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A C₁-symmetric *N*-heterocyclic carbene catalysed oxidative spiroannulation of isatin-derived enals: highly enantioselective synthesis of spirooxindole δ -lactones†

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A C₁-symmetric *N*-heterocyclic carbene (NHC)-catalysed activation of isatin-derived enals under oxidative conditions was achieved. The *in situ* generated α,β -unsaturated acyl azolium species was efficiently trapped by 1,3-dicarbonyl compounds *via* a Michael addition/spiroannulation cascade, delivering a series of synthetically important spirooxindole δ -lactones with up to 96% enantioselectivity.

Spirooxindoles are present in a wide variety of natural products and biologically active molecules.¹ Polycyclic spirooxindole scaffold with multiple stereocenters are particularly intriguing targets in organic synthesis owing to their structural complexity and potential pharmaceutical value.² For example, the spirooxindole δ -lactone constitutes the backbone of many natural products, such as trigolutes A–C (Fig. 1a).³ However, the catalytic asymmetric synthesis of this privileged scaffold in a highly stereocontrolled fashion is still a challenging task, and very limited protocols were established to this end.^{3c,4} In this regard, the development of new, practical and efficient strategy for the highly enantioselective synthesis of spirooxindole δ -lactone scaffold is still in great demand.

Over the past decades, the ability of chiral *N*-heterocyclic carbenes (NHCs)⁵ toward activation of aldehyde and activated carboxylic acid derivatives through formation of transient acyl anion,⁶ enolate,⁷ homoenolate⁸ and acyl azolium⁹ intermediates was well-investigated for enantioselective C–C, C–O and C–N bond formations. Within the context, the oxidative NHC catalysis strategy enabled generation of acyl azolium from aldehydes *via in situ* oxidation, which provided new opportunities for reaction design and selectivity control, thus remarkably extending the scope of NHC catalysis.^{10,11} However, despite fruitful achievements in the oxidative annulation of simple unsaturated aldehydes,¹² oxidative NHC catalysed addition/cyclization of β,β -disubstituted enals, such as isatin-derived enals, was a very attractive yet challenging topic with the

concomitant formation of a congested quaternary center.¹³ Recently, we disclosed that C₁-symmetric biaryl-saturated imidazolium catalyst was a superior NHC for asymmetric oxidative annulation of α -aryl-substituted α,β -unsaturated aldehydes.¹⁴ With our ongoing interest in the development of practical methods toward asymmetric synthesis of chiral spirooxindole scaffolds,^{4b,15} we attempted to explore the catalytic performance of C₁-symmetric NHC in oxidative activation of isatin-derived enals.¹⁶ The *in situ* generated isatin-derived α,β -unsaturated acyl azolium could be trapped by readily available 1,3-dicarbonyl compounds *via* a spiro-quaternary carbon-forming Michael addition/cyclization process, eventually delivering synthetically intriguing chiral spirooxindole δ -lactones (Fig. 1b).¹⁷

Our investigation started with the reaction of isatin-derived α,β -unsaturated aldehyde **1a** (a mixture of *E* and *Z* isomer)

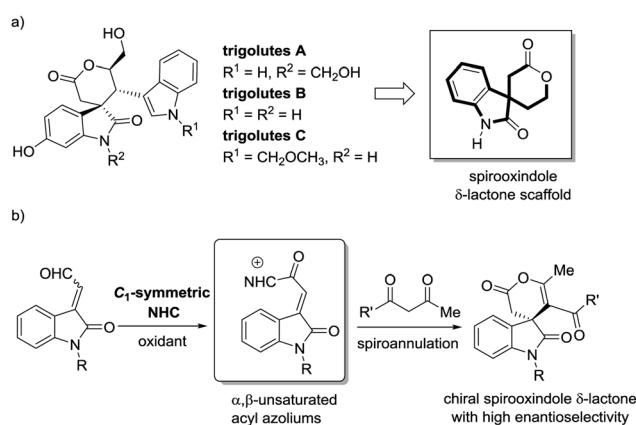


Fig. 1 (a) Examples of spirooxindole-containing natural products. (b) Assembly of spirooxindole δ -lactone scaffolds *via* C₁-symmetric NHC-catalysed spiroannulation (this work).

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with 1,3-dicarbonyl compound **2a** under the catalysis of C_1 -symmetric diaryl-saturated imidazolium catalysts **I–III**.¹⁸ To our delight, the desired oxidative annulation product **3a** and its regioisomer **3a'** (6 : 1 rr) were obtained in 23% yield and with high enantioselectivity (89% ee) when catalyst **I**/DBU combination were employed under the oxidation of 3,3',5,5'-tetra-*tert*-butyldiphenylquinone in THF (Table 1, entry 1), while catalyst **II** and **III** gave a relatively higher stereoselectivity and regioselectivity (Table 1, entries 2 and 3). As a comparison, the catalytic activity of C_2 -symmetric diaryl-saturated catalyst **IV**, triazolium catalysts **V** and **VI** was further examined, and inferior results with respect to reactivity and selectivity were observed (Table 1, entries 4–6). Then different bases including inorganic and organic bases were varied, and we found that DBU give better

stereoselectivity and regioselectivity within a shorter time (Table 1, entries 7–12). Solvent screening revealed that toluene gave rise to a better yield (Table 1, entries 13–17). When the catalyst loading was reduced to 10%, the reaction yield was obviously decreased (Table 1, entry 18). For further investigation, we modulated the ratio of **1** and **2** to 1 : 1, and further improvement in yield was observed (Table 1, entry 19).

With the optimized reaction conditions in hand, the generality of the reaction was explored (Table 2, entries 1–20). A broad range of differently substituted 1,3-dione bearing electron-donating or electron-withdrawing substituents on *meta*- or *para*-position of aromatic rings were well tolerated, whereas the 1,3-dione with substituents on *ortho*-position of aromatic rings gave complex products as a result of steric hindrance. Although no pronounced electronic effect was observed, we discovered that substrates with the nitro group (**2e**, **2n**) gave lower yields and enantioselectivities (Table 2, entries 5 and 14), while substrates with the methyl group (**2g**, **2j**) gave lower regioselectivities but higher yields (Table 2, entries 7, 8 and 10). Heteroaryl-containing substrate (**2o**) also worked well to give the δ -lactone **3o** with excellent regioselectivity in moderate yield (52%) and high ee (83%) (Table 2, entry 15). Aliphatic 1,3-dione **2p** could participate in the oxidative reaction efficiently, giving

Table 1 Optimization of reaction conditions

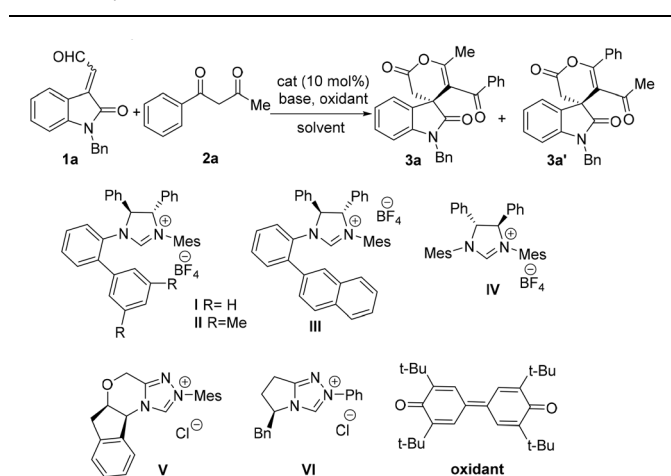


Table 2 Scope of the reaction

Reaction scheme for Table 2: 1 (1,3-dicarbonyl compound) reacts with 2 (1,3-dione) in the presence of catalyst **II** (20 mol%), DBU (20 mol%), and oxidant (200 mol%) in toluene at room temperature to yield product 3.

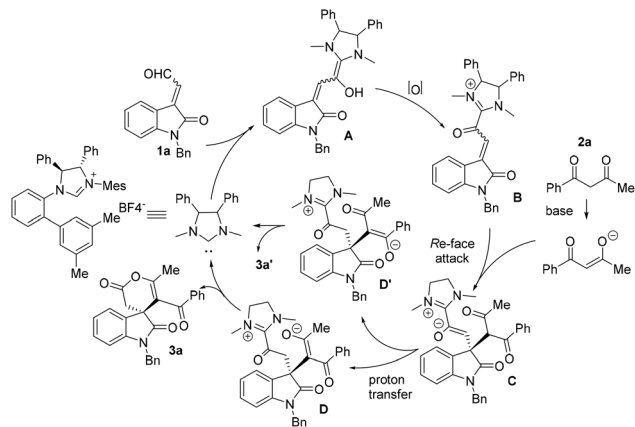
Legend: **1a**, R=Bn; **1b**, R=Me; **1c**, R=allyl; **1d**, R=*n*-Bu.

Entry ^a	R ¹ /R ² (2)	3	Yield ^b [%]	Rr ^c	Ee ^d [%]
1	Ph/Me (2a)	3a	77	12 : 1	94
2	4-BrC ₆ H ₄ /Me (2b)	3b	75	14 : 1	92
3	4-FC ₆ H ₄ /Me (2c)	3c	78	14 : 1	92
4	4-ClC ₆ H ₄ /Me (2d)	3d	69	17 : 1	93
5	4-NO ₂ C ₆ H ₄ /Me (2e)	3e	50	18 : 1	84
6	4-CF ₃ C ₆ H ₄ /Me (2f)	3f	65	14 : 1	90
7	4-MeC ₆ H ₄ /Me (2g)	3g	82	7 : 1	95
8	4- <i>t</i> -BuC ₆ H ₄ /Me (2h)	3h	74	5 : 1	94
9	3-MeOC ₆ H ₄ /Me (2i)	3i	79	12 : 1	94
10	3-MeC ₆ H ₄ /Me (2j)	3j	84	6 : 1	94
11	3-BrC ₆ H ₄ /Me (2k)	3k	69	16 : 1	92
12	3-ClC ₆ H ₄ /Me (2l)	3l	72	16 : 1	92
13	3-FC ₆ H ₄ /Me (2m)	3m	76	17 : 1	93
14	3-NO ₂ C ₆ H ₄ /Me (2n)	3n	45	19 : 1	83
15	2-Thienyl/Me (2o)	3o	52	>25 : 1	83
16	Me/Me (2p)	3p	58	—	90
17 ^e	Ph/Me (2a)	3q	62	>20 : 1	94
18 ^f	Ph/Me (2a)	3r	74	12 : 1	94
19 ^g	Ph/Me (2a)	3s	52	8 : 1	94
20	Ph/Ph (2q)	3t	60	—	96

^a Unless otherwise mentioned, all reactions were performed using **1a** (0.1 mmol) and **2a** (0.2 mmol) in solvent (1.0 mL) in the presence of catalyst (20 mol%), base (20 mol%), and oxidant (0.2 mmol) at room temperature. ^b Combined yields of **3a** and **3a'**. ^c Rr refers to ratio of **3a** and **3a'**, which is determined by ¹H NMR analysis. ^d Determined by HPLC analysis on a chiral column. ^e 10 mol% catalyst was used. ^f 0.1 mmol **1a** and 0.1 mmol **2a** were used.

^a all reactions were performed using **1a** (0.1 mmol) and **2** (0.1 mmol) in toluene (1.0 mL) in the presence of catalyst **B** (20 mol%), DBU (20 mol%), and oxidant (0.2 mmol) at room temperature. ^b Combined yields of **3** and **3'**. ^c Determined by ¹H NMR analysis. ^d Determined of the major isomer by HPLC analysis on a chiral column. ^e **1b** was used. ^f **1c** was used. ^g **1d** was used.





Scheme 1 Proposed catalytic cycle.

rise to the product in 58% yield and 90% ee within 10 minutes (Table 2, entry 16), while biaryl-substituted substrate **2q** give the corresponding product **3t** in 60% yield and 96% ee over 24 h (Table 2, entry 20). Further investigation revealed that this method was compatible with different *N*-substituted enals, affording the desired δ -lactone **3q–3s** in moderate yield with good regioselectivities and excellent enantioselectivities (Table 2, entries 17–19).

The absolute configuration of the product **3a** was assigned to be *S* by X-ray crystallographic analysis (Cu target, see ESI†).¹⁹ Based on our experimental results and previous literature reports, a proposed catalytic cycle is shown in Scheme 1. The free NHC catalyst, derived from precatalyst by *in situ* deprotonation, could react with isatin-derived unsaturated aldehyde **1a**, giving rise to Breslow intermediate **A**, which could be converted to acyl azolium intermediate **B** through two-electron oxidation and deprotonation. Subsequent Re-face Michael attack of **2a**-derived enolate to species **B** along with the proton transfer sequentially produced intermediates **C** and **D**. Intramolecular cyclization of **D** gave the product **3a** and released the NHC catalyst to next catalytic cycle. Mechanistically, the formation of byproduct **3a'** arose from intramolecular annulation of intermediate **D'**, which was generated from Michael adduct **C** via undesired competing proton transfer step.²⁰

Conclusions

In conclusion, we have developed a C_1 -symmetric NHC catalyzed oxidative spiroannulation of isatin-derived enals. The *in situ* generated isatin-derived α,β -unsaturated acyl azolium species was efficiently trapped by 1,3-dicarbonyl compounds via a Michael addition/cyclization process. A series of synthetically intriguing spirooxindole δ -lactones was readily obtained in moderate to high yields with excellent enantioselectivities under mild conditions. Further application of the C_1 -symmetric NHCs to novel and challenging transformations toward synthesis of synthetically or pharmaceutically valuable molecules are currently underway.

Conflicts of interest

There are no conflicts to declare.

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