Morita–Baylis–Hillman reaction of 3-formyl-9H-pyrido[3,4-b]indoles and fluorescence studies of the products

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Abstract

β-Carboline is a privileged class of the alkaloid family and is associated with a broad spectrum of biological properties. 3-Formyl-9H-pyrido[3,4-b]indole is a such potent precursor belonging to this family which can be tailored for installing diversity at various positions of β-carboline to generate unique molecular hybrids of biological importance. The present work is a step towards this and assimilates the results related to the exploration of 3-formyl-9H-β-carbolines for the synthesis of β-carboline C-3 substituted MBH adducts followed by evaluation of their fluorescent characteristic. The effect of contact time, solvent system, concentration and substituents was also studied during investigation of fluorescence properties of these derivatives.

Introduction

Among the polycyclic alkaloids based on indole, the tricyclic structure β-carboline represents a promising class of pyrido-indole alkaloids with a variety of biological activities which make them interesting synthetic targets [1-8]. Alkaloids containing the β-carboline nucleus in their molecular architecture are present ubiquitously in nature and a large number of natural products are reported representing this scaffold [9-16]. The key precursor used in the biosynthesis of β-carboline is L-tryptophan which forms the basis of great abundance of β-carboline-containing natural products [17]. A broad spectrum of biological activities is displayed by this pharmacologically rich nucleus which includes antibacterial, antifungal, anticancer, anxiolytic, antimalarial, antiviral, anti-HIV, anti-Alzheimer, and anticonvulsant activities etc. [18-26]. Potent anticancer activities are shown by the majority of β-carboline-containing compounds [27-30]. Figure 1 summarizes some examples of β-carboline-based drugs and bioactive natural products some of which have even been commercialized successfully showing the importance of this nucleus [31-33]. This pharmacological richness and colossal medicinal importance is the reason that the synthesis of β-carboline-containing derivatives has been an exciting area for researchers [34-40].
The Morita–Baylis–Hillman (MBH) reaction is an astonishing C–C bond forming reaction between a carbonyl electrophile and an activated alkene leading to the formation of allylic alcohol; a highly functionalized product [41-44]. The chemistry of the MBH reaction is decorated with several unique features viz. atom economy, complexity generation and generation of a chiral center from a pro-chiral electrophile. The chemistry of the MBH reaction has gained considerable attention from the past two decades as these MBH adducts are highly functionalized and offer various points of diversity. Due to these amazing features, these MBH adducts act as starting material on which various organic transformations can be performed leading to the synthesis of various natural and synthetic products. MBH adducts itself display diverse biological activities like antifungal, antibacterial, herbicide, antiparasitic and antitumor as reviewed by Lima-Junior et al. (2012) [45].

It was envisaged that in comparison to the traditional methods like Pictet–Spengler (P-S) or Bischler–Napieralski (B-N) cyclisation, introduction of a formyl group at C-1 or C-3 position of the β-carboline frameworks may provide a new route for generating unlimited diversity at C-1 as well as at the C-3 position of β-carbolines. As depicted in Figure 2, 1-formyl-β-carbolines and 3-formyl-β-carbolines are decorated with different sites for
diversification which make these synthons a promising template for the construction of β-carboline-fused frameworks via C-1 N-9, C-1 N-2 and C-3 N-2 cyclisation. Similarly, β-carboline-substituted molecular frameworks can be generated at C-3 position.

Our group has previously explored 1-formyl-β-carbolines and 3-formyl-β-carbolines for the generation of β-carboline-imidazo[1,2-a]azepine conjugates at C-1 as well as C-3 position by the application of the Groebke–Blackburn–Bienaymé (GBB) multicomponent approach [46,47]. Our research group has also investigated the scope of 1-formyl-β-carbolines for generating unique molecular hybrids by application of the Morita–Baylis–Hillman reaction [48-50]. It was also revealed from a detailed literature survey that only limited reports have been documented toward exploration of 3-formyl-9H-β-carbolines for generating diversity at the β-carboline skeleton as outlined in Figure 3 [51-56].

Therefore, we herein report the synthesis of C-3-substituted pyrido[3,4-b]indole MBH adducts from substituted 3-formyl-9H-β-carbolines by the application of the MBH reaction followed by evaluation of their fluorescence properties.

**Results and Discussion**

The current study began with the synthesis of substituted 3-formyl-9H-β-carbolines (6a–e), which was accomplished by modifying the previously disclosed process as presented in Scheme 1 [46,47,57]. Pictet–Spengler (P-S) condensation of L-tryptophan (1) with different aldehydes (a–e) in dry DCM at room temperature yielded tetrahydro-β-carboline derivatives 2a–e, which were then oxidized with KMnO$_4$ in anhydrous DMF for 45 minutes to yield β-carboline derivatives 3a–e. It was encouraging to observe that the P-S condensation with L-tryptophan (1) was much faster than with the tryptophan ester, taking only 45 minutes to complete. Interestingly, KMnO$_4$ oxidation was selective, with no decarboxylation seen. Within

![Figure 3](image-url) Figure 3: A summary of previous reports toward exploration of 3-formyl-9H-β-carbolines.
15 minutes, further treatment of \(3a-e\) with methyl iodide in the presence of \(K_2CO_3\) provided the corresponding methyl ester \(4a-e\) in high yield (83–87%) and ester functionality reduction with LiAlH\(_4\) in dry THF yielded the alcohols \(5a-e\) in excellent yield (90–98%). The required 3-formyl-9H-β-carbolines \(6a-e\) were obtained in 73–88% yield by oxidizing the alcohol derivatives \(5a-e\) with \(MnO_2\) in dry DCM. The present methodology is decorated with several advantages like scalability and selectivity. Additionally, no column chromatographic purification was required at any stage and each step was high yielding.

After the synthesis of starting materials, the Morita–Baylis–Hillman reaction was explored for C-3 functionalization of the β-carbolines \(6a-e\) for the \(β\)-carbolines \(6a-e\) were subjected to MBH reaction with acrylonitrile \(A\) and various acrylates \(B-E\) under neat conditions as depicted in Scheme 2. All the products were furnished smoothly in 27–72% yield. During this study, it was observed that the MBH reaction of \(6b\) with acrylonitrile \(A\) resulted in the formation of product \(8bA\) which evidenced that \(6b\) underwent Morita–Baylis–Hillman reaction at the electrophilic carbonyl center as well as Michael addition reaction at the nucleophilic nitrogen center (N-9). Similar results were obtained when \(6e\) was subjected to MBH reaction with acrylonitrile \(A\) and methylacrylate \(B\) and products \(8eA\) and \(8eB\) were generated as outlined in Scheme 2.

The effect of a substituent at N-9 position on the reactivity of 3-formyl-9H-β-carbolines \(6a-e\) was also investigated during this study. For this purpose, the N-ethyl derivative \(9e\) of \(6e\) was prepared and subjected to MBH reaction with acrylonitrile \(A\) and methylacrylate \(B\) under neat conditions to generate the corresponding MBH adducts (\(10eA\) and \(10eB\)) (Scheme 3). Interestingly, \(9e\) showed more affinity towards this C–C bond forming transformation than \(6e\). It is noteworthy here that all the products were purified by column chromatography.

A small library of C-3-substituted pyrido[3,4-b]indole derivatives was designed and synthesized which is presented in Figure 4. All the products were characterized using NMR, FTIR and mass spectrometry.

**Fluorescence studies**

Fluorescence studies of these C-3-substituted pyrido[3,4-b]indole derivatives were examined and various parameters (contact time, concentration and solvent) were optimized for obtaining the best results using \(7dA\) as a model substrate. Fluorescence emission spectra for optimizing the contact time were recorded in chloroform at different intervals of time (5 min, 15 min, 1 h and 24 h) at \(1 \times 10^{-6}\) M concentration. \(7dA\) displayed the highest fluorescence intensity after 15 minutes and its fluorescence activity lasted even after 24 h with a slight
Scheme 3: Synthesis of C-3 substituted pyrido[3,4-b]indole MBH derivatives 10.

Figure 4: Library of C-3-substituted pyrido[3,4-b]indole MBH derivatives 7, 8 and 10.

decrease in fluorescence intensity. Further, the fluorescence emission profile of 7dA was recorded in chloroform at different concentrations viz. $1 \times 10^{-6}$ M, $2 \times 10^{-6}$ M, $3 \times 10^{-6}$ M, $4 \times 10^{-6}$ M and $5 \times 10^{-6}$ M which indicated that fluorescence intensity was found to increase with increase in concentration and fluorescence spectra above this concentration showed a fluorescence intensity $>1000$ a.u. After optimizing the time and concentration parameters, dilutions of 7dA in different organic solvents such as dichloromethane, DMF and ethyl acetate were prepared for optimizing the solvent for obtaining the best fluo-
rescence results. Fluorescence spectra were recorded after 15 minutes of sample preparation in $5 \times 10^{-6}$ M concentration and fluorescence intensity was observed to be in the following order: CHCl$_3$ > EtOAc > CH$_2$Cl$_2$ > DMF. The results of the optimization studies are presented in Figure 5 and it was concluded from the studies that C-3-substituted pyrido[3,4-b]indole derivative 7dA displayed the maximum fluorescence intensity in chloroform at a concentration of $5 \times 10^{-6}$ M after 15 minutes of sample preparation.

Accordingly, fluorescence studies of all the other derivatives were conducted following these optimized parameters, i.e., time: 15 min; concentration: $5 \times 10^{-6}$ M; solvent: CHCl$_3$. The results of the fluorescence studies of all the C-3 substituted pyrido[3,4-b]indole derivatives are presented in Table 1.

### Structure–fluorescence activity relationships

From the results presented in Table 1, some structure–fluorescence activity relationships were concluded which are outlined in Figure 6. It was concluded from the structure–fluorescence activity relationships of C-3-substituted pyrido[3,4-b]indole derivatives that the products with o-bromophenyl substituent at R$^1$ position (7cB and 7cE) were the most fluorescent derivatives among all. Also, substituted phenyl derivatives were more fluorescent than the dimethoxymethyl substituted derivatives 7aA–7aE. Further, it was observed that an ethyl substituent at N-9 position of β-carboline decreased the fluorescence intensity in 10eA–10eB than 7eD which is a N-unsubstituted derivative while the $\lambda_{emission}$ was red shifted in N-ethyl-substituted derivatives as is clearly indicated from the data presented in Table 1. CO$_2$n-Bu and CO$_2$t-Bu substituents enhanced the fluo-

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**Figure 5:** Results of optimization for fluorescence studies: a) contact time; b) concentration; c) solvent.

| sample | compound | R$^1$ | R$^2$ | $\lambda_{Ex}$ (nm) | $\lambda_{Em}$ (nm) | fluorescence intensity (a.u.) |
|--------|----------|------|------|----------------------|----------------------|--------------------------------|
| 1      | 7aA      | CH(OMe)$_2$ | CN   | 266                 | 384                 | 669.99                         |
| 2      | 7aB      | CH(OMe)$_2$ | CO$_2$Me | 260               | 369                 | 624.79                         |
| 3      | 7aC      | CH(OMe)$_2$ | CO$_2$Et | 374               | 407                 | 223.68                         |
| 4      | 7aD      | CH(OMe)$_2$ | CO$_2$n-Bu | 258               | 378                 | 486.49                         |
| 5      | 7aE      | CH(OMe)$_2$ | CO$_2$t-Bu | 278               | 388                 | 669.30                         |
| 6      | 8bA      | Ph       | CN   | 278                 | 391                 | 592.48                         |
| 7      | 7bB      | Ph       | CO$_2$Me | 278               | 384                 | 768.75                         |
| 8      | 7bC      | Ph       | CO$_2$Et | 278               | 383                 | 766.89                         |
| 9      | 7cB      | 2-Br-C$_6$H$_4$ | CO$_2$Me | 278               | 375                 | >1000                          |
| 10     | 7cE      | 2-Br-C$_6$H$_4$ | CO$_2$t-Bu | 278               | 383                 | 834.06                         |
| 11     | 7dA      | 4-Br-C$_6$H$_4$ | CN   | 278                 | 386                 | 609.70                         |
| 12     | 7dB      | 4-Br-C$_6$H$_4$ | CO$_2$Me | 294               | 386                 | 768.78                         |
| 13     | 8eA      | 4-Cl-C$_6$H$_4$ | CN   | 270                 | 395                 | 385.38                         |
| 14     | 8eB      | 4-Cl-C$_6$H$_4$ | CO$_2$Me | 270               | 401                 | 353.34                         |
| 15     | 7eD      | 4-Cl-C$_6$H$_4$ | CO$_2$n-Bu | 292               | 385                 | 829.04                         |
| 16     | 10eA     | 4-Cl-C$_6$H$_4$ | CN   | 278                 | 403                 | 556.46                         |
| 17     | 10eB     | 4-Cl-C$_6$H$_4$ | CO$_2$Me | 270               | 428                 | 415.32                         |
rescence intensity more than the other substituents (7aE, 7cE and 7eD). It is noteworthy here that in the derivatives prepared from the same aldehydes, a noticeable decrease in fluorescence intensity of product 8 (Morita–Baylis–Hilman + Michael adducts 8bA, 8eA and 8eB) was observed than in case of product 7 (Morita–Baylis–Hilman adducts 7bB, 7bC and 7eD) while their λ_emission was red-shifted in comparison to type 7 compounds. This difference in the fluorescence intensity values of compound 7 and 8 may be attributed to the addition of a substituent at N-9 position of the β-carboline ring after Michael addition reaction (CH₂CH₂CN or CH₂CH₂CO₂Me).

**Conclusion**

In conclusion, we have successfully explored 3-formyl-1-aryl-9H-pyrido[3,4-b]indole derivatives for the C-3 functionalization by application of the MBH reaction to generate C-3-substituted β-carboline MBH adducts. It was revealed from spectroscopic analysis that few derivatives underwent MBH reaction as well as Michael addition reaction to form type 8 compounds. Additionally, the scope of the reaction was further extended and the effect of substituents at the N-9 position on the reactivity of 3-formyl-1-aryl-9H-pyrido[3,4-b]indoles was also investigated. Furthermore, fluorescence properties of these β-carboline conjugates were also studied and they were found to exhibit excellent fluorescence characteristics. Different parameters like contact time, concentration, solvent effects and substituent effects were examined for obtaining the optimal results. It was observed that the MBH derivatives exhibited excellent fluorescence characteristics at a concentration of 5 × 10⁻⁶ M in chloroform solvent after 15 minutes of sample preparation. Derivatives 7eB and 7eC bearing an o-bromophenyl substituent at R¹ position emerged as two most fluorescent compounds in the present series. Furthermore, products of type 7 (Morita–Baylis–Hilman adducts) were more fluorescent than products 8 (Morita–Baylis–Hilman + Michael adducts). Antimicrobial evaluation of the title compounds is underway and will be reported in due course.

**Supporting Information**

Supporting information contains detailed experimental procedure for the synthesis of compounds 6–9 and 10 followed by detailed characterization data and copies of ¹H NMR and ¹³C NMR spectra of newly synthesized compounds 6–10.

**Supporting Information File 1**

Experimental procedures and characterization data. [https://www.beilstein-journals.org/bjoc/content/supplementary/1860-5397-18-92-S1.pdf]

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