three-dimensional architecture, have been demonstrated to correlate with a wide spectrum of biological properties and pharmacological activities (Figure 1). Various synthetic methods for producing functionalized spirooxindole derivatives containing a polycyclic skeleton in asymmetric catalysis have been reported. Despite the substantial advances made thus far, taking into account the importance of the natural-product-like spirocyclic oxindoles in pharmaceutical science and as promising candidates for drug discovery, the development of novel approaches for the efficient synthesis of polyfunctionalized spirooxindoles is in demand.

Since we initially employed 3-isothiocyanato oxindoles as nucleophiles in catalytic asymmetric synthesis, many studies have documented the stereoselective construction of structurally diverse spirocyclic oxindoles using 3-isothiocyanato oxindoles as powerful and versatile precursors via cascade reaction. Notably, 3-isothiocyanato oxindoles can easily undergo Michael/cyclization cascade reactions with some electron-deficient alkenes. In addition, the indoles, bearing two electron-withdrawing substitutions at the N1- and C3-position, have been recognized as a class of electron-deficient alkene reagents. These indoles possess special reaction features; that is, they are readily attacked by nucleophiles at the C2-position and sequentially react with electrophiles at the C3-position (Scheme 1A). We noticed that 3-nitroindoles, a class of potentially promising electrophilic alkenes, had barely been explored in the field of catalytic asymmetric synthesis. In this context, as our continuous interest in developing new synthetic methods for the...

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synthesis of spirocyclic oxindole compounds,\textsuperscript{5,6a−f,10} we envisioned that a novel asymmetric Michael/cyclization cascade reaction between 3-isothiocyanato oxindoles and 3-nitroindoles should take place with an organocatalyst (Scheme 1B). If this strategy is realized, it will permit the rapid construction of a series of polycyclic spirooxindoles containing three contiguous chiral centers with two of them having quaternary stereocenters in a single step. Moreover, this work represents the first example of 3-nitroindoles used in an organocatalytic asymmetric cascade reaction.\textsuperscript{11} Herein we wish to report our preliminary results on this subject.

To examine the feasibility of the proposed Michael/cyclization cascade reaction of 3-isothiocyanato oxindoles and 3-nitroindoles,\textsuperscript{12} our investigation started with the screening of various chiral bifunctional organocatalysts A–D. As shown in Table 1, the reaction of 1a and 2a could proceed to completion within 1 h in mesitylene at 0 °C with 10 mol % A, furnishing the desired spirocyclic oxindole 3a in 98% yield with a 76:24 diastereomeric ratio (dr) and 73% ee for the major diastereomer (entry 1). And then, 3a could be smoothly obtained in 99% yield with 83% ee and 87% ee for the diastereomers using Takemoto’s catalyst B, but the diastereoselectivity was very poor (entry 2). To our delight, a set of good results (99% yield, 97:3 dr, and 92% ee) could be obtained with amino-thiocarbamate catalyst C (entry 3). The 9-thioureia cinchona alkaloid D was inferior to catalyst C for the stereoselectivity (entry 4 vs 3). Afterward, changing the solvent from mesitylene to toluene gave comparable reactivity and stereoselectivity (entry 5), whereas dichloromethane and ethyl ether resulted in a dramatic decrease in yield and stereoselectivity (entries 6–7). After addition of molecular sieves (MS) as an additive, no improvement for the dr and ee was observed (entries 8–9). We were gratified to find that the ee value could be elevated to 96% by increasing the reaction concentration (entry 10). Unfortunately, changing the reaction temperature gave rise to a slightly lower ee value for the major diastereomer (entries 11–13). Ultimately, with 5 and 1 mol % C for the reaction, respectively, 3a also could be obtained with good results (entries 14–15).

The scope of the cascade reaction was then investigated with catalyst C under the optimized conditions used in Table 1, entry 10. First, we focused on the examination of 3-nitroindole substrates by using 3-isothiocyanato oxindole 1a as the donor. As summarized in Table 2, with electron-withdrawing or -donating substituents at the C5-position of 3-nitroindoles, reactions with 1a presented very high reactivity and furnished the corresponding products in 94–99% yield with up to >99:1 dr and 95% ee (entries 1–4). Moreover, different substituents in the C4-, C6-, and C7-position also could give rise to very high reactivity and deliver the desired products with excellent results (entries 5–8). After the investigations on the varied sulfonly protecting groups of the N1- in 3-nitroindoles, it was found that the sulfonly protecting groups, such as Bs-, Ms-, and Ns-, had a dramatic effect on the diastereo- and enantioselectivity, while the reactivity seemed hardly affected (entries 9–11). Similarly, the N-acetylated and N-alkoxy carbonylated 3-nitroindoles generated the expected spirocyclic oxindoles 3m–p in excellent yields with moderate to good dr and ee values (entries 12–15). On the other hand, a survey of 3-isothiocyanato oxindole substrates was also conducted. The reactions of 3-isothiocyanato oxindoles 1b–d, bearing different N-protecting groups, with 3-nitroindole 2a occurred with complete conversion after 2 h, giving good diastereoselectivities and moderate enantioselectivities (entries 16–18). These results revealed the steric hindrance of the N1-position of 3-isothiocyanato oxindoles was crucial to the stereocenters and diastereoselectivity of the Michael and cyclization steps (Scheme 2).

In order to examine the utility of the methodology, a gram scale experiment between 1a and 2a was carried out under the optimized conditions. As shown in Scheme 2, the reaction...
Table 2. Substrate Scope Examinationc

| R1 | R2 | Yield (%) | dr | ee (%) |
|----|----|-----------|----|--------|
| (H, Boc) | (3a) | 3 | 79:21 | 36 |
| (H, Boc) | (3b) | 2 | 30:70 | 60 |
| (H, Boc) | (3c) | 1 | 50:50 | 50 |
| (H, Boc) | (3d) | 0 | 0:100 | 0 |

cUnless noted, the reactions were carried out with 1 (0.1 mmol), 2 (0.1 mmol), and 10 mol % C in 1.0 mL of mesitylene at 0 °C. Yield of isolated product as a mixture of diastereoisomers. Determined by 1H NMR analysis of crude reaction mixture. Ee of major diastereomer was determined by chiral HPLC analysis after the product reacting with CH3I according to the reported procedure.5 The results in parentheses were the ee value for the minor diastereomer. Ts = Toluenesulfonyl; Bn = Benzenesulfonyl; Ms = Methanesulfonyl; Ns = 4-Nitrobenzenesulfonyl; Ac = Acetyl; Cbz = Carbobenzyloxyl; Boc = Butyloxycarboryl.

Scheme 2. Reaction of 1a and 2a on a Gram Scale

proceeded cleanly with 10 mol % C after 2 h and furnished 3a in excellent results, which are similar to the results of the original reaction illustrated in Table 1, entry 10. Moreover, upon enlarging the scale of the original reaction by 10-fold, even in the presence of 1 mol % C, excellent results were still obtained. These observations reflected the present protocol was amenable to large scale production.

To further expand the potential of this methodology, versatile transformations of the product into some other structurally diverse spirocyclic oxindoles were performed (Scheme 3).13 Product 3a could be readily transformed into compound 4 with iodomethane and K2CO3 according to the reported procedure.5 Similarly, reacting 3a with benzyl bromide gave compound 5 bearing a benzylated thiolactam ring. Notably, 4 and 5 could be obtained in nearly quantitative yields and no changes happened in dr and ee values during the conversions. Compound 4 was converted to spirocyclic oxindole 8 containing a single spirostereocenter in 67% yield with 87% ee under strongly acidic conditions. Additionally, the nitro substituent in 4 could be eliminated by nickel boride reduction in MeOH, generating 9 in 67% yield with excellent stereoselectivity. Treatment of 3a with a solution of 30% aqueous hydrogen peroxide and formic acid according to the reported method5 gave compound 6 in 90% yield with >99:1 dr and 96% ee. After reduction by zinc powder and TMSCl, the nitro group in 3a was transformed to an amine functionality, affording product 7 in 61% yield, without loss in stereoselectivity.

The relative and absolute configuration of 4 was unequivocally established by X-ray analysis,13 thus leading to the determination of the configuration of 3a. The remaining configurations of other products were assigned on the assumption of a uniform mechanistic pathway. Based on previous studies on isothiocyanato oxindoles6 and our experimental results, a plausible transition state model is also proposed to account for the observed enantioselectivity (see Supporting Information).

In conclusion, we have developed an efficient asymmetric Michael/cyclization cascade reaction of 3-isothiocyanato oxindoles and 3-nitroindoles for the construction of spirocyclic oxindoles using an amino-thiocarbamate catalyst. With the developed protocol, a wide range of enantioenriched polycyclic spirooxindoles, containing three contiguous chiral centers with two of them having quaternary stereocenters, could be smoothly obtained with satisfactory results (up to 99% yield, >99:1 dr, and 96% ee) under mild conditions. This method is very promising because the reaction is scalable and the versatile transformations of the products into other structurally diverse spirocyclic oxindoles were also feasible. Additionally, this methodology represents the first example regarding organocatalytic enantioselective reactions of 3-nitroindoles. Application of this methodology toward library synthesis and subsequent biological evaluation of its members is underway.

ASSOCIATED CONTENT

 Supporting Information
Experimental details, characterization data for new compounds, X-ray crystal structure and the CIF files of 4. 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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