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# Access to 6-hydroxy indolizines and related imidazo[1,5-*a*]pyridines through the S<sub>N</sub>2 substitution/condensation/tautomerization cascade process†

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A simple and efficient cascade reaction was developed for the construction of hydroxy substituted indolizines from pyrrole-2-carbaldehydes and commercially available 4-halogenated acetoacetic esters. Their optical properties were also evaluated.

## Introduction

Indolizine, a biostere for indole, is commonly found in numerous natural products and pharmaceuticals. Indolizine derivatives exhibit diverse biological activities such as anti-HIV, anti-inflammatory, anti-tubercular, and anticancer activities.<sup>1–5</sup> They are also used in dyes and optical materials owing to their bright colors.<sup>6–11</sup> As a consequence, much effort has been devoted to their synthesis and functionalization, and thus many methods have been developed.<sup>12–14</sup> In addition to classical Scholtz or Tschichibabin reactions, a variety of straightforward and efficient methods have been reported in recent years<sup>15–20</sup> including 1,3-dipolar cycloaddition of pyridinium salts and intramolecular cyclization catalyzed by transition metals and intermolecular cyclization. Despite the efficiency of these methods, they suffer from the requirement of specific pre-organized substrates, necessity of expensive metal catalysts, multistep synthesis, and a lack of product diversity. Moreover, no method has been reported for the preparation of indolizines bearing a hydroxyl group.

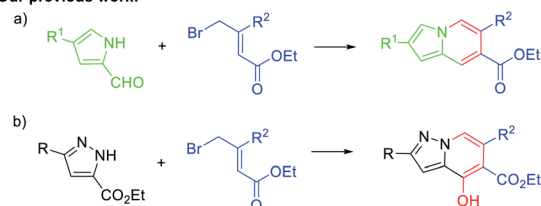
Recently, we synthesized a series of indolizine and related N-bridgehead heterocycles *via* a cascade reaction (Scheme 1a).<sup>21</sup> To achieve the related pyrazolo[1,5-*a*]pyridines through a shorter and convenient route, a simple and efficient synthetic method was also reported subsequently using commercially accessible starting materials (Scheme 1b).<sup>22</sup> Based on the results obtained in our laboratory, we expected that a cascade reaction

of pyrrole-2-carbaldehyde **1** with 4-halogenated acetoacetic ester **2** might be successful in the presence of a weak base (Scheme 1). In continuation of our effort to search for new fluorophores for imaging,<sup>23–26</sup> herein, we report a simple and efficient method for the synthesis of hydroxy substituted indolizines *via* an S<sub>N</sub>2 substitution/condensation/tautomerization cascade process in a metal-free fashion. Their optical properties were also evaluated.

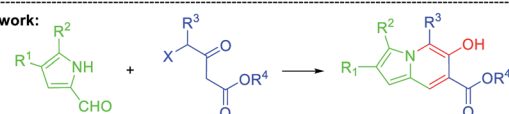
## Results and discussion

Initially, commercially available pyrrole-2-formaldehyde and ethyl 4-chloro-3-oxobutanoate **2a** were selected for the experimental design. However, only the dimerization product of **2a** was obtained. Subsequently, we optimized the reaction conditions using 4-propionyl-pyrrole-2-formaldehyde **1a** and ethyl 4-chloro-3-oxobutanoate **2a** as the model substrates. To our delight, the desired cyclized product **3a** was obtained in an acceptable 56% yield in the presence of K<sub>2</sub>CO<sub>3</sub> (Table 1, entry 1). The reaction even progressed very well at room temperature

Our previous work:



This work:



Scheme 1 Access to indolizines *via* a cascade reaction.

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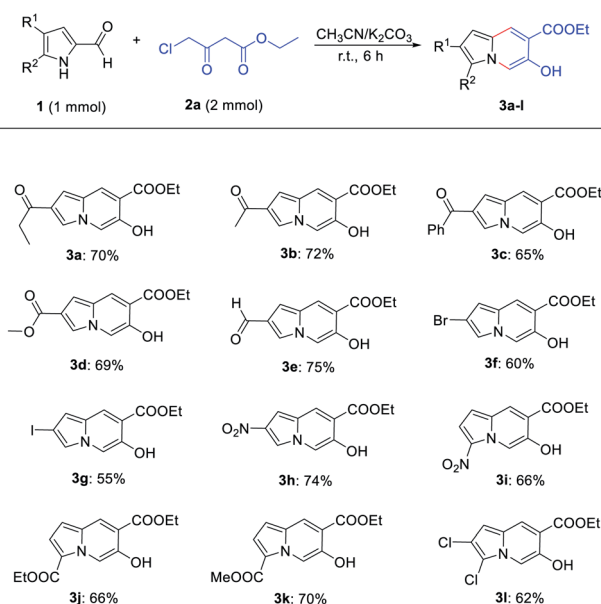
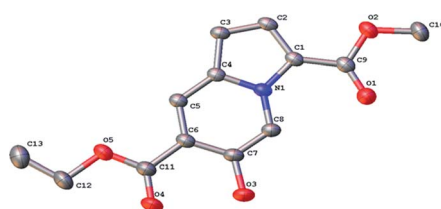
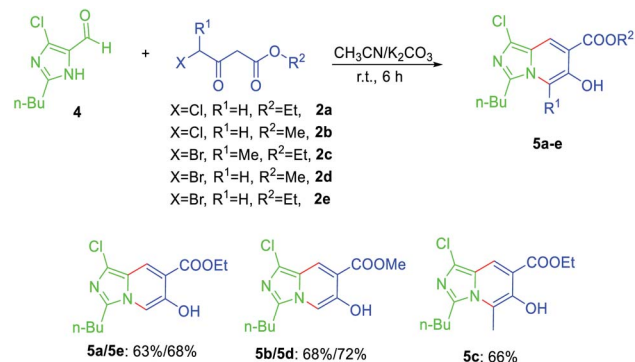
† Electronic supplementary information (ESI) available: <sup>1</sup>H NMR, <sup>13</sup>C NMR and HRMS spectra of compounds **3a–3l** and **5a–5e**. CCDC 2081693. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d1ra04425g

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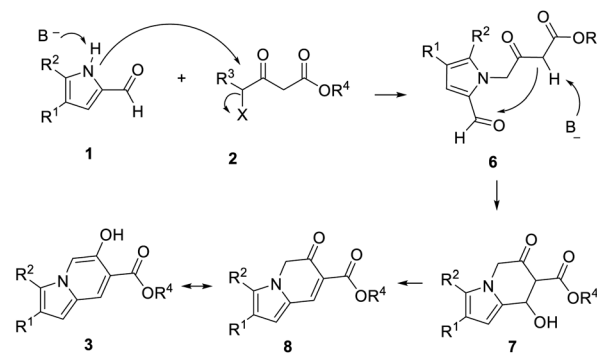
Table 1 Optimization of reaction conditions<sup>a</sup>

| Entry | Base                            | T (°C) | Solvent | Time | Yield (%) |
|-------|---------------------------------|--------|---------|------|-----------|
| 1     | K <sub>2</sub> CO <sub>3</sub>  | 50     | MeCN    | 6 h  | 56        |
| 2     | K <sub>2</sub> CO <sub>3</sub>  | 25     | MeCN    | 6 h  | 70        |
| 3     | DBU                             | 25     | MeCN    | 6 h  | 54        |
| 4     | Cs <sub>2</sub> CO <sub>3</sub> | 25     | MeCN    | 6 h  | 63        |
| 5     | NaOH                            | 25     | MeCN    | 6 h  | 42        |
| 6     | <i>t</i> -BuOK                  | 25     | MeCN    | 6 h  | 59        |
| 7     | NaOAc                           | 25     | MeCN    | 6 h  | 12        |
| 8     | CsOAc                           | 25     | MeCN    | 6 h  | 26        |
| 9     | K <sub>2</sub> CO <sub>3</sub>  | 25     | DMF     | 6 h  | 67        |
| 10    | K <sub>2</sub> CO <sub>3</sub>  | 25     | EtOH    | 6 h  | 60        |
| 11    | K <sub>2</sub> CO <sub>3</sub>  | 25     | Acetone | 6 h  | 62        |

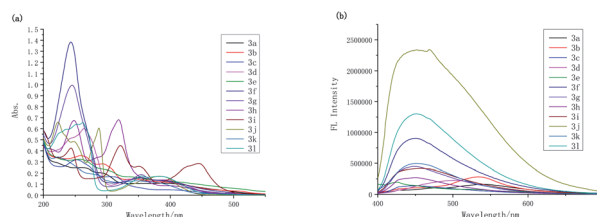
<sup>a</sup> 1 mmol **1a**, 2 mmol **2a**, 3 mmol base, and 10 mL solvent were used.Scheme 2 Substrate scope of pyrrole-2-formaldehyde. Reaction conditions: **1** (1 mmol), **2** (2 mmol), K<sub>2</sub>CO<sub>3</sub> (3 mmol), CH<sub>3</sub>CN (10 mL), 20 °C, 6 h.Fig. 1 X-ray crystal structure of compound **3k**.Scheme 3 Substrate scope studies of 4-halogenated acetoacetic ester. Reaction conditions: **4** (1 mmol), **2** (2 mmol), K<sub>2</sub>CO<sub>3</sub> (3 mmol), CH<sub>3</sub>CN (10 mL), 20 °C, 6 h.

(entry 2). Other bases such as DBU, Cs<sub>2</sub>CO<sub>3</sub>, NaOH, *t*-BuOK, NaOAc, and CsOAc were then evaluated. However, the yields decreased, especially for a weak base (NaOAc) (entries 3–8). Regarding the effect of solvents, no improvement in the yield was obtained when the reaction was carried out in DMF, EtOH, and acetone (entries 9–11).

Then, the reactions of various substituted pyrrole-2-formaldehyde were tested (Scheme 2). Generally, the desired products were obtained in moderate-to-good yields when the pyrroles contained electron-withdrawing groups at the 4- or 5-position. However, no product was obtained for pyrrole-2-formaldehyde, presumably due to the reduction of nucleophilicity of the pyrrole ring. The structure of compound **3k** was confirmed by X-ray crystal structure analysis (Fig. 1, CCDC 2081693†).



Scheme 4 The proposed mechanism.

Fig. 2 UV-vis and FL spectra of compounds **3a–3l**.

Next, the applicability of this cascade reaction was expanded to the synthesis of imidazo[1,5-*a*]pyridine, furnishing the desired product **5a** in 61% yield. The scope of the 4-halogenated acetoacetic ester was also evaluated (Scheme 3). The results indicate that the halogen and alkyl groups on position 4 and the ether group hardly influenced the yields.

Based on the above mentioned results and our previous work, we propose a mechanism as shown in Scheme 4. First, S<sub>N</sub>2 substitution of 4-halogenated acetoacetic ester **2** and pyrrole-2-formaldehyde **1** yields intermediate **6**. Subsequently, cyclized intermediate **7** is formed through intramolecular nucleophilic substitution. Finally, the desired products **3** are obtained through dehydration and tautomerism.

To advance our efforts to search for new fluorophores for cell imaging and their strong luminescence, we investigated the UV-vis and fluorescence spectra of these new compounds (Fig. 2). Compounds **3a–3l** show similar absorptions at *ca.* 250 nm (Table S1†), which should be assigned to the  $\pi$ – $\pi^*$  electronic transition originating from the indolizine ring. Notably, the substituent and their position on the indolizine ring slightly affect these absorption peaks. However, the weak absorption bands between 290 nm and 445 nm due to  $n$ – $\pi^*$  electronic transition are especially different for compounds **3h** and **3i** containing a strong electron-withdrawing group (NO<sub>2</sub>). The maximum emission bands of **3c**, **3e**, and **3f** are similar (425–455 nm, Table S1†), while those of **3a**, **3b**, and **3d** are 540 nm, 535 nm, and 505 nm, respectively, with a much higher red shift.

## Conclusions

In summary, we developed an efficient cascade reaction to construct indolizines and related imidazo[1,5-*a*]pyridines with a hydroxyl group, which is difficult to introduce through other methods. The structure was confirmed by single-crystal X-ray diffraction analysis. The compounds showed strong fluorescence in a dilute solution. Further studies on the optical properties of these indolizines are in progress.

## Conflicts of interest

There are no conflicts to declare.

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