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Silver-catalyzed decarboxylative radical cascade cyclization toward benzimidazo[2,1-*a*]isoquinolin-6(5*H*)-ones†

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A simple and efficient decarboxylative radical addition/cyclization strategy was developed, by which a wide range of benzimidazo[2,1-*a*]isoquinoline-6(5*H*)-ones were prepared in one-pot *via* reaction of functionalized 2-arylbenzimidazoles and carboxylic acids in the presence of $K_2S_2O_8/AgNO_3$ under mild reaction conditions.

Structurally diverse polycycles containing benzimidazo-isoquinoline fused framework **A**, as shown in Scheme 1, compose an important part of privileged synthetic intermediates, biologically active molecules and functional materials.¹ Scheme 1 shows some important polyheterocycles that contain scaffold **A**. Among them, polyheterocycles **B–E** exhibit a wide variety of pharmaceutical activities, such as modulation of potassium ion flux, antitumor, *etc.*² Compound **F** was reported to be a potential candidate for organic electronics.³ Nowadays, the development of efficient

synthetic strategies for quickly accessing biologically and materially valuable benzimidazole-containing polycycles has been enthusiastically pursued and expected in the organic synthetic field.⁴

Most traditional synthetic methods for the construction of the above-mentioned benzimidazo-isoquinoline fused polycycles (Scheme 1) heavily relied on condensation.⁵ However, almost all of those classical methods suffer limitations such as not easily available starting materials, harsh reaction conditions and practical inconvenience. Very recently, Song and co-workers developed an elegant [4+2] annulative reaction for the construction of benzimidazole[2,1-*a*]isoquinolines from 2-arylimidazoles and α -diazoketoesters in the presence of a $Cp^*Rh(III)$ catalyst (Scheme 2a).^{2b} However, this protocol involved the use of potentially explosive diazo compounds,⁶ and an expensive Rh catalyst, which might hinder its synthetic application. Over the past decade, the rapid development of radical cascade reactions for concise construction of structurally diverse polycycles has been witnessed.⁷ The remarkable merits of radical cascade reactions include atom/step economy, together with the reduction of work and time needed to carry them out.⁸ Over the past four decades, carboxylic acids, as commercially available and inexpensive compounds, have been widely used as alkyl radical precursors to construct C-heteroatom and C–C bonds *via* decarboxylation reactions.⁹ In particular, in the past ten years, many highly valuable decarboxylative cascade cyclization reactions have been successfully developed for accessing many different biologically



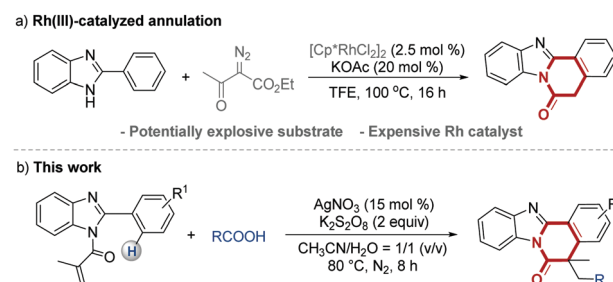
Scheme 1 Examples of benzimidazo-isoquinoline containing polyheterocycles.

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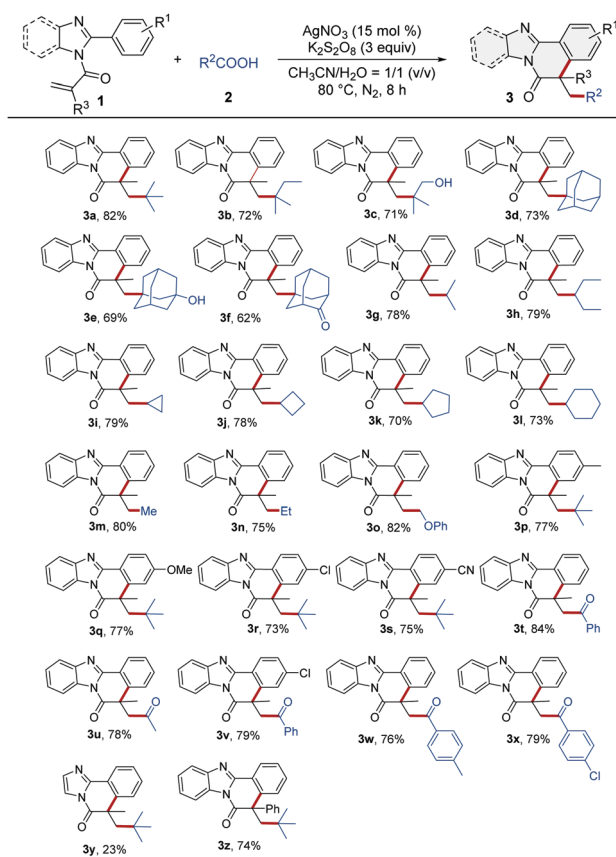
Scheme 2 Synthesis of benzimidazo-isoquinoline fused frameworks.

and materially important heterocycles.¹⁰ As part of our continuing interest in the development of novel alkyl-radical-involved reactions,¹¹ herein, we present an efficient silver-catalyzed decarboxylative cascade radical cyclization strategy, by which a large variety of benzimidazo[2,1-*a*]isoquinolins were prepared *via* reaction of functionalized 2-arylbenzimidazoles with carboxylic acids in the presence of AgNO₃/K₂S₂O₈ in one pot under mild reaction conditions, as illustrated in Scheme 2b. To the best of our knowledge, this is the first example of constructing a large variety of biologically and materially valuable benzimidazo[2,1-*a*]isoquinolins *via* a silver-catalyzed decarboxylative cascade radical cyclization.

We initiated our study by evaluating the model reaction of *N*-methacryloyl-2-phenylbenzimidazole (**1a**) with pivalic acid (**2a**) under different reaction conditions, as summarized in Table S1 (ESI[†]). After an intensive experimentation, the optimal reaction conditions were thus established as follows: **1a** (0.5 mmol), **2a** (1 mmol), AgNO₃ (15 mol%) and K₂S₂O₈ (3 equiv.) in the mixed solvent of CH₃CN/H₂O (v/v = 1/1) at 80 °C for 8 h under a N₂ atmosphere.

With the optimized reaction conditions established, we next explored the substrate scope of this Ag-catalyzed decarboxylative radical cyclization. As it can be seen in Table 1, many tertiary α -substituted aliphatic carboxylic acids reacted well with *N*-methacryloyl-2-phenylbenzimidazole (**1a**), giving the corresponding polycyclic products **3a–3f** in good yields (62–82%). It is especially worth mentioning here that the pharmaceutically attractive¹² adamantane motifs were successfully incorporated into the polycycles as meaningful functional groups of the benzimidazoisoquinolines (**3d–3f**). Many secondary α -substituted carboxylic acids, including isobutyric acid, 2-ethylbutanoic acid, and even 3-, 4-, 5-, and 6-membered ring containing carboxylic acids were also efficiently converted into the corresponding products (**3g–3l**). Notably, three primary carboxylic acids, acetic acid, propionic acid and phenoxy acetic acid which were predicted to produce three unstable primary alkyl radicals *via* a decarboxylation process, were surprisingly feasible for this synthetic procedure, delivering the corresponding polycycles in 75–82% yields (**3m–3o**). It was also worth noticing that a number of functionally unstable groups such as the hydroxy group, ether group, and in particular the cyclopropyl group and cyclobutyl group contained in the starting carboxylic acids were well tolerated in these procedures, affording the corresponding products **3c**, **3e**, **3i**, **3j**, and **3o** in good yields. Afterward, pivalic acid was used to react with various functionalized 2-arylbenzimidazoles bearing electron-donating and electron-withdrawing groups like –Me, –OMe, –Cl, and –CN on the 2-phenyl ring, giving the products **3p–3s** in good yields (73–77%), and no obvious electronic effects were observed in those cases (**3p–3s**). Encouraged by these results, we extended this decarboxylative addition/cyclization reaction to 2-oxopropanoic acid and 2-oxo-2-arylacetic acids. To our delight, the desired products were obtained in 76–79% yields (**3t–3x**). In addition, we carried out a reaction starting from *N*-methacryloyl-2-phenylimidazole and pivalic acid under the optimized reaction conditions, as illustrated in Table 1. However, we only get 23% yield of the corresponding product (**3y**). The low yield of **3y** might

Table 1 Substrate scope for benzimidazo[2,1-*a*]isoquinolin-6(5*H*)-ones^a

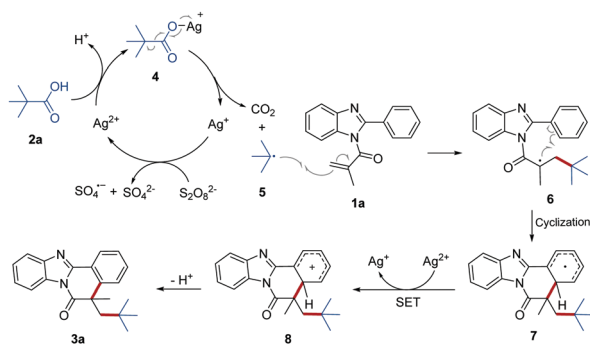


^a Reaction conditions: **1** (0.5 mmol), **2** (1 mmol), AgNO₃ (15 mol%), K₂S₂O₈ (3 equiv.), CH₃CN/H₂O = 1/1 (v/v) (5 mL), 80 °C, 8 h. Isolated yields were given.

be owing to the less favorable conformation of the related intermediate radical (for details, see the ESI[†]). Finally, we performed an experiment by reacting 2-phenyl-1-(2-phenylbenzimidazol-1-yl)prop-2-en-1-one (**1z**) with pivalic acid under the optimized reaction conditions. It was pleasing that 74% yield of **3z** was obtained, indicating that the phenyl group directly attached to the vinyl group in **1z**, basically didn't interfere with the cyclization process toward the formation of the final product **3z**.

To gain further insight into the reaction mechanism, control experiments were conducted (Scheme S1, ESI[†]). 4.0 equiv. of TEMPO (2,2,6,6-tetramethyl-1-piperidinyloxy) and 4.0 equiv. of BHT (2,6-di-*tert*-butyl-4-methylphenol) as radical scavengers were subjected to the reaction under standard conditions, respectively. As a result, the Ag-catalyzed decarboxylative cyclization reactions were remarkably suppressed, indicating that this cascade reaction may involve a radical process.

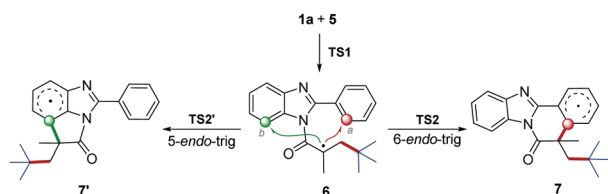
Based on the above experimental results and previous reports, we proposed a plausible mechanism of this silver-catalyzed transformation as described in Scheme 3. Initially, Ag(I) was oxidized by persulfate (S₂O₈²⁻) to generate the Ag(II) cation. Subsequently, the Ag(II) cation reacted with **2a** to afford the silver carboxylate complex **4** with loss of a proton. Complex **4** then was homolytically cleaved to form *t*-butyl radical **5** with release of a



Scheme 3 Plausible mechanism of the decarboxylative cyclization.

CO₂ molecule and regenerating the Ag(I) cation. Following that, the addition of reactive radical **5** to the C=C bond of **1a** delivered radical intermediate **6**, which subsequently underwent an intramolecular cyclization to form intermediate **7**. Intermediate **7** was further oxidized into the carbocation **8** by Ag(II) via an intermolecular single electron transfer (SET) process. Finally, rapid deprotonation of carbocation **8** regenerated the aromatic ring to give the final product **3**.

According to the proposed mechanism and previous reports,¹³ after the generation of radical intermediate **6**, both the 6-*endo*-trig cyclization at the a-position and 5-*endo*-trig cyclization at the b-position, leading to the polycycle **7** and **7'** respectively, would be possible (Scheme 4). However, no **7'** product was obtained from our synthesis. To understand this regioselectivity, we have investigated the competing pathways for the structural transformations from **1a** and **5** to the five- and six-membered ring intermediates **7** and **7'** by performing DFT calculations, which have been frequently carried out for the investigation of organic reaction mechanisms.¹⁴ As shown in Fig. 1, the two regioselective pathways would share the same mechanism for the addition of radical **5** to **1a** to form intermediate **6** via transition state **TS1** with an energy barrier of 8.6 kcal mol⁻¹, and then divorced. Starting from **6**, the one pathway underwent six-membered ring transition state **TS2** to generate **7**, and the other pathway went through five-membered ring transition state **TS2'** to form **7'**. Obviously, the energy barrier associated with **TS2** (16.6 kcal mol⁻¹) is much lower than that associated with **TS2'** (34.2 kcal mol⁻¹), indicating that the six-membered ring product should be the main one in kinetics. Compared with the initial angle C–N–C (124°) in **6**, it would be distorted much more in **TS2'** (110°) than that (127°) in **TS2**, which demonstrates that the less distortion should be responsible for the lower energy of **TS2**. In addition, the relative Gibbs free energy of **7** would be much lower than that of **7'**.



Scheme 4 The possible pathways of cyclization.



Fig. 1 The Gibbs free energy profile for the competing reactions (distance in Å).

Hence, the pathway associated with the six-membered product would be more energetically favourable in both kinetics and thermodynamics, which is in agreement with the experimental observation.

In conclusion, we have disclosed a simple and efficient decarboxylative radical addition/cyclization strategy to construct the benzimidazo[2,1-*a*]isoquinolines from easily available 2-arylbenzimidazoles and carboxylic acids using inexpensive silver catalysis. A large number of carboxylic acids, including tertiary α -substituted tertiary, secondary and even primary carboxylic acids, as well as 2-oxocarboxylic acids were all suitable for being used as the alkyl radical precursors to initiate this radical addition/cyclization reaction. The wide reactant scope of carboxylic acids of this synthesis thus gave a big chance to incorporate many meaningful functional groups into the benzimidazo-quinolines. For example, the pharmaceutically attractive adamantane motifs were successfully introduced from the starting adamantane carboxylic acids into the biological and meaningful polycycle scaffold. Notably, several vulnerable functional groups contained in the starting carboxylic acids, such as the hydroxy group, ether group, and in particular the cyclopropyl group and cyclobutyl group, were well tolerated in this procedure, generating the corresponding polycycles in good yields. DFT calculations proved that the 6-*endo*-trig pathway is more energetically favourable in both kinetics and thermodynamics than the 5-*endo*-trig pathway. To the best of our knowledge, this is the first example of constructing a large variety of biologically and materially valuable benzimidazo[2,1-*a*]isoquinolin-6(5*H*)-ones via a silver-catalyzed decarboxylative cascade radical cyclization. Great advantages of this strategy include wide reactant scope, mild reaction conditions, experimental simplicity and easy workup. Further application of this strategy is currently ongoing in our laboratory.

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Conflicts of interest

There are no conflicts to declare.

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