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Cu-catalyzed mild and efficient oxidation of TH β Cs using air: application in practical total syntheses of perlolyrine and flazin†

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A mild, efficient and environmentally benign method for synthesis of aromatic β -carbolines *via* Cu(II)-catalyzed oxidation of 1,2,3,4-tetrahydro- β -carbolines (TH β Cs) was developed, in which air (O₂) was used as the clean oxidant. This method has advantages such as environmentally friendliness, mildness, very good tolerance of functional groups, high yielding and easy experiment operation. In addition, this new methodology was successfully applied in the efficient and practical total syntheses of β -carboline alkaloids perlolyrine and flazin.

Introduction

The aromatic β -carboline skeleton is ubiquitous in many alkaloids,¹ bioactive congeners,² agrochemicals³ and functional materials.⁴ In view of the significance of β -carbolines in drug discovery, agricultural chemistry and material science, development of practical, efficient and environmentally benign methods for syntheses of β -carbolines is of considerable interest and has attracted much attention from chemists.⁵

Since 1,2,3,4-tetrahydro- β -carbolines (TH β Cs) can be readily and efficiently prepared *via* the Pictet–Spengler reaction,⁶ so the aromatization of TH β Cs *via* dehydrogenation or oxidation is an easy and good method to obtain β -carbolines. Dehydrogenation of TH β Cs usually employed precious metallic catalysts including Pd,⁷ Pt⁸ and complexes of Ru⁹ and Ir.¹⁰ Oxidation of TH β Cs normally needed stoichiometric strong oxidants such as KMnO₄,¹¹ MnO₂,¹² SeO₂,¹³ DDQ,¹⁴ TCCA¹⁵ and IBX.¹⁶ However, uses of expensive metallic catalysts and hazardous strong oxidants are sometimes lack of cheapness, practicality and environmental friendliness, therefore novel practical, efficient and environmentally benign methods for the conversion of TH β Cs to β -carbolines are highly desirable.

Oxygen is an important component of air, which is one of the most abundant resources on the earth, so the air has been used as a clean and eco-friendly oxidant for various oxidation reactions in recent decades.¹⁷ On the other hand, copper is one of the most abundant metals on the earth's crust, thus copper salts are normally very cheap, and moreover copper is a low

toxic transition metal, so an increasing amount of copper-catalyzed reactions have been developed recently.¹⁸ Herein, we describe a new practical and very mild Cu(II)-catalyzed oxidative conversion of TH β Cs to β -carbolines using air as the clean oxidant.

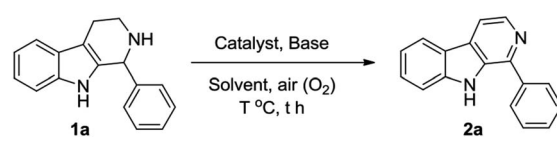
Results and discussion

At first, we attempted to find out the optimized reaction conditions for the Cu-catalyzed oxidative conversion of TH β Cs **1** to β -carbolines **2**. With the conversion 1-phenyl-TH β C **1a** to 1-phenyl- β -carboline **2a** as the model reaction, we tried the reaction under various conditions, and results are summarized in Table 1. As can be seen from Table 1, various copper reagents have been tested as catalysts (Table 1, Entries 2–11), and it was found that cupric bromide is the best catalyst for the reaction. When 0.2 equivalents of CuBr₂ was used as the catalyst, the aerobic oxidation of compound **1a** took place smoothly at room temperature in DMSO in the presence of 2 equivalents of 1,8-diazabicyclo[5,4,0]undec-7-ene (DBU) to afford the desired compound **2a** in high yield (Entry 2). Several amines such as DBU, 1,5-diazabicyclo[4,3,0]non-5-ene (DBN), 4-dimethylaminopyridine (DMAP), pyridine and trimethylamine have been tested as bases (Entries 2 and 12–15), DBU was found to be the most appropriate base for the reaction. Amount of DBU has significant effect on the reaction (Entries 2 and 16–19), almost no desired product **2a** could be obtained in the absence of DBU (Entry 16); and the reaction was hard to be complete even for a longer time, if less than 2.0 equivalents of DBU were used (Entries 17–19). Several other solvents such as *N,N*-dimethylformamide (DMF), acetonitrile, ethanol, tetrahydrofuran (THF), dichloromethane, ethyl acetate and acetone have also been examined (Entries 20–26); DMSO is obviously better than them. Additionally, when the reaction temperature was

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Table 1 Optimization of reaction conditions for the Cu-catalyzed aerobic oxidative conversion of 1-phenyl-TH β C **1a** to 1-phenyl- β -carboline **2a**^a

						
Entry	Catalyst ^b	Base ^c	Solvent	T (°C)	t (h)	Yield ^d (%)
1	None	DBU ^e	DMSO ^f	25	18	0
2	CuBr ₂	DBU	DMSO	25	18	95
3	CuCl ₂	DBU	DMSO	25	18	90
4	Cu(OAc) ₂	DBU	DMSO	25	18	86
5	CuBr	DBU	DMSO	25	18	85
6	CuCl	DBU	DMSO	25	18	84
7	CuI	DBU	DMSO	25	18	82
8	CuSO ₄	DBU	DMSO	25	24	<5
9	CuCO ₃	DBU	DMSO	25	24	11
10	CuO	DBU	DMSO	25	24	<5
11	Cu	DBU	DMSO	25	24	<5
12	CuBr ₂	DBN ^g	DMSO	25	20	88
13	CuBr ₂	DMAP ^h	DMSO	25	20	19
14	CuBr ₂	Py	DMSO	25	20	12
15	CuBr ₂	Et ₃ N	DMSO	25	20	10
16	CuBr ₂	None	DMSO	25	18	0
17	CuBr ₂	DBU (0.5) ⁱ	DMSO	25	60	75
18	CuBr ₂	DBU (1.0) ^j	DMSO	25	30	83
19	CuBr ₂	DBU (1.5) ^k	DMSO	25	25	92
20	CuBr ₂	DBU	DMF ^l	25	24	82
21	CuBr ₂	DBU	CH ₃ CN	25	24	65
22	CuBr ₂	DBU	EtOH	25	24	23
23	CuBr ₂	DBU	THF ^m	25	24	30
24	CuBr ₂	DBU	CH ₂ Cl ₂	25	24	27
25	CuBr ₂	DBU	EtOAc	25	24	12
26	CuBr ₂	DBU	Me ₂ CO	25	24	14
27	CuBr ₂	DBU	DMSO	45	13	90
28	CuBr ₂	DBU	DMSO	65	9	84
29	CuBr ₂	DBU	DMSO	85	6	76

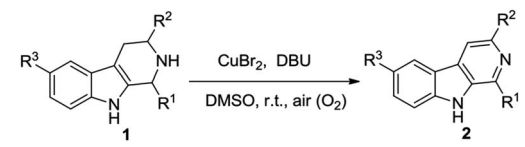
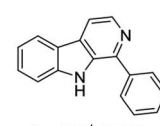
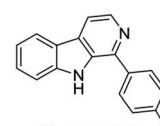
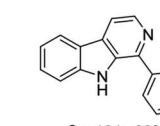
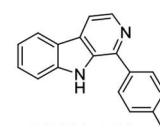
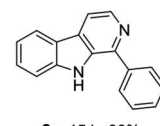
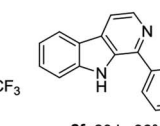
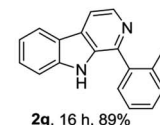
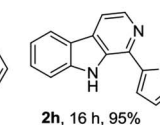
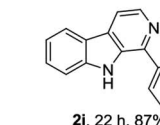
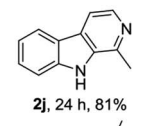
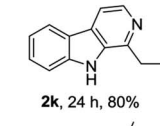
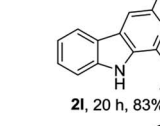
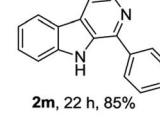
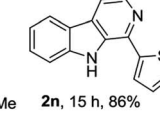
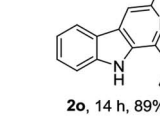
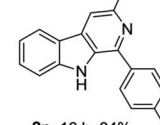
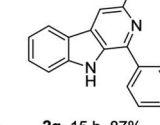
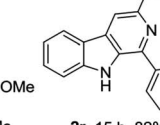
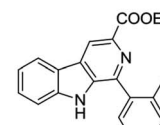
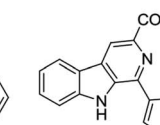
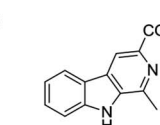
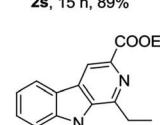
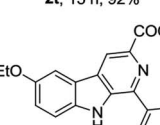
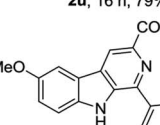
^a All reactions were performed at T °C for t hours under an atmosphere of air. ^b 20% (mol%) of the catalyst was used. ^c 2.0 equivalents of the base were used unless otherwise indicated. ^d Isolated yields. ^e 1,8-Diazabicyclo[5.4.0]undec-7-ene. ^f Dimethyl sulfoxide. ^g 1,5-Diazabicyclo[4.3.0]non-5-ene. ^h 4-Dimethylaminopyridine. ⁱ 0.5 equivalent of DBU was used. ^j 1.0 equivalent of DBU was used. ^k 1.5 equivalents of DBU was used. ^l N,N-Dimethylformamide. ^m Tetrahydrofuran.

elevated, the reaction velocity increased, but the yield of desired product **2a** obviously decreased (Entries 27–29).

Subsequently, we attempted the CuBr₂-catalyzed oxidative conversion of variously substituted TH β Cs **1** to β -carboline **2** in DMSO with DBU as the base, and the results are summarized in Table 2. The scope of the reaction is very wide; almost all of the tested TH β Cs could be smoothly converted to β -carbolines in good to excellent yields. All the conversions were performed under the standard reaction conditions (see footnote of the Table 2), and a total of twenty-four variously substituted β -carbolines **2a–2x** were obtained in 78–96% yields.

A possible mechanism for the CuBr₂-catalyzed oxidative conversion of TH β Cs **1** to β -carbolines **2** is proposed in Scheme

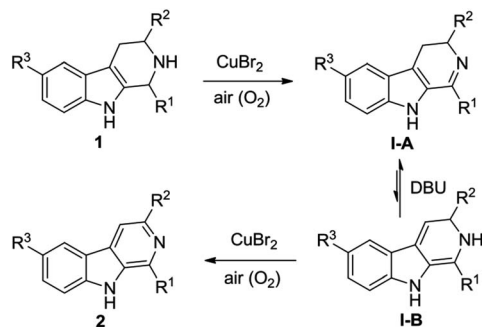
Table 2 CuBr₂-catalyzed oxidation of variously substituted TH β Cs **1** to β -carbolines **2** using air as the clean oxidant^a

		
 2a , 18 h ^b , 95% ^c	 2b , 12 h, 90%	 2c , 16 h, 88%
 2d , 18 h, 91%	 2e , 15 h, 90%	 2f , 20 h, 96%
 2g , 16 h, 89%	 2h , 16 h, 95%	 2i , 22 h, 87%
 2j , 24 h, 81%	 2k , 24 h, 80%	 2l , 20 h, 83%
 2m , 22 h, 85%	 2n , 15 h, 86%	 2o , 14 h, 89%
 2p , 16 h, 84%	 2q , 15 h, 87%	 2r , 15 h, 82%
 2s , 15 h, 89%	 2t , 15 h, 92%	 2u , 16 h, 79%
 2v , 16 h, 78%	 2w , 16 h, 87%	 2x , 18 h, 91%

^a Reaction conditions: TH β Cs **1** (2 mmol), CuBr₂ (0.4 mmol), DBU (4 mmol), DMSO (6 mL), stirring at room temperature (25 °C) under an atmosphere of air. ^b Reaction time. ^c Isolated yields.

1. As can be seen from Scheme 1, TH β Cs **1** would first undergo CuBr₂-catalyzed aerobic oxidation of C–N single bond to C=N double bond to produce 3,4-dihydro- β -carbolines **I-A** according to Adimurthy's reports.¹⁹ 3,4-Dihydro- β -carbolines **I-A** would then undergo DBU-promoted reversible tautomerization²⁰ to form unstable enamine intermediates **I-B**, which would also



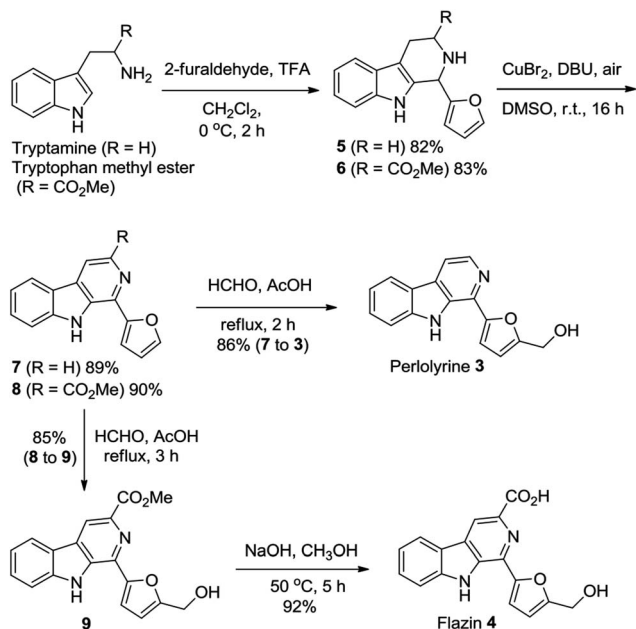


Scheme 1 Possible mechanism for the CuBr_2 -catalyzed oxidative conversion of TH β Cs 1 to β -carbolines 2.

undergo the CuBr_2 -catalyzed aerobic oxidation of C–N single bond to afford β -carbolines 2. Actually, dihydro- β -carbolines **I-A** could be isolated if the reaction was stopped in the midway. For example, when the CuBr_2 -catalyzed oxidative conversion of 1-phenyl-TH β C **1a** to 1-phenyl- β -carboline **2a** was stopped at 8 h, 1-phenyl-3,4-dihydro- β -carboline was obtained in 36% yield.

To showcase the synthetic utility of the above-described method for the CuBr_2 -catalyzed oxidative conversion of TH β Cs **1** to β -carbolines **2**, we have successively applied the methodology to the efficient and practical total syntheses of two β -carboline alkaloids perlopyrine **3** and flazin **4**.

Perlopyrine **3** and flazin **4** are both strongly fluorescent 1-furanyl- β -carboline alkaloids. These two particular alkaloids are widespread in nature, and have been isolated from plants, bacteria, Japanese sake and soy sauce.^{1h,3b,21} Several total syntheses of perlopyrine **3** and flazin **4** have been reported hitherto,²² we herein report novel practical total syntheses of them, which are depicted in Scheme 2. Pictet–Spengler reaction



Scheme 2 Novel concise total syntheses of β -carboline alkaloids perlopyrine **3** and flazin **4**.

of tryptamine and tryptophan methyl ester with 2-furaldehyde first gave TH β Cs **5** and **6** in 82% and 83% yields, respectively. CuBr_2 -catalyzed oxidation of compounds **5** and **6** with air then furnished β -carbolines **7** and **8** in 89% and 90% yields, respectively. Next, hydroxymethylation of compounds **7** and **8** with HCHO in AcOH produced perlopyrine **3** and compound **9** in 86% and 85% yields, respectively. Hydrolysis of the ester **9** afforded flazin **4** in 92% yield. Thus, perlopyrine **3** was synthesized from tryptamine in 3 steps in 63% overall yield; and flazin **4** was synthesized from tryptophan methyl ester in 4 steps in 58% overall yield.

Conclusions

In conclusion, we have developed a novel method for the CuBr_2 -catalyzed oxidative conversion of TH β Cs to β -carbolines. This method has some advantages as follows: (a) the process was carried out at room temperature with air as the clean oxidant, so it is very mild and environmentally benign; (b) CuBr_2 as the catalyst is inexpensive and low-toxic; (c) all products were obtained in good to excellent yields; (d) the reaction has very good tolerance of functional groups, it seemed to be applicable to all the tested substrates; (e) experiment operation is quite easy. In addition, novel practical total syntheses of β -carboline alkaloids perlopyrine and flazin were performed with the above-described CuBr_2 -catalyzed mild oxidation of TH β Cs as the key step.

Experimental

General

^1H NMR and ^{13}C NMR spectra were acquired on a Bruker AM 400 instrument, chemical shifts are given on the δ scale as parts per million (ppm) with tetramethylsilane (TMS) as the internal standard. IR spectra were recorded on a Nicolet Magna IR-550 instrument. Mass spectra were performed with a HP1100 LC-MS spectrometer. Melting points were measured on a Mei-TEMP II melting point apparatus. Column chromatography was performed on silica gel (Qingdao Chemical Factory). All reagents and solvents were analytically pure, and were used as such as received from the chemical suppliers.

General procedure for the preparation of various TH β Cs 1. 1-Substituted tryptamine (3 mmol) was dissolved in CH_2Cl_2 (10 mL), an aldehyde (3.3 mmol) was added. The resulting solution was cooled to 0°C , TFA (0.686 mg, 6.016 mmol) was added. The mixture was then stirred at $0-5^\circ\text{C}$ for 2–6 h. After the reaction was complete (checked by TLC, eluent: $\text{EtOAc}/\text{CH}_2\text{Cl}_2 = 1:4$), solvent was removed under vacuum, the residue was then partitioned between EtOAc (30 mL) and an aqueous solution of K_2CO_3 (10% w/w, 20 mL). Two phases were separated, and the aqueous phase was extracted again with EtOAc (20 mL). Organic extracts were combined, and dried over anhydrous MgSO_4 . Evaporation of solvent under vacuum gave crude product, which was purified by flash chromatography (eluent: $\text{MeOH}/\text{CH}_2\text{Cl}_2 = 1:20-1:10$) to afford TH β Cs **1a–1x** in 70–95% yields.

General procedure for the CuBr_2 -catalyzed oxidative conversion of TH β Cs **1 to β -carbolines **2**.** TH β Cs **1** (2 mmol) was dissolved in DMSO (5 mL). DBU (609 mg, 4.000 mmol) and



anhydrous CuBr₂ (89.5 mg, 0.401 mmol) were added. The resulting solution was then stirred under air at room temperature (around 25 °C) for 12–24 h (see Table 2). After the reaction was complete (checked by TLC, EtOAc/CH₂Cl₂ = 1 : 1 to 1 : 4), the reaction solution was poured into the mixed solution of EtOAc (30 mL) and aqueous ammonia (5% w/w, 25 mL). After the mixture was stirred for 5 minutes, two phases were separated. The aqueous layer was extracted again with EtOAc (20 mL). The organic extracts were combined and washed with brine (10 mL). The organic solution was dried over anhydrous MgSO₄, and then was concentrated under vacuum to give the crude product, which was purified by flash chromatography (eluent: EtOAc/CH₂Cl₂ = 1 : 2 to 1 : 15) to afford pure β-carbolines **2** in 78–96% yield as indicated in the Table 2. Characterization data of the β-carbolines **2a–2x** are given in the ESI† of this paper.

Total syntheses of perlolyrine **3** and flazin **4**

1-(Furan-2-yl)-1,2,3,4-tetrahydro-9H-pyrido[3,4-*b*]indole 5. A mixture of tryptamine (1.604 g, 10.01 mmol), 2-furaldehyde (1.076 g, 11.20 mmol) and CH₂Cl₂ (15 mL) was stirred and cooled to 0 °C by an ice-bath. TFA (2.285 g, 20.04 mmol) was slowly added, and then the reaction mixture was further stirred at 0 °C for 2 h. After the reaction was complete (checked by TLC, eluent: EtOAc/CH₂Cl₂ = 1 : 3), ethyl acetate (60 mL) and an aqueous solution of K₂CO₃ (10% w/w, 30 mL) were added. The mixture was vigorously stirred for 5 min, and then two phases were separated. Organic phase was extracted again with ethyl acetate (20 mL). Organic extracts were combined and dried over anhydrous MgSO₄, and then was concentrated under vacuum to give the crude product, which was purified by flash chromatography (eluent: MeOH/CH₂Cl₂ = 1 : 10) to afford pure compound **5** (1.957 g, 8.213 mmol) in 82% yield as white solid, mp 138–139 °C. ¹H NMR (CDCl₃, 400 MHz) δ 8.17 (s, 1H, NH in indole), 7.94 (s, 1H), 7.64 (d, *J* = 7.8 Hz, 1H), 7.48 (d, *J* = 3.2 Hz, 1H), 7.32 (d, *J* = 8.0 Hz, 1H), 7.18 (dd, *J*₁ = 8.0 Hz, *J*₂ = 7.9 Hz, 1H), 7.11 (dd, *J*₁ = 7.8 Hz, *J*₂ = 7.9 Hz, 1H), 6.96 (brs, 1H, NH), 6.66 (d, *J* = 3.7 Hz, 1H), 6.44 (dd, *J*₁ = 3.7 Hz, *J*₂ = 3.2 Hz, 1H), 3.90 (t, *J* = 7.3 Hz, 2H), 3.17 (t, *J* = 7.3 Hz, 2H). ¹³C NMR (CDCl₃, 100 MHz) δ 154.51, 142.52, 135.87, 131.84, 127.23, 121.91, 119.37, 118.34, 111.04, 110.35, 109.94, 107.76, 50.73, 41.46, 22.31. HRMS (ESI) *m/z* calcd for C₁₅H₁₄N₂ONa [M + Na]⁺: 261.1004, found: 261.1008. IR (KBr, film) 3393, 3145, 2922, 2846, 1450, 1298, 1141, 1010, 739 cm^{−1}.

Methyl1-(furan-2-yl)-1,2,3,4-tetrahydro-9H-pyrido[3,4-*b*]indole-3-carboxylate 6. Compound **6** was prepared from tryptophan methyl ester and 2-furaldehyde according to the same procedure as above for compound **5**. Compound **6** was obtained as an epimeric mixture of *cis* and *trans* isomers (*cis/trans* = 3 : 5) in a combined yield of 83% as offwhite solid. Two epimers can be separated by chromatography, but herein the epimeric mixture was used as such for the next step. Data for *cis* epimer: white solid, mp 141–142 °C. ¹H NMR (CDCl₃, 400 MHz) δ 7.98 (s, 1H, NH in indole), 7.50 (d, *J* = 7.6 Hz, 1H), 7.41 (d, *J* = 3.5 Hz, 1H), 7.25 (d, *J* = 7.8 Hz, 1H), 7.15 (dd, *J*₁ = 7.6 Hz, *J*₂ = 7.7 Hz, 1H), 7.11 (dd, *J*₁ =

7.7 Hz, *J*₂ = 7.8 Hz, 1H), 6.32–6.36 (m, 2H), 5.38 (s, 1H), 3.91 (dd, *J*₁ = 9.8 Hz, *J*₂ = 4.5 Hz, 1H), 3.79 (s, 3H), 3.17 (dd, *J*₁ = 15.0 Hz, *J*₂ = 4.5 Hz, 1H), 2.95 (dd, *J*₁ = 15.0 Hz, *J*₂ = 9.8 Hz, 1H), 2.44 (s, 1H, NH). ¹³C NMR (CDCl₃, 100 MHz) δ 173.06, 153.02, 142.85, 136.03, 131.79, 127.01, 122.18, 119.68, 118.28, 111.07, 110.48, 108.61, 107.78, 56.51, 52.33, 51.64, 25.46. HRMS (ESI) *m/z* calcd for C₁₇H₁₆N₂O₃Na [M + Na]⁺: 319.1059, found: 319.1057. IR (KBr, film) 3393, 2950, 1735, 1436, 1351, 1271, 1218, 1174, 1141, 1009, 740 cm^{−1}. Data for *trans* epimer: white solid, mp 145–146 °C. ¹H NMR (CDCl₃, 400 MHz) δ 8.30 (s, 1H, NH in indole), 7.49 (d, *J* = 7.5 Hz, 1H), 7.34 (d, *J* = 3.4 Hz, 1H), 7.16 (d, *J* = 7.8 Hz, 1H), 7.11 (dd, *J*₁ = 7.5 Hz, *J*₂ = 7.6 Hz, 1H), 7.06 (dd, *J*₁ = 7.6 Hz, *J*₂ = 7.8 Hz, 1H), 6.23 (dd, *J*₁ = 4.2 Hz, *J*₂ = 3.4 Hz, 1H), 5.96 (d, *J* = 4.2 Hz, 1H), 5.25 (s, 1H), 3.88 (dd, *J*₁ = 9.5 Hz, *J*₂ = 4.7 Hz, 1H), 3.69 (s, 3H), 3.13 (dd, *J*₁ = 15.2 Hz, *J*₂ = 4.7 Hz, 1H), 2.88 (*J*₁ = 15.2 Hz, *J*₂ = 9.5 Hz, 1H), 2.59 (brs, 1H, NH). ¹³C NMR (CDCl₃, 100 MHz) δ 173.82, 154.40, 142.67, 136.26, 130.97, 126.78, 122.14, 119.49, 118.27, 111.20, 110.28, 108.65, 108.14, 52.24, 52.23, 49.10, 25.17. HRMS (ESI) *m/z* calcd for C₁₇H₁₆N₂O₃Na [M + Na]⁺: 319.1059, found: 319.1055. IR (KBr, film) 3395, 2945, 1736, 1435, 1352, 1271, 1218, 1175, 738 cm^{−1}.

1-(Furan-2-yl)-9H-pyrido[3,4-*b*]indole 7. Compound **5** (0.962 g, 4.037 mmol) was dissolved in DMSO (8 mL), DBU (1.225 g, 8.047 mmol) and anhydrous CuBr₂ (0.179 g, 0.801 mmol) were added. The resulting solution was then stirred under air at 25 °C for 16 h. After the reaction was complete (checked by TLC, eluent: EtOAc/CH₂Cl₂ = 1 : 1), the reaction solution was poured into the mixed solution of EtOAc (50 mL) and aqueous ammonia (5% w/w, 20 mL). After the mixture was stirred for 5 minutes, two phases were separated. The aqueous layer was extracted again with EtOAc (20 mL). The organic extracts were combined and washed with brine (10 mL). The organic solution was dried over anhydrous MgSO₄, and then was concentrated under vacuum to give the crude product, which was purified by flash chromatography (eluent: EtOAc/CH₂Cl₂ = 1 : 3) to afford pure compound **7** (0.842 g, 3.594 mmol) in 89% yield as white solid, mp 166–167 °C. ¹H NMR (DMSO-*d*₆, 400 MHz) δ 11.50 (s, 1H, NH in indole), 8.38 (d, *J* = 5.2 Hz, 1H), 8.25 (d, *J* = 7.8 Hz, 1H), 8.08 (d, *J* = 5.2 Hz, 1H), 7.99 (d, *J* = 1.6 Hz, 1H), 7.77 (d, *J* = 8.2 Hz, 1H), 7.58 (dd, *J*₁ = 7.6 Hz, *J*₂ = 7.8 Hz, 1H), 7.23–7.32 (m, 2H), 6.79 (dd, *J*₁ = 3.5 Hz, *J*₂ = 3.2 Hz, 1H). ¹³C NMR (CDCl₃, 100 MHz) δ 155.09, 143.34, 141.09, 139.47, 134.09, 131.97, 130.83, 129.17, 122.25, 121.87, 120.72, 114.26, 112.92, 112.25, 109.32. HRMS (ESI) *m/z* calcd for C₁₅H₁₁N₂O [M + H]⁺: 235.0871, found: 235.0872. IR (KBr, film) 3458, 2998, 2920, 1624, 1557, 1493, 1424, 1377, 1235, 997, 754, 459 cm^{−1}.

Methyl1-(furan-2-yl)-9H-pyrido[3,4-*b*]indole-3-carboxylate 8. Compound **8** was prepared from compound **6** according to the same procedure as above for compound **7**. Compound **8** was obtained in 90% yield as white solid, mp 168–169 °C. ¹H NMR (DMSO-*d*₆, 400 MHz) δ 11.93 (s, 1H, NH in indole), 8.90 (s, 1H), 8.44 (d, *J* = 8.0 Hz, 1H), 8.06 (d, *J* = 3.3 Hz, 1H), 7.84 (d, *J* = 8.2 Hz, 1H), 7.65 (dd, *J*₁ = 8.0 Hz, *J*₂ = 8.1 Hz, 1H), 7.32–7.40 (m, 2H), 6.85 (dd, *J*₁ = 3.3 Hz, *J*₂ = 3.7 Hz, 1H), 3.96 (s, 3H). ¹³C NMR



(DMSO-*d*₆, 100 MHz) δ 165.76, 152.11, 144.23, 141.48, 136.37, 132.76, 132.11, 129.51, 128.79, 121.90, 120.82, 120.48, 116.41, 112.97, 112.34, 109.92, 52.07. HRMS (ESI) *m/z* calcd for C₁₇H₁₃N₂O₃ [M + H]⁺: 293.0926, found: 293.0922. IR (KBr, film) 3440, 3060, 2951, 1702, 1624, 1493, 1430, 1368, 1346, 1260, 738, 494 cm⁻¹.

1-[5-(Hydroxymethyl)furan-2-yl]-9H-pyrido[3,4-*b*]indole (perlolyrine 3). Compound 7 (0.470 g, 2.006 mmol) was dissolved in acetic acid (3 mL). An aqueous solution of formaldehyde (37% w/w, 1 mL) was added. The resulting solution was then heated and stirred at reflux for 2 h. After the reaction was complete (checked by TLC, eluent: EtOAc/CH₂Cl₂ = 1 : 3), the solvent was removed by vacuum distillation to give a viscous residue, which was partitioned between ethyl acetate (30 mL) and an aqueous solution of potassium carbonate (10% w/v, 20 mL). Two phases were separated, and aqueous phase was extracted again with ethyl acetate (20 mL). The organic extracts were combined and washed twice with brine (2 × 10 mL). After having been dried with anhydrous MgSO₄, the organic solution was concentrated under vacuum to give crude solid, which was purified by flash chromatography (eluent: MeOH/CH₂Cl₂ = 1 : 10) to give pure perlolyrine 3 (0.457 g, 1.729 mmol) in 86% yield as pale yellow solid, mp 165–166 °C. ¹H NMR (DMSO-*d*₆, 400 MHz) δ 11.24 (s, 1H, NH in indole), 8.38 (d, *J* = 5.1 Hz, 1H), 8.27 (d, *J* = 7.8 Hz, 1H), 8.08 (d, *J* = 5.1 Hz, 1H), 7.78 (d, *J* = 8.2 Hz, 1H), 7.61 (dd, *J*₁ = 7.8 Hz, *J*₂ = 8.0 Hz, 1H), 7.29 (dd, *J*₁ = 8.0 Hz, *J*₂ = 8.2 Hz, 1H), 7.23 (d, *J* = 3.3 Hz, 1H), 6.60 (d, *J* = 3.3 Hz, 1H), 5.50 (t, *J* = 6.0 Hz, 1H, OH), 4.68 (d, *J* = 6.0 Hz, 2H). ¹³C NMR (DMSO-*d*₆, 100 MHz) δ 156.73, 152.14, 140.92, 138.16, 133.14, 130.47, 129.42, 128.39, 121.59, 120.61, 119.68, 113.60, 112.40, 109.62, 109.04, 55.95. HRMS (ESI) calcd for C₁₆H₁₃N₂O₂ [M + H]⁺: 265.0977, found: 265.0974. IR (KBr, film) 3370, 2920, 1629, 1568, 1426, 1317, 1234, 1014, 799, 743, 634 cm⁻¹.

Methyl 1-(5-(hydroxymethyl)furan-2-yl)-9H-pyrido[3,4-*b*]indole-3-carboxylate 9. Compound 9 was prepared from compound 8 according to the same procedure as above for perlolyrine 3. Compound 9 was obtained in 85% yield as white solid, mp 205–206 °C. ¹H NMR (DMSO-*d*₆, 400 MHz) δ 11.63 (s, 1H, NH in indole), 8.87 (s, 1H), 8.44 (d, *J* = 7.9 Hz, 1H), 7.82 (d, *J* = 8.2 Hz, 1H), 7.66 (dd, *J*₁ = 7.9 Hz, *J*₂ = 8.0 Hz, 1H), 7.35 (dd, *J*₁ = 8.0 Hz, *J*₂ = 8.2 Hz, 1H), 7.29 (d, *J* = 3.4 Hz, 1H), 6.63 (d, *J* = 3.4 Hz, 1H), 5.50 (t, *J* = 6.1 Hz, 1H, OH), 4.68 (d, *J* = 6.1 Hz, 2H), 3.94 (s, 3H). ¹³C NMR (DMSO-*d*₆, 100 MHz) δ 165.92, 157.42, 151.21, 141.45, 136.65, 132.99, 132.14, 129.65, 129.01, 122.11, 121.08, 120.71, 116.43, 112.94, 110.99, 109.36, 56.07, 52.23. HRMS (ESI) *m/z* calcd for C₁₈H₁₅N₂O₄ [M + H]⁺: 323.1032, found: 323.1033. IR (KBr, film) 3406, 3297, 2922, 1727, 1565, 1435, 1351, 1251, 1118, 1005, 746 cm⁻¹.

1-(5-(Hydroxymethyl)furan-2-yl)-9H-pyrido[3,4-*b*]indole-3-carboxylic acid (flazin 4). Compound 9 (0.323 g, 1.002 mmol) was dissolved in methanol (5 mL), and then an aqueous solution of NaOH (3 mol L⁻¹, 1 mL) was added. The mixture was then heated and stirred at 50 °C for 5 h. After the reaction was complete (checked by TLC, eluent: EtOAc/CH₂Cl₂ = 1 : 1), ethyl acetate (25 mL) and water (20 mL) were added. The mixture was vigorously stirred, and pH value of the aqueous

solution was adjusted to 4–5 by adding citric acid. Two phases were separated, and the aqueous solution was extracted twice with ethyl acetate (2 × 25 mL). Organic extracts were combined and washed twice with brine (2 × 10 mL). After having been dried over anhydrous MgSO₄, the organic solution was concentrated under vacuum to give a crude solid product, which was purified by flash chromatography (eluent: MeOH/CH₂Cl₂ = 1 : 5) to afford pure flazin 4 (0.284 g, 0.921 mmol) in 92% yield as pale yellow solid, mp 239–240 °C. ¹H NMR (DMSO-*d*₆, 400 MHz) δ 11.69 (s, 1H, NH in indole), 8.84 (s, 1H), 8.42 (d, *J* = 7.8 Hz, 1H), 7.87 (d, *J* = 8.2 Hz, 1H), 7.65 (dd, *J*₁ = 7.8 Hz, *J*₂ = 8.0 Hz, 1H), 7.42 (d, *J* = 3.2 Hz, 1H), 7.34 (dd, *J*₁ = 8.0 Hz, *J*₂ = 8.2 Hz, 1H), 6.62 (d, *J* = 3.2 Hz, 1H), 4.69 (s, 2H). ¹³C NMR (DMSO-*d*₆, 100 MHz) δ 166.64, 157.45, 151.40, 141.53, 137.13, 132.63, 132.06, 129.98, 129.05, 122.17, 121.11, 120.69, 115.85, 112.94, 111.22, 109.37, 56.11. HRMS (ESI) *m/z* calcd for C₁₇H₁₂N₂O₄Na [M + Na]⁺: 331.0695, found: 331.0697. IR (KBr, film) 3242, 2922, 1735, 1624, 1612, 1576, 1360, 1321, 1018, 742 cm⁻¹.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- (a) W.-H. Jiao, G.-D. Chen, H. Gao, J. Li, B.-B. Gu, T.-T. Xu, H.-B. Yu, G.-H. Shi, F. Yang, X.-S. Yao and H.-W. Lin, *J. Nat. Prod.*, 2015, **78**, 125; (b) K. Ohishi, K. Toume, M. A. Arai, T. Koyano, T. Kowithayakorn, T. Mizoguchi, M. Itoh and M. Ishibashi, *J. Nat. Prod.*, 2015, **78**, 1139; (c) A. Furusato, H. Kato, T. Nehira, K. Eguchi, T. Kawabata, Y. Fujiwara, F. Losung, R. E. P. Mangindaan, N. J. d. Voogd, M. Takeya, H. Yokosawa and S. Tsukamoto, *Org. Lett.*, 2014, **16**, 3888; (d) M. Kitajima, S. Ohara, N. Kogure, D. Santiarworn and H. Takayama, *Tetrahedron*, 2013, **69**, 9451; (e) H. Huang, Y. Yao, Z. He, T. Yang, J. Ma, X. Tian, Y. Li, C. Huang, X. Chen, W. Li, S. Zhang, C. Zhang and J. Ju, *J. Nat. Prod.*, 2011, **74**, 2122; (f) W. D. Inman, W. M. Bray, N. C. Gassner, R. S. Lokey, K. Tenny, Y. Y. Shen, K. TenDyke, T. Suh and P. Grews, *J. Nat. Prod.*, 2010, **73**, 255; (g) W.-H. Jiao, H. Gao, C.-Y. Li, F. Zhao, R.-W. Jiang, Y. Wang, G.-X. Zhou and X.-S. Yao, *J. Nat. Prod.*, 2010, **73**, 167; (h) S. Fotso, R. P. Maskey, D. Schroder, A. S. Ferrer, I. Grun-Wollny and H. Laatsch, *J. Nat. Prod.*, 2008, **71**, 1630; (i) R. Cao, W. Peng, Z. Wang and A. Xu, *Curr. Med. Chem.*, 2007, **14**, 479.
- (a) A. Kamal, M. Sathish, V. L. Nayak, V. Srinivasulu, B. Kavitha, Y. Tangella, D. Thummuri, C. Bagul, N. Shankaraiah and N. Nagesh, *Bioorg. Med. Chem.*, 2015, **23**, 5511; (b) F. C. Savariz, M. A. Foglio, A. L. T. G. Ruiz, W. F. d. Costa, M. d. M. Silva, J. C. C. Santos,



- I. M. Figueiredo, E. Meyer, J. E. d. Carvalho and M. H. Sarragiotto, *Bioorg. Med. Chem.*, 2014, **22**, 6867; (c) B. Shi, R. Cao, W. Fan, L. Guo, Q. Ma, X. Chen, G. Zhang, L. Qiu and H. Song, *Eur. J. Med. Chem.*, 2013, **60**, 10.
- 3 (a) H. Song, Y. Liu, Y. Liu, L. Wang and Q. Wang, *J. Agric. Food Chem.*, 2014, **62**, 1010; (b) Y. Li, M. Zhao and K. L. Parkin, *J. Agric. Food Chem.*, 2011, **59**, 2332; (c) T. Herraiz, *J. Agric. Food Chem.*, 2007, **55**, 8534.
- 4 (a) Y. Im and L. Y. Lee, *Chem. Commun.*, 2013, **49**, 5948; (b) B. K. Paul, N. Ghosh and S. Mukherjee, *RSC Adv.*, 2016, **6**, 9984.
- 5 (a) F. Nissen, V. Richard, C. Alayrac and B. Witulski, *Chem. Commun.*, 2011, **47**, 6656; (b) S. Ding, Z. Shi and N. Jiao, *Org. Lett.*, 2010, **12**, 1540; (c) G. Verniest, D. England, N. D. Kimpe and A. Padawa, *Tetrahedron*, 2010, **66**, 1496; (d) H. Zhang and R. C. Larock, *J. Org. Chem.*, 2002, **67**, 7048; (e) N. Kanekiyo, T. Kuwada, T. Choshi, J. Nobuhiro and S. Hibino, *J. Org. Chem.*, 2001, **66**, 8793.
- 6 (a) E. D. Cox and J. M. Cook, *Chem. Rev.*, 1995, **95**, 1797; (b) J. Stockigt, A. P. Antonchick, F. Wu and H. Waldmann, *Angew. Chem., Int. Ed.*, 2011, **50**, 8538.
- 7 (a) R. S. Kusurkar, N. A. H. Alkobati, A. S. Gokule and V. G. Puranik, *Tetrahedron*, 2008, **64**, 1654; (b) A. Kulkarni, M. Abid, B. Torok and X. Huang, *Tetrahedron Lett.*, 2009, **50**, 1791; (c) S. Eagon and M. O. Anderson, *Eur. J. Org. Chem.*, 2014, 1653.
- 8 D. Ge, L. Hu, J. Wang, X. Li, F. Qi, J. Lu, X. Cao and H. Gu, *ChemCatChem*, 2013, **5**, 2183.
- 9 S. Muthaiah and S. H. Hong, *Adv. Synth. Catal.*, 2012, **354**, 3045.
- 10 J. Wu, D. Talwar, S. Johnston, M. Yan and J. Xiao, *Angew. Chem., Int. Ed.*, 2013, **52**, 6983.
- 11 (a) M. Zheng, Y. Yang, M. Zhao, X. Zhang, J. Wu, G. Chen, L. Peng, Y. Wang and S. Peng, *Eur. J. Med. Chem.*, 2011, **46**, 1672; (b) P. Ashok, S. Chander, J. Balzarini, C. Pannecouque and S. Murugesan, *Bioorg. Med. Chem. Lett.*, 2015, **25**, 1232.
- 12 W. Yin, P. V. V. S. Sarma, J. Ma, D. Han, J. L. Chen and J. M. Cook, *Tetrahedron Lett.*, 2005, **46**, 6363.
- 13 M. Cain, O. Campos, F. Guzman and J. M. Cook, *J. Am. Chem. Soc.*, 1983, **105**, 907.
- 14 K. Kondo, H. Shigemori, Y. Kikuchi, M. Ishibashi, T. Sasaki and J. Kobayashi, *J. Org. Chem.*, 1992, **57**, 2480.
- 15 B. Bai, L. Shen, J. Ren and H. J. Zhu, *Adv. Synth. Catal.*, 2012, **354**, 354.
- 16 J. D. Panarese and S. P. Waters, *Org. Lett.*, 2010, **12**, 4086.
- 17 (a) Z. Shi, C. Zhang, C. Tang and N. Jiao, *Chem. Soc. Rev.*, 2012, **41**, 3381; (b) S. E. Allen, R. R. Walvoord, R. Padilla-Salinas and M. C. Kozlowski, *Chem. Rev.*, 2013, **113**, 6234; (c) S. D. McCann and S. S. Stahl, *Acc. Chem. Res.*, 2015, **48**, 1756; (d) A. E. Wendlandt, A. M. Suess and S. S. Stahl, *Angew. Chem., Int. Ed.*, 2011, **50**, 11062; (e) A. Shaabani, S. Shaabani and H. Afaridoun, *RSC Adv.*, 2016, **6**, 48396; (f) C. Li, S. An, Y. Zhu, J. Zhang, Y. Kang, P. Liu, Y. Wang and J. Li, *RSC Adv.*, 2014, **4**, 49888; (g) H. Tian, H. Qiao, C. Zhu and H. Fu, *RSC Adv.*, 2014, **4**, 2694; (h) A. E. Fernandes, O. Riant, A. M. Jonas and K. F. Jensen, *RSC Adv.*, 2016, **6**, 36602; (i) C. Zhang, P. Zhao, Z. Zhang, J. Zhang, P. Yang, P. Gao, J. Gao and D. Liu, *RSC Adv.*, 2017, **7**, 47366.
- 18 (a) C. Zhang, C. Tang and N. Jiao, *Chem. Soc. Rev.*, 2012, **41**, 3464; (b) C. Sambiagio, S. P. Marsden, A. J. Blacker and P. C. McGowan, *Chem. Soc. Rev.*, 2014, **43**, 3525; (c) X.-X. Guo, D.-W. Gu, Z. Wu and W. Zhang, *Chem. Rev.*, 2015, **115**, 1622.
- 19 (a) R. D. Patil and S. Adimurthy, *Adv. Synth. Catal.*, 2011, **353**, 1695; (b) R. D. Patil and S. Adimurthy, *RSC Adv.*, 2012, **2**, 5119.
- 20 (a) R. A. Clark and D. C. Parker, *J. Am. Chem. Soc.*, 1971, **93**, 7257; (b) J. J. Eisch and F. J. Gadek, *J. Org. Chem.*, 1971, **36**, 2065.
- 21 (a) T. Liu, W. Liang and G. Tu, *Planta Med.*, 1988, **54**, 472; (b) B.-N. Su, L. C. Chang, E. J. Park, M. Guendet, B. D. Santarsiero, A. D. Mesecar, R. G. Mehta, H. H. S. Fong, J. M. Pezzuto and A. D. Kinghorn, *Planta Med.*, 2002, **68**, 730; (c) J. A. D. Jeffreys, *J. Chem. Soc. C*, 1970, 1091; (d) S.-i. Nakatsuka, B.-n. Feng, T. Goto and K. Kihara, *Tetrahedron Lett.*, 1986, **27**, 3399; (e) B. H. Han, M. H. Park, Y. N. Han and L. K. Woo, *Arch. Pharmacol. Res.*, 1986, **9**, 21; (f) B. H. Han, J. H. Park, M. H. Park and Y. N. Park, *Arch. Pharmacol. Res.*, 1985, **8**, 249.
- 22 (a) N. Wu, F. Song, L. Yan, J. Li and J. You, *Chem. – Eur. J.*, 2014, **20**, 3408; (b) B. Dassonneville, B. Witulski and H. Detert, *Eur. J. Org. Chem.*, 2011, 2836; (c) W. P. Gessner and A. Brossi, *Arch. Pharm.*, 1988, **321**, 95; (d) F. Bracher and D. Hildebrand, *Liebigs Ann. Chem.*, 1992, 1315.

