Synthesis of Optically Active Oxazolines by an Organocatalytic Isocyanoacetate Aldol Reaction with α-Keto Esters

Fei Wang
Jiean Chen*
Yong Huang*

Key Laboratory of Chemical Genomics, School of Chemical Biology and Biotechnology, Peking University, Shenzhen Graduate School, Shenzhen, 518055, P. R. of China
chenja@pkusz.edu.cn
huangyong@pkusz.edu.cn

Abstract
An enantioselective [3+2] cyclization is reported for the construction of a chiral oxazoline skeleton in moderate yield and up to 97% ee. The reactivity and stereochemical discrimination originate from the noncovalent interaction and orientation of a bifunctional catalyst. The novel combination of an α-keto ester and an α-isocyanoacetate establishes an oxazoline which could be a potential chiral ligand for metal-mediated catalysis, and also could be easily converted into an optically active β-hydroxy-α-amino acid.

Key words organocatalysis, bifunctional catalyst, noncovalent interaction, asymmetric cyclization, chiral oxazoline

Nitrogen-containing heterocycles are important in pharmaceutical investigations, especially for small-molecule drug design.1 Aimed at target skeletons of this sort, α-isocyanoacetate has been frequently used as versatile synthon providing dipolar reactivity in a [3+2] cyclization process.2 Mediated by organocatalysis, several prominent manipulations have been used for the construction of analogues of pyrrole3 and imidazole.4 However, the straightforward assembly of optically active oxazolines, which act as ligands for transition metals and are the core framework in numerous natural products, is highly dependent on chiral metal complexes. In 1986, Ito and Hayashi reported the first asymmetric aldol-type reaction for α-isocyanoacetate using a chiral Au(I) complex.5 Subsequent investigations focused on disparate transition metal catalyst systems and often led to the formation of trans-substituted oxazolines.6 In 2011 Dixon et al. utilized a chiral amino phosphine Ag(I) complex to obtain oxazolines with excellent cis selectivity.7 In sharp contrast, the organocatalytic variant for such conversions has been studied and only a single example was presented by Gong et al.8 There has been no report of the combination involving a ketone group, which would afford direct access to oxazolines bearing one or two quaternary asymmetric centers. In this paper, we describe the first asymmetric aldol-type transformation of α-keto esters with α-isocyanoacetate catalyzed by a thiourea/amine bifunctional catalyst and leading to a precursor of β-hydroxy-α-amino acids.9a-c,7,9

We began by evaluating the efficiency of the bifunctional catalyst system.10 Progress in noncovalent catalysis furnishes findings, abundant and well-documented, concerning versatile methodologies and selectivities.11 With thiourea/amine-type bifunctional catalysts, the Brønsted basicity of a tertiary amine affords the activation energy for reaction with nucleophilic compounds and simultaneously the thiourea moiety acts as a hydrogen donor to interact with the corresponding electrophile.12 This cooperative mode takes advantage of both the substrate proximity effect and a sterically well-defined transition state rendering high synergy to the catalysis. We hypothesized that the dual carbonyl groups of α-keto esters would permit a well-organized cyclic interaction with the thiourea moiety of the bifunctional catalyst, as shown in Scheme 1. Simultaneously, the acidic C–H bond of α-isocyanoacetate spontaneously enters into a noncovalent interaction with the alkaline site and interacts intramolecularly with the keto group of the α-keto ester.

The inception of the aldol-cyclization cascade is initiated by the reaction of ethyl phenylglyoxylate (1a) with methyl isocyanoacetate (2a) which is promoted by various thiourea amines. Takemoto’s catalyst (3a; Table 1, entry 1), the simplest bifunctional analogue of chiral 1,2-diaminocyclohexane, delivers fair asymmetric induction along with moderate differentiation of diastereoisomers. The results indicate that thioureas containing a strong basic tertiary amine two carbon atoms removed from the thio carbonyl group could lead to activation and the resulting stereochemistry.

The comparable structure in the cinchona alkaloid system could lead to activation and the resulting stereochemistry.

Central to this bifunctional catalyst system is the thiourea moiety, which acts as a hydrogen bond acceptor and provides high synergism to the catalysis. The thiourea/amine-type bifunctional catalyst is designed to afford compounds with high selectivity. The thiourea moiety provides high synergism to the catalysis. The thiourea moiety of the bifunctional catalyst is designed to afford compounds with high selectivity. The thiourea moiety of the bifunctional catalyst is designed to afford compounds with high selectivity.
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loids (3c–g) was systematically investigated in the reaction and the essential role of the thiourea moiety was also evaluated (3b). Catalysts lacking thiourea can nevertheless catalyze the oxazoline synthesis, but with noticeably diminished enantioselectivity and yield (Table 1, entry 2, 48% yield, 33% ee). The aryl substituents of the thiourea group demonstrate no obvious discrepancy and promote the cyclization with moderate efficiency and enantioselectivity (entries 3 and 4, Table 1, 68% ee vs. 76% ee). In a brief screening of the solvent system, toluene was identified as a suitable solvent, delivering acceptable reactivity and selectivity (Table 1, entries 8 and 9).

The substituent effect was evaluated by modifying the R1, R2 and R3 groups in the starting materials. The substituent in the α-position of the isocyanoacetate (R3) triggered selectivity which might result from excessive crowding in the transition state (entry 10, Table 1; methyl substituent for 53% ee). When manipulating the steric size of ester groups in the isocyanoacetates, a proportional increase of enantioselectivity indicates that the bulky tert-butyl ester could enhance the facial discrimination and generate an appreciable ee value (Table 1, entries 13 and 14). The absolute configurations of the thiourea and quinuclidine moieties were found to control the steric orientation of the transition state and the reaction furnished the desired product with determinable conformation. The same reactions for cinchonidine (3e), quinidine (3f) and cinchonine (3g) showed no significant difference in both catalytic ability and steric discrimination (Table 1, entries 15–17). Catalysts 3f and 3g generated the desired product with a structure which is the mirror image of that in 3c. In view of the experimental data and the demonstrated catalytic model, the absolute configuration of the title product was assigned and is shown in Scheme 2.

Under the optimized reaction conditions, diverse aromatic α-keto esters were examined and the corresponding oxazolines were synthesized in good yields and with excellent ee values.13 As shown in Table 2, ethyl phenylglyoxylate structures containing electron-donating substituents (Me, OMe) or electron-withdrawing groups (F, Cl, Br) gave the corresponding products with appreciable enantioselectivity (Table 2, entries 2–9). Different patterns of substituents (m- and p-) of substituents with different electronic characteristics are tolerated in the reaction. Compounds with a multisubstituted aryl group also performed well giving an excellent ee value for the title oxazoline (entry 10). Furthermore, α-keto esters with a heteroaromatic ring and a fused ring can also be converted smoothly into the desired products with appropriate selectivity (Table 2, entries 11 and 12).

Derived from Nuclear Overhauser Effect Spectroscopy (NOESY) analysis (cis product)14 and the well-understood interaction pattern between the thiourea moiety and α-keto ester,15 a proposed transition state is shown in Scheme 2. The ester carbonyl group of the α-keto ester enjoys an H-bonding interaction with the N atom bearing an aryl group and the ketone carbonyl interacts with the residual N atom of the thiourea. The nucleophilic α-isocyanoacetate is activated by the quinuclidine moiety and attacks from above the rigid ring system. As a consequence, the substitu-
Table 1 Reaction Optimization

<table>
<thead>
<tr>
<th>Entry</th>
<th>R1</th>
<th>R2</th>
<th>R3</th>
<th>Solvent</th>
<th>Yield (%)</th>
<th>dr</th>
<th>ee (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>3a</td>
<td>Et, Me, H</td>
<td>(2a)</td>
<td>CH2Cl2</td>
<td>53</td>
<td>1:8:1</td>
<td>70</td>
</tr>
<tr>
<td>2</td>
<td>3b</td>
<td>Et, Me, H</td>
<td>(2a)</td>
<td>CH2Cl2</td>
<td>48</td>
<td>1:2:1</td>
<td>33</td>
</tr>
<tr>
<td>3</td>
<td>3c</td>
<td>Et, Me, H</td>
<td>(2a)</td>
<td>CH2Cl2</td>
<td>53</td>
<td>1:3:1</td>
<td>68</td>
</tr>
<tr>
<td>4</td>
<td>3d</td>
<td>Et, Me, H</td>
<td>(2a)</td>
<td>CH2Cl2</td>
<td>58</td>
<td>1:4:1</td>
<td>76</td>
</tr>
<tr>
<td>5</td>
<td>3e</td>
<td>Et, Me, H</td>
<td>(2a)</td>
<td>EtO</td>
<td>56</td>
<td>1:4:1</td>
<td>56</td>
</tr>
<tr>
<td>6</td>
<td>3f</td>
<td>Et, Me, H</td>
<td>(2a)</td>
<td>THF</td>
<td>trace</td>
<td>n.d.</td>
<td>n.d.</td>
</tr>
<tr>
<td>7</td>
<td>3g</td>
<td>Et, Me, H</td>
<td>(2a)</td>
<td>EtOAc</td>
<td>36</td>
<td>1:5:1</td>
<td>76</td>
</tr>
<tr>
<td>8</td>
<td>3h</td>
<td>Et, Me, H</td>
<td>(2a)</td>
<td>toluene</td>
<td>65</td>
<td>2:1:1</td>
<td>84</td>
</tr>
<tr>
<td>9</td>
<td>3i</td>
<td>Et, Me, H</td>
<td>(2a)</td>
<td>toluene</td>
<td>70</td>
<td>2:1:1</td>
<td>79</td>
</tr>
<tr>
<td>10</td>
<td>3j</td>
<td>Et, Me, Me</td>
<td>(2b)</td>
<td>toluene</td>
<td>62</td>
<td>2:1:1</td>
<td>53</td>
</tr>
<tr>
<td>11</td>
<td>3k</td>
<td>Et, Et, H</td>
<td>(2c)</td>
<td>toluene</td>
<td>46</td>
<td>1:7:1</td>
<td>84</td>
</tr>
<tr>
<td>12</td>
<td>3l</td>
<td>Et, t-Bu, H</td>
<td>(2d)</td>
<td>toluene</td>
<td>75</td>
<td>2:1:1</td>
<td>94</td>
</tr>
<tr>
<td>13</td>
<td>3m</td>
<td>Me, t-Bu, H</td>
<td>(2e)</td>
<td>toluene</td>
<td>67</td>
<td>2:1:1</td>
<td>93</td>
</tr>
<tr>
<td>14</td>
<td>3n</td>
<td>Bn, t-Bu, H</td>
<td>(2f)</td>
<td>toluene</td>
<td>60</td>
<td>1:1:1</td>
<td>94</td>
</tr>
<tr>
<td>15</td>
<td>3o</td>
<td>Et, t-Bu, H</td>
<td>(2d)</td>
<td>toluene</td>
<td>75</td>
<td>2:1:1</td>
<td>91</td>
</tr>
<tr>
<td>16</td>
<td>3p</td>
<td>Et, t-Bu, H</td>
<td>(2d)</td>
<td>toluene</td>
<td>75</td>
<td>2:1:1</td>
<td>91</td>
</tr>
<tr>
<td>17</td>
<td>3q</td>
<td>Et, t-Bu, H</td>
<td>(2d)</td>
<td>toluene</td>
<td>75</td>
<td>2:1:1</td>
<td>89</td>
</tr>
</tbody>
</table>

*Unless otherwise noted, all reactions were carried out using 1a–I (0.1 mmol), 2a (0.12 mmol) and the catalyst (10 mol%) in solvent (0.5 mL) at 26 °C for 24 h.

Table 2 Scope of the α-Keto Esters

<table>
<thead>
<tr>
<th>Entry</th>
<th>R</th>
<th>Yield (%)</th>
<th>dr</th>
<th>ee (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Ph (1a)</td>
<td>75</td>
<td>2:1</td>
<td>94 (81e)</td>
</tr>
<tr>
<td>2</td>
<td>4-MeC6H4 (1b)</td>
<td>73</td>
<td>2:1</td>
<td>94</td>
</tr>
<tr>
<td>3</td>
<td>3-MeC6H4 (1c)</td>
<td>67</td>
<td>2:1</td>
<td>91</td>
</tr>
<tr>
<td>4</td>
<td>4-MeOC6H4 (1d)</td>
<td>78</td>
<td>2:1</td>
<td>97</td>
</tr>
<tr>
<td>5</td>
<td>3-MeOC6H4 (1e)</td>
<td>72</td>
<td>2:1</td>
<td>94</td>
</tr>
<tr>
<td>6</td>
<td>4-FC6H4 (1f)</td>
<td>70</td>
<td>2:1</td>
<td>92</td>
</tr>
<tr>
<td>7</td>
<td>3-FC6H4 (1g)</td>
<td>76</td>
<td>2:1</td>
<td>91</td>
</tr>
<tr>
<td>8</td>
<td>4-ClC6H4 (1h)</td>
<td>73</td>
<td>2:1</td>
<td>93</td>
</tr>
<tr>
<td>9</td>
<td>4-BrC6H4 (1i)</td>
<td>77</td>
<td>2:1</td>
<td>91</td>
</tr>
<tr>
<td>10</td>
<td>3-F-6-MeC6H3 (1j)</td>
<td>75</td>
<td>2:1</td>
<td>97</td>
</tr>
<tr>
<td>11</td>
<td>2-naphthyl (1k)</td>
<td>84</td>
<td>2:1</td>
<td>90</td>
</tr>
<tr>
<td>12</td>
<td>2-thienyl (1l)</td>
<td>80</td>
<td>2:1</td>
<td>89</td>
</tr>
</tbody>
</table>

*Unless otherwise noted, all reactions were carried out using 1a–I (0.1 mmol), 2d (0.12 mmol) and 3c (10 mol%) in toluene (0.5 mL) at 26 °C for 24 h.

In summary, an asymmetric isocyanoacetate aldol reaction of α-keto esters was realized by a bifunctional catalytic strategy. Aromatic and heteroaromatic α-keto esters are converted into oxazolines in good yield (up to 84%) and with high enantioselectivity (up to 97% ee). This provides practical synthetic access to chiral oxazolines which can be precursors to β-hydroxy-α-amino acids. Further efforts are aimed at improving the diastereoselectivity and elucidating the synthetic utility of the reaction in pharmaceutical chemistry.

Acknowledgment

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Supporting Information

Supporting information for this article is available online at https://doi.org/10.1055/s-0036-1588718.
References and Notes


(13) For selected reviews of thiourea-amine-type esters such as ethyl 2-cyclohexyl-2-oxoacetate were also used, but only gave 59% ee.

(14) For selected reviews of thiourea-amine-type esters such as ethyl 2-cyclohexyl-2-oxoacetate were also used, but only gave 59% ee.


(16) General Procedure for the Synthesis of Chiral Oxazoline: Phenylglyoxylate (1 mol mmol), tert-butyl isocyanatoacetate (2d; 11.9 mg, 0.12 mmol) and bifunctional catalyst 3c (5.9 mg, 0.01 mmol) were stirred in toluene (0.5 mL) at 26 °C for 24 h. The mixture was separated by silica gel chromatography (10% EtOAc/petroleum ether) and gave product 4.
Analysis data for compound 4a: ¹H NMR (400 MHz, CDCl₃): \(\delta = 1.20-1.24\) (m, 3 H), 1.52 (s, 9 H), 4.11-4.24 (m, 2 H), 4.88 (d, \(J = 2.0\) Hz, 1 H), 7.14 (d, \(J = 2.0\) Hz, 1 H), 7.31–7.56 (m, 2 H), 7.58–7.60 (m, 2 H). ¹³C NMR (100 MHz, CDCl₃): \(\delta = 14.0, 28.0, 62.5, 79.7, 83.0, 89.4, 125.3, 129.0, 138.7, 155.5, 168.2, 168.6\). HPLC (Chiralcel IB, hexane/i-PrOH = 95:5, flow rate: 1.0 mL/min, \(\lambda = 210\) nm): \(t_R\) (major) = 18.08 min, \(t_R\) (minor) = 20.22 min. ESI–HRMS: \(m/z\) [M + H] calcd for C₁₇H₂₁NO₅: 320.1498; found: 320.1492.