Lewis acid-catalyzed asymmetric reactions of β, γ-unsaturated 2-acyl imidazoles

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The investigation of diverse reactivity of β,γ-unsaturated carbonyl compounds is of great value in asymmetric catalytic synthesis. Numerous enantioselective transformations have been well developed with β,γ-unsaturated carbonyl compounds as nucleophiles, however, few example were realized by utilizing them as not only nucleophiles but also electrophiles under a same catalytic system. Here we report a regioselective catalytic asymmetric tandem isomerization/α-Michael addition of β,γ-unsaturated 2-acyl imidazoles in the presence of chiral N,N′-dioxide metal complexes, delivering a broad range of optically pure 1,5-dicarbonyl compounds with two vicinal tertiary carbon stereocenters in up to >99% ee under mild conditions. Meanwhile, stereodivergent synthesis is disclosed to yield all four stereoisomers of products. Control experiments suggest an isomerization process involved in the reaction and give an insight into the role of NEt3. In addition, Mannich reaction and sulfur-Michael addition of β,γ-unsaturated 2-acyl imidazoles proceed smoothly as well under the same catalytic system.
The exploration of reaction diversity from β,γ-unsaturated carbonyl compounds is interesting and of great synthetic value. These compounds and their analogs bearing one potential enolization have been demonstrated as highly active nucleophiles in a number of catalytic asymmetric reactions for the synthesis of natural products and bioactive compounds 1–16. Especially, γ-addition as dienolate pronucleophiles with either metal catalysis 17–28 or organocatalysis 29–36 has been widely documented during the past several years, and the maintained π-conjugation of γ-addition process leading to thermodynamically stable conjugated products (Fig. 1a, A). The regioselectivity changing from γ-addition to α-addition seems to be plaguing 37,38, and α-addition of specific substrates, such as γ,γ-disubstituted ones, has been reported 39–43. Notably, in some cases, C=C isomerization occurred after α-addition which further expanded the reaction diversity (Fig. 1a, B) 44–46.

Although versatile catalytic asymmetric reactions have been demonstrated by utilizing β,γ-unsaturated carbonyl compounds as mentioned above, however, few examples were investigated by employing them as electrophiles upon isomerization to conjugated α,β-unsaturated carbonyl compounds (Fig. 1a, C) 47,48. We envision that, by careful design of β,γ-unsaturated carbonyl compounds, these could serve not only as nucleophiles but also electrophiles. Based on this assumption, here we report the synthesis of a series of β,γ-unsaturated 2-acyl imidazoles by introducing an imidazole moiety which would address the...

Fig. 1 Strategies for γ- and α-addition of β,γ-unsaturated carbonyl compounds. a Regioselectivity of deconjugated carbonyl compounds. b Our strategies for diverse reactivity of β,γ-unsaturated 2-acyl imidazoles.
following two points: (1) bidentate coordination with a Lewis acid of acyl imidazole exhibits good stereocontrol\(^{49-55}\) and (2) the strong coordination facilitates isomerization of the \(\beta,\gamma\)-unsaturated ketone to an \(\alpha,\beta\)-unsaturated ketone. Chiral \(\text{N},\text{N}'\)-dioxide-metal\(^{56-59}\) complexes catalyze diverse reactions of \(\beta,\gamma\)-unsaturated 2-acyl imidazoles, including tandem isomerization/\(\alpha\)-Michael addition (Fig. 1b, D), Mannich reaction (Fig. 1b, B), and sulfur-Michael addition (Fig. 1b, C) with high efficiency and stereoinduction. In addition, stereodivergent catalysis\(^{60-63}\) is also disclosed and provides a unified and predictable route for the access to all four stereoisomers of 1,5-dicarbonyl compounds by matching the configuration between the Lewis acid catalysts and substrates.

### Results

#### Optimization of the reaction conditions.

We began our study by employing \(\beta,\gamma\)-unsaturated 2-acyl imidazole \(E\)-1a as the model substrate to optimize the reaction conditions. Several metal salts coordinated with the \(\text{N},\text{N}'\)-dioxide ligand \(L_3\text{-RaPr}_2\) (Fig. 2) were evaluated, such as \(\text{Sc(O Tf)}_3\), \(\text{Ni(O Tf)}_2\), and \(\text{Mg(O Tf)}_2\); however, only trace amount of the self-\(\alpha,\beta\)-addition product \(2a\) was observed, which was generated from \(\alpha\)-addition of \(E\)-1a with the corresponding \(\alpha,\beta\)-unsaturated 2-acyl imidazole upon C–C isomerization (Table 1, entry 1). Pleasingly, the \(\text{Y(O Tf)}_3/L_3\text{-RaPr}_2\) complex was efficient to promote the tandem isomerization/\(\alpha\)-Michael addition and provided the corresponding product \(2a\) with 60% yield, 2.2:1 \textit{anti:syn} ratio, and 96% ee in \(\text{CH}_2\text{Cl}_2\).

![Fig. 2 Representative chiral N,N'-dioxide ligands used in the study.](image)

**Table 1 Optimization of the reaction conditions.**

| Entry | Metal Salt | Yield (%) | \textit{anti:syn} \text{Ratio} | ee (%) |
|-------|------------|-----------|-------------------------------|--------|
| 1     | \(\text{Sc(O Tf)}_3\)/\(\text{Ni(O Tf)}_2\)/\(\text{Mg(O Tf)}_2\) | Trace |  —                          |  —     |
| 2     | \(\text{Y(O Tf)}_3\) | 60 | 2.2:1                       | 96/−34 |
| 3     | \(\text{La(O Tf)}_3\) | 58 | 2.7:1                       | 92/63  |
| 4     | \(\text{Yb(O Tf)}_3\) | 46 | 2.2:1                       | 84/13  |
| 5\(d\) | \(\text{Y(O Tf)}_3\) | 73 | 5.2:1                       | 97/0   |
| 6\(d\) | \(\text{Y(O Tf)}_3\) | 74 | 10:1                        | 98/N.D.|

Unless otherwise noted, all reactions were performed with metal salt/ligand (1:1, 2.5 mol%), \(E\)-1a (0.20 mmol) in \(\text{CH}_2\text{Cl}_2\) (1.0 mL) at 25 °C under \(N_2\) atmosphere for 24 h. \(d\)Isolated yield of anti-synmer. \(e\)Determined by \(^1\)H NMR analysis of crude products. \(f\)Determined by HPLC analysis on a chiral stationary phase. \(g\)Toluene was used as solvent. \(h\)Addition of \(\text{NEt}_3\) (10 mol%) and for 12 h.

Substrate scope in isomerization/\(\alpha\)-Michael addition reaction.

The generality of the tandem isomerization/\(\alpha\)-Michael addition reaction was investigated under the optimized conditions (Fig. 3). An array of \(\beta,\gamma\)-unsaturated 2-acyl imidazoles bearing different substituents on the \(\gamma\)-phenyl group (both electron-withdrawing and electron-donating groups at the \textit{para}, \textit{meta}, or \textit{ortho}-positions) were converted into the corresponding dimerization products (entry 2). Lanthanide metal salts \(\text{La(O Tf)}_3\) and \(\text{Yb(O Tf)}_3\) could also mediate the reaction but gave lower yields and ee values (entries 3 and 4). The screening of chiral backbones and steric hindrance of the amide moiety on the \(\text{N},\text{N}'\)-dioxide ligands afforded no better results (for details, see Supplementary Table 1). When toluene was used as solvent instead, the isolated yield of anti-\(2a\) was increased to 73% with 5.2:1 dr and 97% ee (entry 5). To our delight, the diastereoselectivity could be improved to 10:1 with addition of \(\text{NEt}_3\) (entry 6). Other common chiral ligands such as Box, Pybox, and BINAP were also explored, and 32% yield, 5:1 dr with 60% ee were observed as the best results (for details, see Supplementary Table 3).

![Fig. 3 Substrate scope in isomerization/\(\alpha\)-Michael addition reaction.](image)
exhibited high tolerance as well, generating the desired products 2n–2q with a high level of yields (63–84%) and stereoselectivities (9:1 to >19:1 dr; 92–99% ee). Estrone-derived 1r could be transformed into 2r smoothly in 69% yield, 2.8:1 E/Z, >19:1 dr, and 99% ee for E-isomer. Other Michael acceptors such as α,β-unsaturated 2-acyl imidazole and ethyl vinyl ketone were also suitable in this reaction, delivering 2s–2w with good yields (60–71%) and stereoselectivities (6:1 dr, 91–99% ee). The absolute configuration of 2j was determined to be (2S, 3R) by X-ray crystallography analysis.

Substrate scope in α-Mannich reaction of β,γ-unsaturated 2-acyl imidazoles and imines. The reaction described above indicated that β,γ-unsaturated 2-acyl imidazoles performed both α-addition reaction and β-addition upon isomerization under proper Lewis acid catalysts. Next, to extend the scope of α-addition of β,γ-unsaturated 2-acyl imidazoles, several types of imines 3 were explored as the electrophiles. By switching the catalyst to La(OTf)_3/L_3-PiBu complex (for detailed screening of the conditions, see Supplementary Table 4), the Mannich reaction between E-1 and isatin-derived ketimines 3a–3h was successfully realized to deliver the desired β-amino 2-acyl imidazoles 4a–4h as single isomers in 75–99% yields and 88–91% ee (Fig. 4a).

Substrate scope in isomerization/sulfur-Michael reaction. Inspired by the isomerization process of β,γ-unsaturated 2-acyl imidazoles into α,β-unsaturated 2-acyl imidazoles, we next enlarged the diverse reactivity of β,γ-unsaturated compounds as the electrophiles under the current catalytic system. However, only a trace amount of desired tandem isomerization/sulfur-Michael addition product 6a was achieved if E-1a reacted with thiophenol 5a. After examination of the reaction conditions (for details, see Supplementary Table 5), Z-1a was used instead, and 6a could be obtained in 89% yield with 90% ee (Fig. 5). The scope of isomerization/sulfur-Michael reaction was investigated next.
Substrate scope with pyrazolinone-derived ketimins.

**Gram-scale synthesis and derivatization of products.** To evaluate the synthetic utility of this methodology, a gram-scale synthesis of 2a was conducted. The current reaction could be carried out at 7.0 mmol scale without loss of yield (70%), diastereoselectivity (10:1 dr), and ee value (98%) (Fig. 6a). Furthermore, hydrogenation of 2a in the presence of Pd/C and H2 afforded derivative 7 in 98% yield with 98% ee (Fig. 6b). Chiral sulfone motif is found in numerous biological compounds as well as drug candidates. Upon treatment of 6a with m-CPBA, the oxidized sulfone product 8 was obtained in 85% yield with 90% ee. Moreover, 6a went through further transformations to afford sulfone 9 in 50% yield with 85% ee (Fig. 6c).

Mechanistic studies. To gain insight into the mechanism of tandem isomerization/α-Michael addition, some control experiments were carried out. Firstly, we wondered why the addition of NEt3 led to an increase in diastereoselectivity (Table 1, entry 6). Treating the product 2a (2.9:1 dr, 85%/12% ee) under the standard conditions for 12 h (for details, see Supplementary Note 5), no change of enantioselectivity and diastereoselectivity was observed, which ruled out the possibility that the diastereoselectivity increased via epimerization of syn-2a (Fig. 6d).

Thiolphenols and alkyl-substituted thiols could be converted into the final products in 39–95% yields with 70–93% ee values. For the Michael acceptors, aryl- and alkyl-substituted βγ-unsaturated 2-acyl imidazoles were also tolerated in this reaction, giving 6j–6p in 60–92% yields with 80–92% ee.

![Diagram](https://example.com/diagram.png)
Moreover, when \(Z-\alpha,\beta\)-unsaturated 2-acyl imidazole \(Z-10\) was used to react with \(E-1\), the product \(2\) was obtained in 1:5.2 \(E\):\(Z\) after 2 h, and decreased to 1:2.8 \(E\):\(Z\) after 5 h (Fig. 7b). These experiments confirmed the isomerization of \(\beta,\gamma\)-unsaturated \(C=\)C bond into \(\alpha,\beta\)-unsaturated \(C=\)C bond in the presence of \(N,N\)-dioxide-metal complexes, and this process was likely to be the rate-determining step. It also suggests the diastereoselectivity was mainly controlled by the \(E/Z\)-configuration of the \(\alpha,\beta\)-unsaturated 2-acyl imidazole intermediate, and the addition of \(\text{NEt}_3\) might improve the \(E/Z\) ratio during the isomerization process. As a result of equilibrium between \(E-1\), \(E-10\), and \(Z-10\) (Fig. 7c), the use of \(E-10\) as the starting substrate alone, albeit unstable yielded the corresponding \(E-2\) as the major product in 98% ee after 3 h (Fig. 7d), while the reaction from only \(Z-10\) gave the \(Z-2\) product in 60% isolated yield and 92% ee (Fig. 7e). In addition, operando IR experiments were also performed to interpret the reaction process (for details, see Supplementary Note 7). Furthermore, we set out to establish the

### Fig. 5 Substrate scope in isomerization/sulfur-Michael reaction.

Unless otherwise noted, all reactions were performed with \(\text{Y(OTf)}_2/\text{L}_3\text{-PrPr}_2\) (1:1, 5 mol%), \(1\) (0.25 mmol), \(5\) (0.10 mmol) in \(\text{CH}_2\text{ClCHCl}_2\) (1.0 mL) at 25 °C for 17 h. [a] \(Z/E\) mixture of \(\beta,\gamma\)-unsaturated 2-acyl imidazole was used for 6n. The reaction time was 5 days.

### Fig. 6 Gram-scale synthesis and derivatization of products.

**a** Gram-scale synthesis of \(2\). **b** Hydrogenation of \(2\). **c** The derivatization of \(6\). (1) \(\Phi\text{MgBr}, \text{THF}\); (2) \(\text{MeOTf}, \text{MeCN}\); (3) \(\text{K}_2\text{CO}_3\) (aq); (4) \(m\text{-CPBA}, \text{CH}_2\text{Cl}_2\).
availability of stereodivergent access to 2a. All four stereoisomers of 2a could be readily obtained in good yields (67–85%) and diastereoselectivities (8:1–>19:1) with excellent ee values by matching the E/Z-configured 10 and the chiral ligand (Fig. 7f).

**Proposed catalytic cycle.** Based on the absolute configuration of the product 2j, control experiments and our previous studies, a possible catalytic cycle with a transition-state model was proposed (Fig. 8). First, the coordination of chiral N,N'-dioxide L3-RaPr2 and metal salt in situ to form chiral metal complex (Y*). Then, the β,γ-unsaturated ketone E-1a attaches to Y* as a dianolate of NEt3 to give the intermediate T1, and which partly transforms into the α,β-unsaturated ketone E/Z-10 upon 1,5-proton shift. Next, the catalyst-bonded dienolate will react with the newly formed Michael acceptors. The α-Re-face of β,γ-unsaturated 2-acyl imidazoles E-1a is strongly shielded by the nearby aryl ring of the ligand. Therefore, the dienolate prefers to attack E/Z-10 from its α-Si-face (T2). Finally, the desired product 2a dissociates after a protonation of the intermediate T3, and the catalyst is regenerated to accomplish one catalytic cycle.

**Discussion**

In summary, we have disclosed the diverse transformation of β,γ-unsaturated 2-acyl imidazoles in the presence of chiral Lewis acid catalysts, involving catalytic asymmetric tandem isomerization/Michael addition, sulfur-Michael addition, and direct Mannich reaction. A wide range of chiral 1,5-dicarbonyl and functionalized carbonyl compounds was afforded with good to excellent levels yields, diastereoselectivities, and enantioselectivities.

![Mechanistic studies](https://doi.org/10.1038/s41467-020-17681-9)
into useful compounds with good results under mild conditions. Further studies on this methodology are ongoing.

**Methods**

**Tandem isomerization/α-Michael addition.** Y(OTf)₃ (0.005 mmol), L₃-RaPr₂ (0.005 mmol), β,γ-unsaturated 2-acyl imidazole E-1a (0.2 mmol), and NEt₃ (0.02 mmol) were dissolved in 1.0 mL of toluene under N₂ atmosphere. The mixture was stirred at 25 °C for 12 h and subjected to column chromatography on silica to afford the product 2a (Pet/EtOAc = 1:1 as eluent) as a colorless foam.

**Mannich reaction with isatin-derived ketimines.** A dry reaction tube was charged with L₃-PitBu (2.2 mg, 5 mol%), La(OTf)₃ (2.9 mg, 5 mol%), 3 Å M.S. (30 mg), and E-1a (27.1 mg, 0.12 mmol) in CH₂ClCHCl₂ (1.0 mL). The mixture was stirred at 30 °C for 30 min, and then 3a (0.10 mmol, 26.0 mg) was added at 0 °C. After 3a was consumed (detected by thin-layer chromatography (TLC)), the residue was purified by column chromatography on silica gel to afford the product 4a (Pet/EtOAc = 1:1 as eluent) as a colorless foam.

**Mannich reaction with pyrazolinone-derived ketimines.** A dry reaction tube was charged with L₃-RaPr₂ (3.5 mg, 5 mol%), La(OTf)₃ (2.9 mg, 5 mol%), E-1a (24.9 mg, 0.11 mmol), and pyrazolinone-derived ketimine (34.9 mg, 0.10 mmol) in CHCl₃ (1.0 mL). After ketimine was consumed (detected by TLC), the residue was purified by column chromatography on silica gel to afford the product 4i (Pet/EtOAc = 2:1 as eluent) as a colorless foam.

**Mannich reaction with aldimines.** A dry reaction tube was charged with L₃-RaPr₂ (7.0 mg, 10 mol%), La(OTf)₃ (5.9 mg, 10 mol%), E-1a (24.9 mg, 0.10 mmol), 4 Å M.S. (20 mg), and benzaldehyde-derived aldimine (30.8 mg, 0.15 mmol) in CH₂ClCHCl₂ (1.0 mL). After E-1a was consumed (detected by TLC), the residue was purified by column chromatography on silica gel to afford the product 4r (Pet/EtOAc = 2:1 as eluent) as a colorless oil.

**Isomerization/sulfur-Michael reaction.** A dry reaction tube was charged with L₃-PePr₃ (4.2 mg, 5 mol%), Dy(OTf)₃ (3.0 mg, 5 mol%), and Z-1a (56.5 mg, 0.25 mmol) in CH₂ClCHCl₂ (1.0 mL). PhSH (0.10 mmol) was added and the mixture was stirred at 25 °C for 17 h. After PhSH was consumed (detected by TLC), the residue was purified by column chromatography on silica gel to afford the product 6a (Pet/EtOAc = 3:1 as eluent) as a pale yellow oil.

**Data availability**

The X-ray crystallographic coordinates for structures reported in this study have been deposited at the Cambridge Crystallographic Data Centre (CCDC), under deposition numbers CCDC 1972987 (2j), 2001513 (4r), and 1972937 (11). These data can be...
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**Author contributions**

T.K. performed experiments and prepared the Supplementary Information and paper. L.H. took part in the reaction development and synthesized several substrates. S.R. repeated some experiments. W.C. and X.L. helped with modifying the paper and Supplementary Information. X.F. conceived and directed the project.

**Competing interests**

The authors declare no competing interests.

**Additional information**

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