

Cite this: *Chem. Sci.*, 2023, 14, 6663

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Cu(II)-mediated direct intramolecular cyclopropanation of distal olefinic acetate: access to cyclopropane-fused γ -lactones†

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Cyclopropane-fused ring scaffolds represent one of the most appealing structural motifs in organic chemistry due to their wide presence in bioactive molecules and versatility in organic synthesis. These skeletons are typically prepared from olefinic diazo compounds *via* transition-metal catalysed intramolecular carbenoid insertion, which suffers from prefunctionalization of starting materials and limited substrate scope. Herein, we disclose a practical copper-mediated direct intramolecular cyclopropanation of distal olefinic acetate to synthesize cyclopropane-fused γ -lactones and lactams. This cascade reaction is postulated to proceed through a hydrogen atom transfer event induced radical cyclization and copper-mediated cyclopropanation sequence. The protocol features high atom- and step-economy, excellent diastereoselectivity, broad tolerance of functional groups, and operational simplicity.

Received 5th April 2023

Accepted 24th May 2023

DOI: 10.1039/d3sc01752d

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Introduction

Cyclopropane represents one of the most appealing structural units with distinctive properties, containing three coplanar but highly strained π -character C–C bonds and stronger C–H bonds than common alkanes.¹ These features made cyclopropane not only a highly attractive building block in synthetic chemistry,² but also a valuable structure motif in medicinal chemistry. In particular, cyclopropane-fused γ -lactone and lactam units are prevalent in various bioactive naturally occurring products and drug molecules, which constitute key pharmacophores³ (Fig. 1).

Therefore, the chemical synthesis of cyclopropane-fused ring scaffolds has evoked considerable interest, and extensive efforts have been made in this field over the past few decades.⁴ Among the reported synthetic procedures, cyclopropanation of olefins with diazo compounds *via* metal-carbenoid intermediates represents one of the most frequently investigated methods. Robust variants of intramolecular cyclopropanation of olefins have been realized using allylic diazoacetates or diazoamides with transition-metal catalysts such as Rh, Ru, Co, Fe and Cu (Scheme 1a).⁵ Although they are efficient and provide a handle for accessing cyclopropane derivatives, the prefunctionalization

required to install the diazo moiety and the potential hazardous properties of diazo compounds limit their further application. In contrast, direct intramolecular dehydrogenative annulation of distal alkenes represents an ideal approach from the atom- and step-economy point of view.⁶ Several elegant examples of this intramolecular oxidative cyclopropanation of alkenes were reported in the 1980s.⁷ However, stoichiometric amounts of Mn(OAc)₃ or iodine were required for these transformations. Very recently, Xu and co-workers achieved such a transformation with an organo-electrocatalysis protocol, which provided convenient and efficient access to several types of cyclopropane-fused rings (Scheme 1b).⁸ In addition, reactions of ylides with electron-deficient alkenes⁹ and intermolecular cyclopropanation of allylic alcohols with aryldiazoacetates¹⁰ were also developed to build cyclopropane-fused skeletons. Despite these elegant

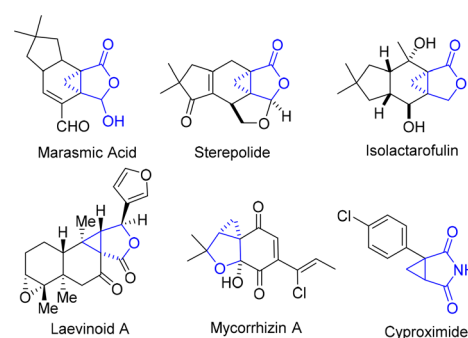
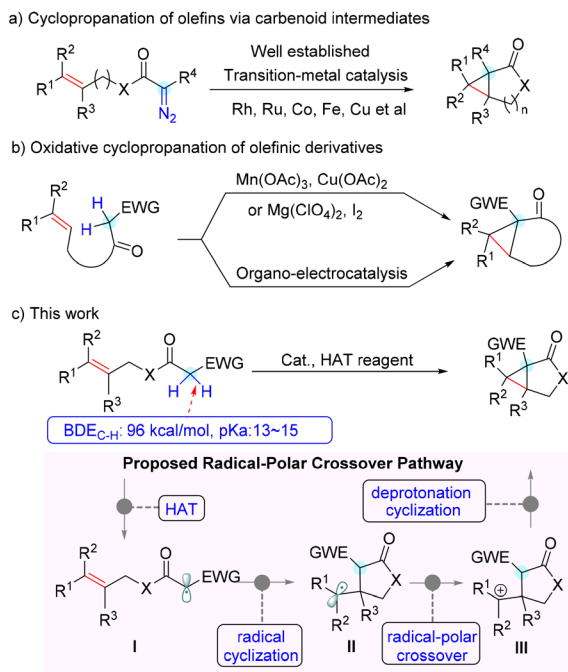


Fig. 1 Selected naturally occurring cyclopropane-fused γ -lactones and lactams.

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† Electronic supplementary information (ESI) available: CCDC 2254091. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3sc01752d>

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Scheme 1 Examples of intramolecular cyclopropanation and our method.

achievements, it is still of great significance to develop concise and versatile methodologies to construct cyclopropane-fused rings from readily available starting substrates in a step-, atom-, and cost-economical fashion.

Recently, we have been interested in radical chemistry¹¹ and heterocycle synthesis.¹² We envisioned that the relatively weak bond dissociation energy (~ 96 kcal mol⁻¹) of the specific α -C-H bond of allyl acetate activated by an electron-withdrawing group would be abstracted by a suitable hydrogen atom transfer (HAT) reagent to generate the active radical intermediate (Scheme 1c, I). On the other hand, the rather acidic proton in the substrate would be easily deprotonated to engage in a nucleophilic addition. Thus, we hypothesized that a radical cyclization followed by a nucleophilic annulation *via* a radical-polar crossover process would be highly feasible to construct cyclopropane-fused rings (Scheme 1c). As the outcome of this effort, we report herein an efficient pathway for direct intramolecular alkene cyclopropanation *via* a Cu(II)-mediated cascade process, which provides rapid and scalable access to cyclopropane-fused γ -lactones and -lactams.

Given the rather high O-H bond dissociation energy (BDE_{O-H} = 105 kcal mol⁻¹) in alcohol as well as its ready availability, an alkoxy radical derived from peroxide was selected as the HAT reagent to initiate the reaction. We probed the hypothesis by treating cinnamyl 2-(phenylsulfonyl)acetate **1aa** with di-*tert*-butyl peroxide (DTBP) in the presence of CuBr. Gratifyingly, the expected cascade cyclopropanation reaction readily occurred to afford cyclopropane-fused lactone **2aa** in a 64% isolated yield with high diastereoselectivity (Table 1, entry 1). Further optimization of the metal catalysts revealed that CuBr₂ is superior to others (for details, see the ESI†), and the catalyst loading had

Table 1 Optimization of the reaction conditions^a

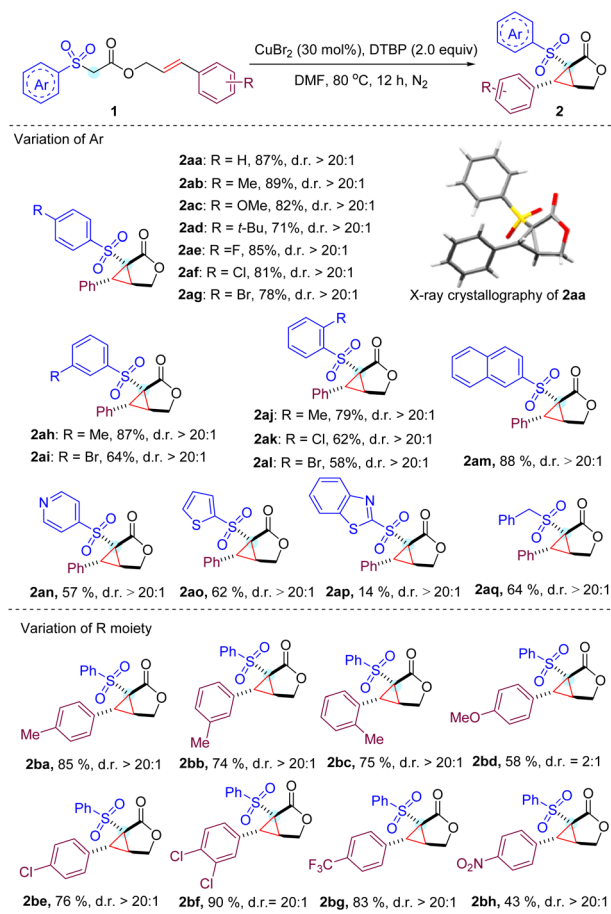
Entry	Variation of reaction conditions	Yield of 2aa ^b (%)
1 ^c	CuBr instead of CuBr ₂	64
2 ^c	CuI instead of CuBr ₂	71
3	None	87
4	20 mol% of CuBr ₂	61
5	10 mol% of CuBr ₂	22
6 ^c	TBPP or LPO instead of DTBP	Trace
7 ^c	DCP instead of DTBP	80
8 ^c	Conducting at 40 °C	N.R.
9 ^c	Conducting at 60 °C	81
10 ^c	CH ₃ CN as the solvent	83
11 ^c	DMSO as the solvent	58
12 ^c	THF, DCE or toluene	N.D.
13	Conducting in air	Trace

^a Reaction conditions: **1aa** (0.2 mmol, 1.0 equiv.), CuBr₂ (30 mol%) and DTBP (0.4 mmol, 2.0 equiv.) in DMF (1 mL) stirred under N₂ at 80 °C unless otherwise specified. ^b Isolated yield. ^c 40 mol% CuBr₂ was employed.

a distinct influence on the outcome of this reaction. 30 mol% of CuBr₂ is required to achieve a high 87% yield, while decreasing the loading of CuBr₂ resulted in a dramatic drop in the yield (Table 1, entries 3–5). Other peroxides, including *tert*-butyl peroxybenzoate (TBPP), lauroyl peroxide (LPO), and dicumyl peroxide (DCP), were also surveyed, but none of them gave better results than DTBP (Table 1, entries 6 and 7). Heating was also important to this domino reaction, as conducting the reaction at a lower temperature led to a lower yield because of reduced substrate conversion (Table 1, entries 8 and 9). Presumably, heating benefits the conversion through several pathways, including facilitating the homolysis of peroxide, increasing mass transfer, and enhancing catalyst turnover.¹² Further studies showed that polar solvents (CH₃CN and DMSO) were tolerated well in the reaction, while less polar THF, DCE, or toluene were not compatible (Table 1, entries 10–12). Moreover, when the reaction was conducted in air, only trace **2aa** was observed (Table 1, entry 13). It is worth mentioning that neither the Mn(OAc)₃ mediated oxidative annulation^{7a} nor the halocyclization protocol^{7b} could efficiently produce the expected cyclopropane-fused γ -lactone **2aa** from phenylsulfonylacetate **1aa**, which highlighted the unique efficiency of our copper-catalysed reaction system.

With the optimized conditions in hand, we next studied a series of substituted cinnamyl acetates with different aryl sulfonyl groups to explore the reaction scope (Table 2). To our delight, all reactions of cinnamyl acetate with α -aryl sulfonyl groups bearing electron-donating and -withdrawing groups on the *para* and *meta* positions proceed smoothly and afford the expected products (**2aa–2ai**) in high yields with excellent diastereoselectivity (d.r. > 20 : 1).¹³ Halogens were well tolerated in the substrates, which enabled the possible further derivatization



Table 2 Substrate scope investigation^a

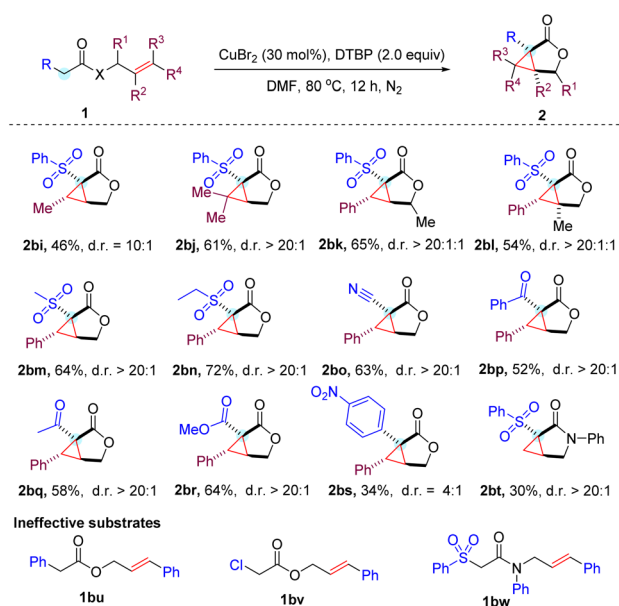
^a Reaction conditions: **1** (0.2 mmol, 1.0 equiv.), CuBr₂ (30 mol%) and DTBP (0.4 mmol, 2 equiv.) in DMF (1 mL) stirred under N₂ at 80 °C and isolated yields are given. d.r. was determined by crude ¹HNMR spectroscopy, for details see the ESI.

of the cyclopropane-fused γ -lactone by classical cross coupling methods. Overall, a minor effect on the reaction yield of *ortho*-substituted phenyl sulfonyl acetate was observed, and the expected products (**2aj–2al**) were obtained in slightly reduced yields, which might result from the steric hindrance of the substrates. In addition, we noticed that the reaction worked well by replacing the phenyl ring with a 2-naphthyl ring in the substrate to deliver product **2am** in 88% yield under the standard conditions. It is noteworthy that heteroaryl bearing substituents such as pyridine (**2an**), thiophene (**2ao**) and benzothiazole (**2ap**) were also viable substrates, albeit with lower yields.

Then, the substituent effect on the cinnamyl moiety of the substrate was investigated. Methyl groups at the *para*, *meta*, and *ortho* positions did not significantly affect the reaction efficiency. In contrast, a strong electron-donating methoxy group at the *para* position of the cinnamyl phenyl ring lowered the yield as well as the diastereoselectivity (**2bd**). It is assumed that rather than participating in the cascade cyclization cycle, the relatively electron-rich benzylic radical involved in this transformation is more likely to be over oxidized to produce a number of unknown

byproducts.¹⁴ Gratifyingly, 4-chlorophenyl, 3,4-dichlorophenyl and 4-trifluorophenyl substituents proved compatible with current reaction conditions, resulting in the corresponding products (**2be–2bg**) in good yields. Notably, the strong electron-withdrawing nitro group was also tolerated to provide product **2bh** at 43% yield.

In addition to cinnamyl acetates, the CuBr₂ mediated cyclopropanation was proven to be effective for allyl acetates (Table 3). However, the corresponding cyclopropane-fused γ -lactones (**2bi–2bl**) were generated in relatively lower yields. This is presumably due to the lower stability of the generated alkyl radical compared to the benzyl radical derived from the cinnamyl substrates. Notably, the steric effect in the allyl moiety likely played an important role in the diastereoselectivity of the reaction, as product **2bi** with one methyl group was obtained in reduced 10 : 1 d.r., while both the phenyl and dimethyl substituted products (**2aa** and **2bj**) were all obtained with excellent >20 : 1 diastereoselectivities. Alkylsulfonyl acetates were also capable of producing the corresponding cyclopropane-fused γ -lactones **2bm** and **2bn** with a moderate yield and good diastereoselectivities, demonstrating that the electron-withdrawing group (EWG) in the substituted acetates is not restricted to the arylsulfone unit. Additionally, the cyano, ketone, and ester groups were all compatible with the reaction conditions, resulting in the expected products (**2bo–2br**) being produced in moderate yields with good diastereoselectivities. It is worth mentioning that the electron-deficient aromatic system (4-nitrophenyl) was also tolerated to give the desired product in a synthetically useful 34% yield. However, phenyl or chloro substituted acetates (**1bu** and **1bv**) were ineffective substrates under the standard conditions. This indicated that an appropriate second electron-withdrawing group

Table 3 Substrate scope investigation^a

^a Reaction conditions: **1** (0.2 mmol, 1.0 equiv.), CuBr₂ (30 mol%) and DTBP (0.4 mmol, 2 equiv.) in DMF (1 mL) stirred under N₂ at 80 °C and isolated yields are given. d.r. was determined by crude ¹HNMR spectroscopy, for details see the ESI.

is required for this copper-catalyzed cyclopropanation reaction. Presumably, it plays an important role in activating the protons of the substrates for deprotonation to engage in the cyclization cascade. We also succeeded in converting allyl acetamide into a cyclopropane-fused γ -lactam (**2bt**), albeit with a poor 30% yield. Unfortunately, the efficiency of current reaction conditions to prepare lactams is limited to terminal alkenes. Internal alkene **1bw** only gave trace amounts of the expected products.

To illustrate the potential practicability of this methodology, a gram-scale (4.5 mmol) synthesis of cyclopropane-fused γ -lactone **2aa** was conducted, and 81% yield (1.14 g) of the expected product was isolated with good diastereoselectivity (Fig. 2a). Remarkably, the cyclopropane-fused γ -lactone scaffold could serve as a versatile handle for further synthetic transformations to provide a wide variety of highly functionalized cyclic products. Reduction of **2aa** with LiAlH_4 afforded diol **3** with a 72% yield. Aminolysis of lactone **2aa** with aniline easily generated the highly substituted cyclopropane **4** as a single diastereomer in a good 89% yield. Treatment of **2aa** with EtMgBr in the presence of CuI provided hemiketal **5** in a moderate 53% yield. On the other hand, the strained cyclopropane ring can be deconstructed with Et_3SiH under Lewis acid conditions, delivering the γ -lactone **6** in 59% yield (Fig. 2b).

To gain insight into the reaction mechanism, a series of control experiments were carried out (Fig. 3). First, the reaction completely shut down in the absence of either the peroxide (DTBP) or CuBr_2 , which indicated that both were crucial to this transformation (Fig. 3A). When a commonly used radical inhibitor, such as BHT or TEMPO, was introduced into the reaction, the cyclopropanation process was completely suppressed. Moreover, the TEMPO adduct **7** was isolated in 22% yield. These results indicated that a radical process might be involved in this transformation (Fig. 3B and C). To prove the involvement of the radical-polar crossover process in the cyclopropanation reaction, we next turned to capture the possible carbocation species under the standard conditions with various nucleophiles such as MeOH, H_2O , TMSCN, aniline, and indole. Unfortunately, those attempts were unsuccessful, indicating that carbocation species were not likely involved (Fig. 3D). These results led us to speculate that the cyclopropane ring might be formed *via* another pathway rather

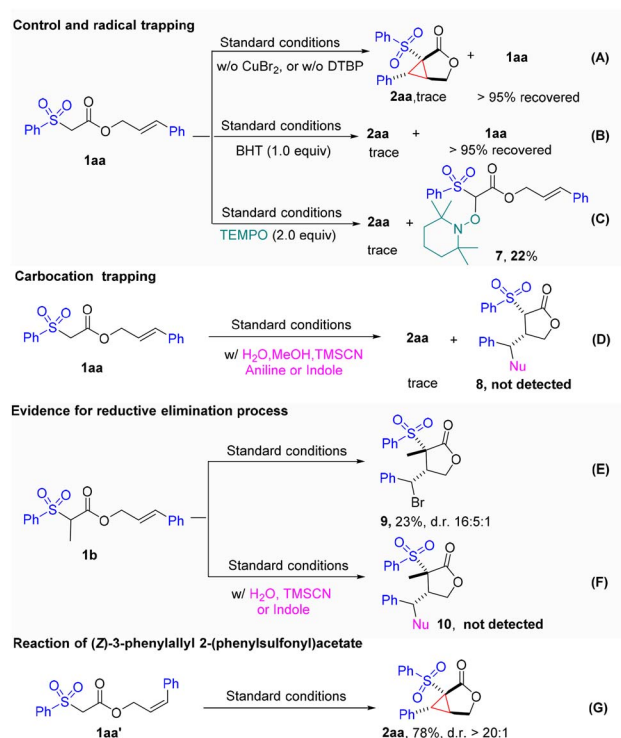


Fig. 3 Control experiments and mechanism investigation (A-G).

than the initially proposed radical-polar crossover mechanism. Intriguingly, we found that when cinnamate **1b** was subjected to the standard reaction conditions, the unexpected cyclized and bromo-substituted lactone **9** was obtained in a 23% yield (Fig. 3E). The addition of other nucleophiles led to suppression of the reaction, and no nucleophilic attached products formed (Fig. 3F). These results implied that the Br on lactone **9** might be introduced by reductive elimination from a hypervalent copper species. To determine whether the cyclopropanation process is stepwise or concerted, we examined the reaction of *cis*-allylic acetate **1aa'**. The identical product, **2aa**, was produced in a yield of 78% with excellent diastereoselectivity, suggesting that a stepwise mechanistic pathway may be at play (Fig. 3G).

On the basis of these results and the literature,¹⁵ a reaction mechanism was proposed, as depicted in Scheme 2. And it was further analyzed by density functional theory (DFT) calculations to probe the energetics of the individual reaction steps. Initially, hydrogen atom transfer (HAT) from **1aa** to the *tert*-butoxy radical generated from thermal homolysis of DTBP or Cu(i)-mediated SET reduction delivers the electrophilic radical **INT1** ($-3.9 \text{ kcal mol}^{-1}$). The transition state of the HAT step is located as **TS1**, which has an energy barrier of $18.8 \text{ kcal mol}^{-1}$. Intramolecular radical cyclization affords the benzyl radical **INT2** ($-24.4 \text{ kcal mol}^{-1}$), which reacts with Cu(II) species to produce the Cu(III) species **INT3** ($-49.2 \text{ kcal mol}^{-1}$). No transition state can be located suggesting that this cyclization process is concerted and spontaneous.¹⁶ The deprotonation of **INT3** induces ligand exchange, leading to the formation of bicyclic **INT4** ($-88.2 \text{ kcal mol}^{-1}$). The high exergonicity ($39.0 \text{ kcal mol}^{-1}$) of this step indicates that this process is very energetically favourable.

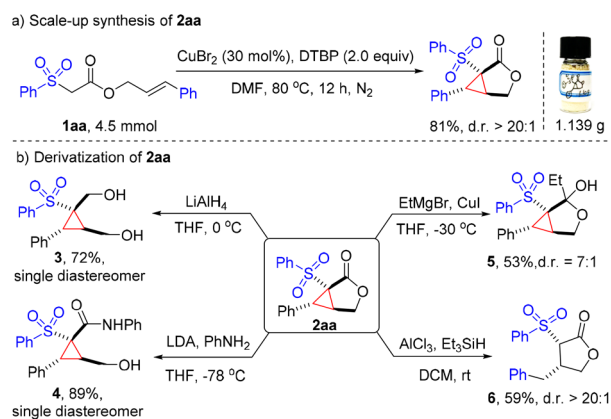
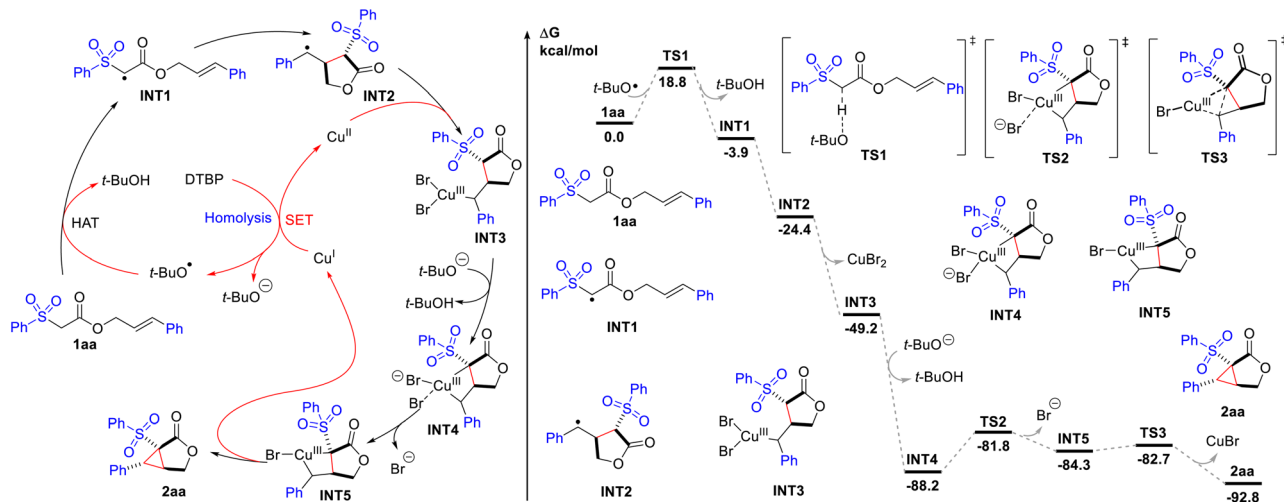


Fig. 2 Scale-up synthesis (a) and derivatization of **2aa** (b).





Scheme 2 Proposed mechanism and computed energy profile for the Cu-catalysed intramolecular cyclopropanation of **2aa**. Energies are in kcal mol⁻¹ and bond distances are labelled in Å.

Subsequently, bromide dissociates (**TS2**) with a barrier of 6.4 kcal mol⁻¹, forming **INT5** (−84.3 kcal mol⁻¹). Finally, reductive elimination of **INT5** gives the final cyclopropane-fused γ -lactone **2aa** (−92.8 kcal mol⁻¹), regenerating the reactive Cu(I) species. The transition state for this step is located as **TS3**, requiring only an energy barrier of 1.6 kcal mol⁻¹. For substrate **1b** which blocked the deprotonation and ligand exchange step (**INT3** to **INT4**), direct reductive elimination affords bromolactone **9**.

Conclusions

In conclusion, we have developed a novel copper-catalyzed protocol to access cyclopropane-fused γ -lactones and lactams with good yields and diastereoselectivities. The practical advantages of this method include the use of readily available starting materials, the avoidance of prefunctionalized precursors of carbenes, broad substrate compatibility, and high diastereoselectivity. Further control experiments and DFT calculations indicated that the HAT event-induced radical cyclization and copper-mediated cyclopropanation cascade processes are likely operational. Given the ubiquity and intermediacy of cyclopropane-fused ring scaffolds in bioactive natural products and drugs, this efficient and scalable method would be of high interest to synthesis communities.

Data availability

General information, detailed experimental procedures, characterization data for all new compounds, and NMR spectra are in the ESI.† Data for the crystal structure reported in this paper have been deposited at the Cambridge Crystallographic Data Centre (CCDC) under the deposition number CCDC 2254091.

Author contributions

L. Liu conceptualized the project. L. Liu and X.-H. Duan supervised the investigation. Y. Wang, C. He, Y. Zhou and K.

Zhang performed the research. S. Shen and Y. Su conducted the DFT calculations. B. Rao and T. Han analysed the X-ray structure of **2aa**. L. Liu and Y. Wang co-wrote the paper. All authors analysed the data, discussed the results and commented on the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

L. Liu thanks the National Natural Science Foundation of China (21901199) and Xi'an Jiaotong University (7121192002) for financial support. Y. Su acknowledges the “Young Talent Support Plan” of Xi'an Jiaotong University. Supercomputing facilities were provided by the Hefei Advanced Computing Center. We thank Prof. Li-Na Guo and Miss Kehui Wang at Xi'an Jiaotong University for valuable discussion. We thank Miss. Lu Bai and Miss. Chao Feng at the Instrument Analysis Center of Xi'an Jiaotong University for HRMS and NMR analysis.

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