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TCCA-mediated oxidative rearrangement of tetrahydro- β -carbolines: facile access to spirooxindoles and the total synthesis of (\pm) -coerulescine and (\pm) -horsfiline†

Manda Sathish, ‡^{ad} Akash P. Sakla, ‡^b Fabiane M. Nachtigall, c Leonardo S. Santos (10 **d and Nagula Shankaraiah (10 **b)

Multi-reactive centered reagents are beneficial in chemical synthesis due to their advantage of minimal material utilization and formation of less by-products. Trichloroisocyanuric acid (TCCA), a reagent with three reactive centers, was employed in the synthesis of spirooxindoles through the oxidative rearrangement of various N-protected tetrahydro- β -carbolines. In this protocol, low equivalents of TCCA were required to access spirooxindoles (up to 99% yield) with a wide substrate scope. Furthermore, the applicability and robustness of this protocol were proven for the gram-scale total synthesis of natural alkaloids such as (\pm) -coerulescine (1) and (\pm) -horsfiline (2) in excellent yields.

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Introduction

Spirooxindoles are exceptional and versatile scaffolds, and thus have been extensively studied in the fields of synthetic and pharmaceutical chemistry.1 The unique three-dimensional spiro system may be the main cause for the bioactivities of spirooxindoles.^{1,2} Especially, spirooxindoles have been considered to exhibit antitumor,3 antimicrobial,4 antioxidant,5 antiinflammatory,6 antiviral1a and other bioactivities. Moreover, many of their derivatives have been extensively used in clinical trials,7 and some representative examples of bioactive spirooxindoles are depicted in Fig. 1. For example, horsfiline (2) is an intoxicating snuff,8 rhynchophylline (4) is potent towards various cancer cell lines,9 spirotryprostatin A prevents G2M progression in mammalian tsFT210 cells, 10 and corynoxine and corynoxine B show potential for the treatment of Parkinson's disease.11 Due to all these pharmaceutical potencies of spirooxindole derivatives, chemists have been inspired to establish various synthetic routes for this class of compounds.

Fig. 1 Representative examples of bioactive spirooxindoles (1–4).

Several approaches have been proposed in the literature to build spirooxindoles, which mainly involve two ways: (i) multistep synthesis14,8,12 and (ii) oxidative rearrangement.13 However, oxidative rearrangement reaction is more beneficial than multistep synthesis, because it only involves a single step, avoids the use of various toxic reagents, and ultimately time saving. A few examples of oxidants have been reported previously to afford spirooxindoles, which involve the use of mild organic oxidants such as oxone (Fig. 2),13f N-bromosuccinimide (NBS), 13c t-BuOCl13a,13d and transition metal oxidants such as Pb(OAc)₄, ^{13a} CrO₃ ^{13f} and OsO₄. ^{13b} However, the latter oxidants are more likely to produce highly toxic by-products. In continuation of our efforts in the development of new synthetic strategies, 14 we established a simple method employing the safer multi-reactive centered reagent (MRCR) chloroisocyanuric acid (TCCA), a highly desirable catalyst that requires less equivalents, thereby reducing the by-products.

R = H, Coerulescine (1)
R = OMe, Horsfiline (2)

Me

Me

OMe

NH

Me

OMe

Rhynchophylline (4)

^aInstituto de Investigación Interdisciplinaria, Vicerrectoría Académica, Universidad de Talca, Campus Lircay, Talca 3460000, Chile

^bDepartment of Medicinal Chemistry, National Institute of Pharmaceutical Education and Research (NIPER), Hyderabad 500 037, India. E-mail: shankar@niperhyd.ac.in ^cInstituto de Ciencias Químicas Aplicadas, Universidad Autónoma de Chile, Sede Talca 3467987, Chile

^aLaboratory of Asymmetric Synthesis, Instituto de Química de Recursos Naturales, Universidad de Talca, Campus Lircay, Talca 3460000, Chile. E-mail: lssantos@ utalca.cl

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[‡] M. Sathish and A. P. Sakla contributed equally.

Previous oxidants: (a) oxone (>1.0 equiv.) (b) OsO₄ (>1.0 equiv.) (c) NBS (>1.0 equiv.)/AcOH (d) Pb(OAc)₄ (>1.0 equiv.) This work: TCCA (0.35 equiv.) THF/water (1:1) rt, 30 min., up to 99%

Fig. 2 Comparison of present work with previous methods.

Particularly, TCCA is inexpensive and produces essentially the nontoxic cyanuric acid as a by-product, which can be easily separated from the reaction mixture.15 TCCA is a versatile reagent with three reactive N-Cl centers, which helps to utilize only 0.33-0.5 equivalents in the reaction. In scaled-up reactions, it is also possible to re-generate TCCA by passing chlorine gas in the aqueous mixture of cyanuric acid. 16 Moreover, it is widely used in several applications such as chlorination and mild oxidation.17 Studer and co-workers18 utilized TCCA to obtain α-chloro aldehydes and α-chloro ketones from their corresponding 1° and 2° alcohols. Bathini and co-workers also reported the TCCA-mediated decarboxylative/dehydrogenative aromatization of tetrahydro-β-carbolines (THBCs), which was applied for the total synthesis of β-carboline alkaloids. 19 Veisi and co-workers developed a method to produce nitriles through the direct oxidation of the corresponding amines, alcohols, aldehydes, and benzyl halides using TCCA.20 Thus, based on these significant findings such as bioactivity and research interest on the synthesis of spirooxindoles, we developed an efficient protocol to construct spirooxindole via oxidative rearrangement using TCCA, which is a cost effective and versatile reagent.

Results and discussion

All the key substrates 5a-z were synthesized according to known reports.21-23 Initially, we performed a model reaction with tetrahydro-β-carboline (5a) and TCCA (1 equiv.) in an acidic solvent mixture of THF/water/AcOH (1:1:1) for 4 h at 0 °C. The formation of spirooxindole 6a was observed with 45% yield (entry 1, Table 1). Next, we considered to avoid the use of acidic solvents and executed another reaction without AcOH, which afforded spirooxindole 6a with slightly improved yield (58%, entry 2, Table 1). We realized that the acidic nature of AcOH and HCl liberated from excess equivalents of TCCA may affect the yield of 6a. Then, after reducing the TCCA equivalents to 0.33 at 0 °C for 2 h (entry 3, Table 1), we achieved 88% of 6a. However, a slight improvement in the yield (90%) was seen after minimizing the reaction time to 30 min at room temperature (entry 4, Table 1). Considering our enthusiasm, another trial with similar reaction conditions was performed by taking 0.35 equiv. of TCCA, and surprisingly 6a was isolated with excellent yield (99%, entry 5, Table 1). Hence, we established that 0.35 equiv. of TCCA is optimal for this oxidative rearrangement reaction. Further investigation using other immiscible solvent mixtures such as DCE/water, DCM/water, and EtOAc/water achieved poor isolated yields of **6a** (entries 6, 7 and 8, respectively, Table 1), thus showing the incompatibility of immiscible solvent mixtures for this reaction. However, considerable yields of 6a were identified in the case of miscible solvent mixtures such as MeOH/water (82%) and MeCN/water (90%) (entries 9 and 10, respectively, Table 1).

Moreover, two reactions were tested in pure THF and distilled water as solvents, but only 40% of **6a** was isolated in the case of water (entry 11, Table 1) and no product (**6a**) was

Table 1 TCCA-mediated oxidative rearrangement^a

Entry	TCCA (equiv.)	Solvent	Temp. (°C)	Yield (%)
1	1	THF/water/AcOH (1:1:1)	0	45^b
2	1	THF/water (1:1)	0	58^b
3	0.33	THF/water $(1:1)$	0	88^b
4	0.33	THF/water $(1:1)$	rt	90
5	0.35	THF/water (1 : 1)	rt	99
6	0.35	DCE/water $(1:1)$	rt	50
7	0.35	DCM/water(1:1)	rt	52
8	0.35	EtOAc/water(1:1)	rt	55
9	0.35	MeOH/water(1:1)	rt	82
10	0.35	MeCN/water (1:1)	rt	90
11	0.35	Water	rt	40^c
12	0.35	THF	rt	_

^a All reactions were performed using tetrahydro-β-carboline **5a** (1 equiv.) and TCCA for 30 min. ^b The reaction was stirred for 2 h. ^c The remaining starting material was recovered.

bserved in THF (entry 12, Table 1). Based on all these obserations, the optimized protocol for the oxidative rearrangement of tetrahydro-β-carbolines to spirocyindoles is 0.35 equiv of

observed in THF (entry 12, Table 1). Based on all these observations, the optimized protocol for the oxidative rearrangement of tetrahydro- β -carbolines to spirooxindoles is 0.35 equiv. of TCCA and THF/water (1:1) as the solvent at room temperature for 30 min. During the reaction optimization, we identified that a homogenous reaction mixture is required before adding TCCA to achieve the complete conversion of the substrate. Accordingly, it is recommended to use 2:1 of THF/water if precipitation of tetrahydro- β -carbolines occurs in a 1:1 mixture of THF/water to ensure a homogeneous reaction mixture is obtained.

Having the optimized protocol in hand, we explored a sequential study on different substituted/protected THBCs 5a-z, which delivered the corresponding spirooxindoles 6a-x, 1 and 2 in good to excellent yields (78–99%). Totally, the protocol was executed on five types of THBC substrates based on their substitution/protection pattern as follows: (i) simple THBCs with N2-protection (5a-i); (ii) THBCs with N2-protection and N2-protection (5j-m); (iii) THBCs with N2-protection and N2-protection (5n-t); (iv) THBCs with N2-protection and C3-substitutions (5u and 5v); and (v) THBCs with C1-substitution, N2-protection and C3-substitutions (5w-z).

The sulfonyl groups ($R^2 = \text{tosyl}$ and mesyl) on N2 of THBCs (5a and 5b) smoothly underwent oxidative rearrangement to produce the corresponding spirooxindoles 6a and 6b with excellent yields (99% and 94%, Table 2), respectively. However, other electron-withdrawing groups (EWG) such as acetyl, benzoyl, CO2Et, Boc and Cbz at N2-position of simple THBCs (5c-g) were only slightly affected to give lower yields (82%, 81%, 83%, 88% and 88%) of the corresponding spirooxindoles (6c-g, Table 2), respectively, which may be due to the acid sensitivity of the amide and carbamate functionalities. Furthermore, the reaction of 5c with TCCA was performed at 0 °C, and a low yield (50%) was observed at room temperature, affording de-protected 5c. However, an electron-donating group (EDG) such as methyl on N2 of THBC (5h and 5i) gave good yields of natural spirooxindoles (\pm)-coerulescine (1, 93%) and (\pm)-horsfiline (2, 90%), respectively, as depicted in Table 2.

Further, our attempts on the oxidative rearrangement of THBCs 5k-m with indole nitrogen protection succeeded and achieved the corresponding spirooxindoles 6h-k in high yields, respectively. The EDGs (R^4 = benzyl and methyl) at the indole nitrogen of THBCs delivered 6h and 6i with satisfactory yields (93% and 95%), respectively. Interestingly, our protocol also worked well on EWG ($R^4 = \text{tosyl}$ and Boc)-protected THBCs (51) and 5m) to provide spirooxindoles 6j (78%) and 6k (82%) without affecting the protecting groups (Table 2), respectively. A recent report13f stated that the presence of EDGs such as hydrogen, alkyl and benzyl at the indolyl nitrogen (N-R4) is essential to achieve oxidative rearrangement. It is noticeable that our protocol is very advantageous given that it works well on both EDGs and EWGs on the indole nitrogen. Next, we explored our protocol on C1 alkyl (R^1 = methyl, ethyl and isopropyl)- and phenyl-substituted THBCs 5n-t, but only alkylsubstituted spirooxindoles 6l-q were obtained in appreciable yields (80-89%, Table 2). A low yield was observed for 6r (27%) at room temperature from 1-phenyl substituted THBC (5t), and

TCCA (0.35 equiv.) THF/water (1:1) 6a (99%) 6b (94%) 6d (89%) 6c (82%)b .COOF Boo 6e (83%) 6f (88%) **6g** (88%) 1 (93%) .COOEt 2 (90%) 6h (93%) 6i (95%) **6j** (78%) Boc 6k (82%) **6m** (89%, dr 1:4)^c 6n (85%) 6I (87%, dr 1:1.5)^o **6o** (87%) **6p** (80%) 6q (85%) 6r (45%, dr 1:7.9)b,c ·N Bn **6u** (95%, *dr* 1.7:1)^c 6t (93%, dr 3:1) 6s (95%, dr 1.9:1)^c 6v (94%, dr 1:8.3)^c 6w (89%, dr 4.5:1)c 6x (91%, dr 1:6.2)

^a All reactions were performed using tetrahydro-β-carboline 5 (1 equiv.) and TCCA (0.35 equiv.) for 30 min at room temperature. ^b The reaction was stirred at 0 °C. ^c The diastereomeric ratio (dr) measured by ¹H NMR analysis.

interestingly another ring-opened side product (**6r**′, 64%) was isolated and confirmed by comparing its NMR data (for NMR and plausible mechanism see ESI†) with the available literature. However, a slight improvement in yield (45%) was observed with good diastereoselectivity (**6r**, dr 1 : 7.9, Table 2) when the reaction was carried out at 0 °C and the yield of the ring-opened side product (**6r**′) decreased to 50%. The diastereomers of spirooxindoles **6l-q** could not be separated by column chromatography and the ratio was determined by ¹H

Scheme 1 (A) Imine formation of N2 unprotected THBCs. (B) Plausible reaction mechanism for the oxidative rearrangement of N2-protected THBCs.

Scheme 2 Gram-scale synthesis of (\pm) -coerulescine (1) and (\pm) -horsfiline (2).

NMR. Nevertheless, the THBCs $5\mathbf{u}$ and $5\mathbf{v}$ generated from L-tryptophan smoothly underwent oxidative rearrangement to produce spirooxindoles $6\mathbf{s}$ and $6\mathbf{t}$ in excellent yields and good diastereoselectivities (95%, dr 1.9:1 and 93%, dr 3:1 respectively, Table 2). Finally, the diastereomers of $5\mathbf{w}$ – \mathbf{z} with C1 ($\mathbf{R}^1=$ methyl or ethyl) and C3 ($\mathbf{R}^3=\mathrm{CO_2Me}$) substitutions were easily separated by column chromatography and employed for oxidative rearrangement to afford spirooxindole products $6\mathbf{u}$ – \mathbf{x} in good yields with prominent diastereomeric ratios (95%, dr 1.7:1, 94%, dr 1:8.3, 89%, dr 4.5:1 and 91%, dr 1:6.2, respectively) as shown in Table 2.

Further, we tested this protocol on C1-substituted (R^1 = methyl and phenyl) and N2-unprotected (R^2 = hydrogen) THBCs 7a and 7b and isolated dihydro- β -carbolines 8a and 8b,

albeit with very poor yields (35% and 30%, respectively) and no oxidative rearrangement product was observed. This can be attributed to dehydrohalogenation (I and II, Scheme 1A) instead of chlorohydrin formation, oxidative rearrangement, and thereby stabilization of the dihydro- β -carbolines by conjugation from imine, as shown in Scheme 1A.

Notably, in the reaction path of oxidative rearrangement, one mole of TCCA is sufficient to produce three moles of spirooxindoles. The reaction mechanism can be explained as initial TCCA chlorination of THBC to form Cl-THBC (III, Scheme 1B), formation of chlorohydrin (IV, Scheme 1B) and dehydrohalogenation followed by semi-pinacol-type rearrangement. The by-product cyanuric acid can be easily separated in the work-up given that it is soluble in water.

To highlight the utility of this protocol, we accomplished the gram-scale total synthesis of two bioactive spirooxindole natural products, (\pm) -coerulescine (1) and (\pm) -horsfiline (2) from 5h and 5i (Scheme 2), respectively. The gram-scale reaction of bioactive natural products is highly desirable from a commercial perspective. Specifically, 5h (1.2 g) and 5i (1.1 g) were reacted with TCCA to produce (\pm) -coerulescine (1) and (\pm) -horsfiline (2) in high yields (1.21 g), 93% and (1.07 g), (1.2 g) and (1.07 g), (1.2 g) and (1.2 g) are spectively. It is important to highlight that the quantity of TCCA used for this gram-scale synthesis is very low (0.52 g) for (1.2 g) and (1.2 g) for (1.2 g) and (1.2 g) for (1.2 g) for (1.2 g) for (1.2 g) and (1.2 g) for (1.2 g) for (1.2 g) for (1.2 g) and (1.2 g) for (1.2 g)

Conclusions

In conclusion, we developed an operationally simple and high yield protocol for the synthesis of spirooxindoles from the corresponding tetrahydro- β -carbolines (THBCs) using the inexpensive TCCA. The reaction proceeds *via* oxidative rearrangement and produced spirooxindoles in good to excellent yields (up to 99%). Diversely substituted spirooxindoles and naturally occurring (\pm)-coerulescine and (\pm)-horsfiline were furnished in excellent yields using the optimized reaction conditions. To demonstrate the commercial importance of this protocol, we also carried out gram-scale reactions to produce (\pm)-coerulescine (1) and (\pm)-horsfiline (2) with yields of 93% and 90%, respectively. In addition, this synthetic strategy is amenable for the generation of a library spiro-compounds and their derivatives, which can be further utilized in the drug discovery process.

Experimental

General

All solvents used for the reactions and purifications were of commercial grade or were purified prior to use when necessary. NMR spectral analyses were done on a Bruker 400 MHz or 500 MHz spectrometer for 1 H and 100 MHz or 125 MHz spectrometer for 13 C spectra, and calibrated to either TMS ($\delta=0$ for 1 H) or residual DMSO ($\delta=2.50$ for 1 H and $\delta=39.51$ for 13 C) and residual CHCl₃ ($\delta=7.26$ for 1 H and $\delta=77.23$ for 13 C). Spin multiplicities are described as s (singlet), br s (broad singlet),

d (doublet), t (triplet), dd (double doublet), td (triple doublet), q (quartet), and m (multiplet) and the coupling constants are reported in hertz (Hz). TLC analyses for the reactions and chromatography purifications were performed using silica gel plates (0.25 mm, E. Merck, 60 F254) with iodine and a UV lamp for visualization. Mass spectral measurements were recorded *via* electrospray ionization mass spectrometry (ESI-MS). HRMS was performed on a Varian QFT-ESI instrument. Melting points were measured on an electrothermal melting point apparatus and are uncorrected.

All the THBCs 5a, ^{21a,22a} 5b, ^{21b} 5c, ^{21b} 5d, 5e, ^{21b} 5f, ^{21b} 5g, ^{21c} 5h, ^{13e} 5i, ^{13e} 5j, 5k, ^{21a} 5l, 5m, 5n, 5o, ^{22a} 5p, 5q, ^{21b} 5r, 5s, ^{22b} 5t, ²³ 5u, ^{23a} 5v, ^{23c} 5x, ^{23c} 5y (ref. 23*d* and *e*) and 5z (ref. 23*d* and *e*) were synthesized using known procedures in the literature from their corresponding starting materials such as tryptamine (1a), tryptophan methyl ester (1b) and 5-methoxy tryptamine (1c). For the preparation of 5a-i and 5n-z, initially we reacted 1 with the respective aldehydes to achieve Pictet-Spengler cyclized products (NH-THBCs). Later, N-protection of THBCs was performed with the corresponding protecting groups in the presence of bases such as triethylamine and *N*,*N*-diisopropylethylamine, as mentioned in the respective literature reports. ^{13e,21-23}

General procedure for N9-protection for the synthesis of 5jm. Briefly, 5j, 5k and 5m were synthesized from 5a, whereas 5l was synthesized from 5e using protecting groups (PG) such as BnBr, MeI, TsCl and Boc₂O, respectively. To a stirred solution of 5a (1 equiv.) in dry THF, 60% NaH (1.5 equiv.) was added at 0 °C. After stirring for 15 min at room temperature, the protecting group (1.1 equiv. of BnBr for 5j, MeI for 5k and TsCl for 5m) was added at 0 °C and the reaction was stirred at room temperature for 8 h. The progress of the reaction was monitored by TLC analysis using EtOAc: hexane or MeOH: CH2-Cl₂ solvent systems and the ninhydrin charring technique. After completion, the reaction was quenched with saturated NH₄Cl solution and extracted with EtOAc and the combined organic phases were washed with water. The organic phase was evaporated in vacuo after drying over Na2SO4 to give the crude product, which was purified by silica gel column chromatography using 20-50% EtOAc/hexane to afford the pure products 5j, 5k and 5m. The same procedure was used for the synthesis of 51 from 5e using Boc₂O as the protecting group.

2-Tosyl-2,3,4,9-tetrahydro-1*H*-pyrido[3,4-*b*]indole (5a). (21*b*,22*a* White solid (340 mg, 90% yield), mp: 110–115 °C; ¹H NMR (400 MHz, DMSO- d_6): δ 10.75 (s, 1H), 7.71 (d, J = 8.3 Hz, 2H), 7.40 (d, J = 8.1 Hz, 2H), 7.34 (d, J = 7.8 Hz, 1H), 7.29 (d, J = 8.0 Hz, 1H), 7.03 (t, J = 8.0 Hz, 1H), 6.94 (t, J = 8.0 Hz, 1H), 4.27 (s, 2H), 3.39 (t, J = 5.7 Hz, 2H), 2.73 (t, J = 5.6 Hz, 2H), 2.37 (s, 3H); ¹³C NMR (100 MHz, DMSO- d_6 + CDCl₃): δ 143.4, 135.9, 133.5, 129.7, 129.1, 127.1, 126.1, 120.9, 118.4, 117.5, 111.0, 106.1, 43.9, 43.3, 20.9, 20.9; HRMS (ESI) calcd for $C_{18}H_{19}N_2O_2S$ m/z 327.1162 [M + H]⁺, found 327.1174.

2-(Methylsulfonyl)-2,3,4,9-tetrahydro-1*H*-pyrido[3,4-*b*]indole (5b). Light yellow solid (360 mg, 98% yield), mp: 200–202 °C; ¹H NMR (500 MHz, CDCl₃): δ 7.88 (s, 1H), 7.49 (d, J = 7.8 Hz, 1H), 7.35 (d, J = 8.1 Hz, 1H), 7.20 (t, J = 8.1 Hz, 1H), 7.13 (t, J = 8.1 Hz, 1H), 4.57 (s, 2H), 3.72 (t, J = 5.8 Hz, 2H), 2.93 (t, J =

5.8 Hz, 2H), 2.85 (s, 3H); 13 C NMR (125 MHz, CDCl₃): δ 136.3, 128.6, 126.9, 122.5, 120.1, 118.3, 111.1, 108.5, 44.0, 43.4, 37.5, 21.2; HRMS (ESI) calcd for $C_{12}H_{15}N_2O_2S$ m/z 251.0849 [M + H]⁺, found 251.0895.

Phenyl(1,3,4,9-tetrahydro-2*H*-pyrido[3,4-*b*]indol-2-yl)

methanone (5d). Light yellow solid (375 mg, 87% yield), mp: 153–155 °C; ¹H NMR (500 MHz, CDCl₃): δ 8.37 (s, 0.76H), 7.80 (s, 0.24H), 7.50–7.44 (m, 6H), 7.31 (d, J=7.3 Hz, 1H), 7.16 (t, J=7.4 Hz, 1H), 7.11 (t, J=7.3 Hz, 1H), 4.95 (s, 1.5H), 4.60 (s, 0.5H), 4.13 (s, 0.5H), 3.73 (s, 1.5H), 2.93 (s, 0.5H), 2.84 (s, 1.5H); ¹³C NMR (125 MHz, CDCl₃): δ 171.7, 136.4, 136.2, 130.0, 128.7, 127.0, 127.0, 122.1, 122.0, 119.8, 118.0, 111.1, 108.1, 46.1, 41.2, 22.2; HRMS (ESI) calcd for C₁₅H₁₈NOS₂ m/z 277.1335 [M + H]⁺, found 277.1350.

2-Methyl-2,3,4,9-tetrahydro-1*H*-pyrido[3,4-*b*]indole (5h). ^{13e} Off-white solid (298 mg, 93% yield), mp: 207–212 °C; ¹H NMR (400 MHz, DMSO- d_6 + CDCl₃): δ 9.70 (s, 1H), 7.42 (d, J = 7.6 Hz, 1H), 7.29 (d, J = 8.1 Hz, 1H), 7.12–6.96 (m, 2H), 3.59 (s, 2H), 2.87–2.75 (m, 4H), 2.50 (s, 3H); ¹³C NMR (100 MHz, DMSO- d_6 + CDCl₃): δ 135.8, 132.0, 126.6, 120.2, 118.2, 117.1, 110.5, 106.5, 52.6, 52.0, 45.3, 21.1; HRMS (ESI): calcd for C₁₂H₁₅N₂ m/z 187.1230 [M + H]⁺, found 187.1238.

6-Methoxy-2-methyl-2,3,4,9-tetrahydro-1*H***-pyrido**[3,4-*b*]**í ndole** (5**i**). ^{13e} Pale yellow oil (302 mg, 91% yield); ¹H NMR (400 MHz, DMSO- d_6): δ 10.56 (s, 1H), 7.20 (d, J = 8.0 Hz, 1H), 6.90 (s, 1H), 6.69 (d, J = 8.0 Hz, 1H), 3.78 (s, 3H), 3.55 (s, 2H), 2.71 (s, 4H), 2.44 (s, 3H); ¹³C NMR (100 MHz, DMSO- d_6): δ 152.4, 133.0, 130.3, 126.4, 110.8, 109.2, 105.4, 99.2, 54.7, 52.1, 51.6, 44.9, 20.8; HRMS (ESI): calcd for C₁₃H₁₇N₂O m/z 217.1335 [M + H]⁺, found 217.1343.

9-Benzyl-2-tosyl-2,3,4,9-tetrahydro-1*H*-pyrido[3,4-*b*]indole

(5j). Pale yellow solid (210 mg, 83% yield); mp: 213–215 °C; 1 H NMR (400 MHz, DMSO- d_{6}): δ 7.65 (d, J = 8.1 Hz, 2H), 7.38 (dd, J = 12.8, 8.2 Hz, 4H), 7.32–7.20 (m, 3H), 7.07 (t, J = 7.5 Hz, 1H), 7.02–6.97 (m, 3H), 5.36 (s, 2H), 4.27 (s, 2H), 3.38 (s, 2H), 2.73 (s, 2H), 2.35 (s, 3H); 13 C NMR (100 MHz, DMSO- d_{6}): δ 143.5, 138.0, 136.5, 133.7, 130.3, 129.8, 128.6, 127.3, 127.2, 126.4, 125.9, 121.3, 119.0, 117.9, 109.7, 106.8, 45.8, 43.7, 42.7, 20.9, 20.8; HRMS (ESI): calcd for $C_{25}H_{25}N_{2}O_{2}$ S m/z 417.1631 [M + H]⁺, found 417.1629.

Ethyl 9-tosyl-1,3,4,9-tetrahydro-2*H*-pyrido[3,4-*b*]indole-2-carboxylate (5l). Brown oil (286 mg, 85% yield); ¹H NMR (400 MHz, CDCl₃): δ 8.13 (d, J = 8.3 Hz, 1H), 7.71 (br s, 2H), 7.35 (d, J = 7.6 Hz, 1H), 7.33–7.27 (m, 1H), 7.24 (d, J = 7.4 Hz, 1H), 7.20 (d, J = 8.2 Hz, 2H), 4.95 (s, 2H), 4.20 (q, J = 7.1 Hz, 2H), 3.75 (s, 2H), 2.69 (s, 2H), 2.33 (s, 3H), 1.31 (t, J = 7.1 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 155.8, 145.0, 136.2, 136.1, 135.6, 130.0, 129.7, 129.6, 126.6, 124.7, 123.6, 118.4, 114.4, 61.8, 43.4, 41.1, 29.8, 21.6, 14.8; HRMS (ESI): calcd for C₂₁H₂₃N₂O₄S m/z 399.1373 [M + H]⁺, found 399.1381.

tert-Butyl 2-tosyl-1,2,3,4-tetrahydro-9*H*-pyrido[3,4-*b*]indole-9-carboxylate (5m). White solid (276 mg, 86% yield), mp: 223–228 °C; ¹H NMR (400 MHz, CDCl₃): δ 8.12 (d, J = 8.1 Hz, 1H), 7.74 (d, J = 8.0 Hz, 2H), 7.34 (dd, J = 19.0, 7.8 Hz, 3H), 7.28–7.18 (m, 2H), 4.54 (s, 2H), 3.46 (t, J = 5.6 Hz, 2H), 2.80 (t, J = 5.3 Hz, 2H), 2.41 (s, 3H), 1.68 (s, 9H); ¹³C NMR (100 MHz, CDCl₃): δ 149.9, 143.8, 135.8, 134.2, 130.0, 129.8, 128.6, 127.6, 124.4,

122.9, 117.9, 115.6, 114.8, 84.3, 45.8, 43.2, 28.3, 21.6, 21.6 HRMS (ESI): calcd for $C_{23}H_{26}N_2O_4S$ m/z 426.1613 $[M + H]^+$, found 426.1617.

1-Methyl-2-(methylsulfonyl)-2,3,4,9-tetrahydro-1*H***-pyrido** [3,4-*b*]indole (5n). White solid (333 mg, 91% yield), mp: 176–178 °C; ¹H NMR (500 MHz, CDCl₃): δ 7.92 (s, 1H), 7.49 (d, J = 7.8 Hz, 1H), 7.35 (d, J = 8.1 Hz, 1H), 7.20 (t, J = 7.8 Hz, 1H), 7.13 (t, J = 7.8 Hz, 1H), 5.09 (q, J = 6.7 Hz, 1H), 4.17 (dd, J = 14.5, 5.7 Hz, 1H), 3.49–3.40 (m, 1H), 2.96–2.87 (m, 1H), 2.82 (s, 3H), 2.79 (dd, J = 16.0, 4.3 Hz, 1H), 1.60 (d, J = 6.7 Hz, 3H); ¹³C NMR (125 MHz, CDCl₃): δ 135.9, 133.7, 126.6, 122.4, 119.8, 118.2, 111.0, 107.4, 48.6, 40.2, 39.0, 21.4, 20.8; HRMS (ESI) calcd for C₁₃H₁₇N₂O₂S m/z 265.1005 [M + H]⁺, found 265.0983.

1-Methyl-2-tosyl-2,3,4,9-tetrahydro-1*H*-**pyrido**[3,4-*b*]**indole** (50). ^{22a} Yellow solid (353 mg, 81% yield), mp: 206–207 °C; 1 H NMR (400 MHz, CDCl₃): δ 7.94 (s, 1H), 7.67 (d, J = 8.3 Hz, 2H), 7.35 (d, J = 7.8 Hz, 1H), 7.28 (d, J = 8.0 Hz, 1H), 7.18–7.10 (m, 3H), 7.06 (t, J = 8.3 Hz, 1H), 5.26 (q, J = 6.6 Hz, 1H), 4.16–4.09 (dd, J = 16.0, 4.0 Hz, 1H), 3.42–3.34 (m, 1H), 2.63–2.45 (m, 2H), 2.31 (s, 3H), 1.54 (d, J = 6.7 Hz, 3H); 13 C NMR (100 MHz, CDCl₃): δ 143.3, 138.3, 136.0, 134.0, 129.7, 126.8, 126.7, 122.1, 119.6, 118.2, 111.0, 107.7, 48.9, 39.3, 21.5, 21.5, 20.7; HRMS (ESI) calcd for $C_{19}H_{21}N_2O_2$ S m/z 341.1318 [M + H]⁺, found 341.1326.

1-Ethyl-2-(methylsulfonyl)-2,3,4,9-tetrahydro-1*H*-**pyrido**[3,4-*b*]**indole** (5**p**). White solid (301 mg, 78% yield), mp: 160–162 °C;

¹H NMR (500 MHz, CDCl₃): δ 7.91 (s, 1H), 7.49 (d, J = 7.8 Hz, 1H), 7.35 (d, J = 8.1 Hz, 1H), 7.21 (t, J = 8.1 Hz, 1H), 7.13 (t, J = 7.9 Hz, 1H), 4.81 (dd, J = 8.9, 5.1 Hz, 1H), 4.21 (dd, J = 14.9, 6.0 Hz, 1H), 3.50–3.42 (m, 1H), 2.99–2.90 (m, 1H), 2.82–2.75 (m, 1H), 2.76 (s, 3H), 1.97–1.83 (m, 2H), 1.14 (t, J = 7.4 Hz, 3H);

¹³C NMR (125 MHz, CDCl₃): δ 135.9, 133.1, 126.7, 122.4, 119.9, 118.3, 111.0, 107.4, 54.3, 40.0, 39.4, 28.6, 20.2, 10.9; HRMS (ESI) calcd for C₁₄H₁₉N₂O₂S m/z 279.1162 [M + H]⁺, found 279.1132.

1-Ethyl-2-tosyl-2,3,4,9-tetrahydro-1*H***-pyrido**[3,4-*b*]indole (**5q**). ^{21b} Pale yellow solid (366 mg, 89% yield), mp: 68–70 °C; ¹H NMR (500 MHz, CDCl₃): δ 7.81 (s, 1H), 7.63 (d, J = 8.3 Hz, 2H), 7.30 (t, J = 7.4 Hz, 2H), 7.14 (t, J = 7.8 Hz, 1H), 7.09 (d, J = 8.2 Hz, 2H), 7.04 (t, J = 7.5 Hz, 1H), 5.06 (dd, J = 8.4, 5.1 Hz, 1H), 4.13 (dd, J = 14.7, 5.7 Hz, 1H), 3.45–3.34 (m, 1H), 2.49 (dd, J = 15.6, 4.1 Hz, 1H), 2.4–2.3 (m, 1H), 2.28 (s, 3H), 1.99–1.82 (m, 2H), 1.12 (t, J = 7.4 Hz, 3H); ¹³C NMR (125 MHz, CDCl₃): δ 143.1, 138.2, 135.8, 133.0, 129.4, 126.7, 126.6, 121.9, 119.4, 118.1, 110.8, 107.8, 54.4, 39.7, 29.0, 21.4, 19.7, 10.8; HRMS (ESI) calcd for $C_{20}H_{23}N_2O_2S$ m/z 355.1475 [M + H]⁺, found 355.1482.

1-Isopropyl-2-(methylsulfonyl)-2,3,4,9-tetrahydro-1*H***-pyrido** [3,4-*b*]indole (5r). White solid (386 mg, 87% yield), mp: 128–129 °C; ¹H NMR (500 MHz, CDCl₃): δ 7.92 (s, 1H), 7.50 (d, J = 7.8 Hz, 1H), 7.36 (d, J = 8.1 Hz, 1H), 7.21 (t, J = 8.1 Hz, 1H), 7.13 (t, J = 7.5 Hz, 1H), 4.51 (d, J = 8.3 Hz, 1H), 4.23 (dd, J = 15.1, 6.5 Hz, 1H), 3.59–3.50 (m, 1H), 3.04–2.92 (m, 1H), 2.79 (dd, J = 16.3, 5.2 Hz, 1H), 2.66 (s, 3H), 2.20–2.07 (m, 1H), 1.18 (d, J = 6.7 Hz, 3H), 1.12 (d, J = 6.8 Hz, 3H); ¹³C NMR (125 MHz, CDCl₃): δ 135.9, 132.4, 126.7, 122.6, 120.0, 118.4, 111.1, 107.9, 58.8, 40.1, 39.9, 33.7, 20.3, 20.2, 20.1; HRMS (ESI) calcd for C₁₅H₂₁N₂O₂S m/z 293.1318 [M + H]⁺, found 293.1288.

1-Isopropyl-2-tosyl-2,3,4,9-tetrahydro-1*H***-pyrido**[**3,4-***b*]**indole** (5**s**). White solid (363 mg, 91% yield), mp: 211–215 °C; ¹H

NMR (500 MHz, CDCl₃): δ 7.77 (s, 1H), 7.53 (d, J = 8.2 Hz, 2H), 7.29 (d, J = 8.1 Hz, 1H), 7.24 (d, J = 7.9 Hz, 1H), 7.13 (t, J = 7.6 Hz, 1H), 7.02 (t, J = 7.8 Hz, 1H), 6.97 (d, J = 8.2 Hz, 2H), 4.79 (d, J = 7.8 Hz, 1H), 4.13 (dd, J = 15.0, 6.5 Hz, 1H), 3.52–3.43 (m, 1H), 2.44 (dd, J = 15.9, 4.8 Hz, 1H), 2.32–2.23 (m, 1H), 2.19 (s, 3H), 2.17–2.08 (m, 1H), 1.17 (d, J = 6.7 Hz, 3H), 1.11 (d, J = 6.8 Hz, 3H); 13 C NMR (125 MHz, CDCl₃): δ 143.2, 138.0, 135.8, 132.3, 129.3, 126.8, 126.7, 122.1, 119.5, 118.1, 110.8, 108.2, 59.1, 40.2, 34.1, 21.4, 20.2, 20.1, 19.4; HRMS (ESI) calcd for $C_{21}H_{25}N_2O_2S$ m/z 369.1631 [M + H]⁺, found 369.1647.

1-Phenyl-2-tosyl-2,3,4,9-tetrahydro-1*H***-pyrido**[3,4-*b*]indole (5t). White solid (393 mg, 93% yield), mp: 157–162 °C; ¹H NMR (400 MHz, DMSO- d_6): δ 10.82 (s, 1H), 7.63 (d, J = 8.1 Hz, 2H), 7.38–7.13 (m, 9H), 7.05 (t, J = 7.5 Hz, 1H), 6.94 (t, J = 7.4 Hz, 1H), 6.24 (s, 1H), 3.95 (dd, J = 14.5, 5.6 Hz, 1H), 3.26–3.14 (m, 1H), 2.61 (dd, J = 15.7, 4.1 Hz, 1H), 2.48–2.35 (m, 1H), 2.22 (s, 3H); ¹³C NMR (100 MHz, DMSO- d_6): δ 143.0, 139.8, 137.6, 136.0, 130.6, 129.5, 128.4, 128.0, 128.0, 126.5, 126.0, 121.3, 118.5, 117.8, 111.2, 107.6, 55.3, 39.1, 20.8, 19.5; HRMS (ESI) calcd for $C_{24}H_{23}N_2O_2S$ m/z 403.1475 [M + H]⁺, found 403.1489.

Methyl (*S*)-2-tosyl-2,3,4,9-tetrahydro-1*H*-pyrido[3,4-*b*]indole-3-carboxylate (5u). ^{23α} White solid (386 mg, 94% yield), mp: 138–143 °C; ¹H NMR (400 MHz, CDCl₃): δ 8.06 (br s, 1H), 7.71 (d, J = 8.3 Hz, 2H), 7.41 (d, J = 7.6 Hz, 1H), 7.28–7.22 (m, 3H), 7.16–7.04 (m, 2H), 5.15 (d, J = 6.4 Hz, 1H), 4.82 (d, J = 15.0 Hz, 1H), 4.64 (d, J = 15.1 Hz, 1H), 3.43 (s, 3H), 3.30 (d, J = 15.7 Hz, 1H), 3.06 (dd, J = 15.7, 6.6 Hz, 1H), 2.37 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 171.0, 143.8, 136.3, 129.7, 128.4, 127.2, 126.7, 122.2, 119.7, 118.1, 111.1, 105.6, 54.1, 52.4, 41.0, 24.6, 21.6; HRMS (ESI) calcd for C₂₀H₂₁N₂O₄S m/z 385.1217 [M + H]⁺, found 385.1219.

General procedure for the synthesis of 5w, 5x, 5y and 5z. The diastereomeric mixtures of THBC esters (5w and 5x) and (5y and 5z) were synthesized according to the procedures given in previous literature reports. These diastereomeric mixtures were separated by silica gel column chromatography using EtOAc/hexane (20–30%) prior to use for TCCA-mediated oxidative rearrangement.

Methyl (1*R*,3*S*)-2-benzyl-1-methyl-2,3,4,9-tetrahydro-1*H*-pyrido[3,4-*b*]indole-3-carboxylate (5x)^{22c}. Brown oil (312 mg, 92% yield); ¹H NMR (400 MHz, CDCl₃): δ 7.70 (s, 1H), 7.51 (d, J = 7.3 Hz, 1H), 7.43 (d, J = 7.4 Hz, 2H), 7.34–7.25 (m, 3H), 7.23 (t, J = 7.3 Hz, 1H), 7.17–7.06 (m, 2H), 4.24 (q, J = 6.8 Hz, 1H), 4.01 (d, J = 4.5 Hz, 2H), 3.90 (t, J = 5.7 Hz, 1H), 3.62 (s, 3H), 3.26 (dd, J = 15.2, 6.1 Hz, 1H), 2.99 (dd, J = 17.2, 5.7 Hz, 1H), 1.34 (d, J = 6.9 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 174.2, 140.0, 136.1, 135.5, 128.6, 128.3, 127.2, 127.0, 121.7, 119.6, 118.3, 110.8, 106.6, 60.0, 54.9, 52.9, 51.8, 22.3, 18.3; HRMS (ESI) calcd for $C_{21}H_{23}N_2O_2$ m/z 335.1754 [M + H]⁺, found 335.1762.

Methyl (1*R*,3*S*)-2-benzyl-1-ethyl-2,3,4,9-tetrahydro-1*H*-pyrido [3,4-*b*]indole-3-carboxylate (5z). ^{23*d*,23*e*} Brown oil (368 mg, 89% yield); ¹H NMR (400 MHz, CDCl₃): δ 7.74 (s, 1H), 7.52 (d, *J* = 7.5 Hz, 1H), 7.44 (d, *J* = 7.3 Hz, 2H), 7.34–7.29 (m, 2H), 7.29–7.22 (m, 2H), 7.16–7.07 (m, 2H), 3.93 (d, *J* = 1.5 Hz, 2H), 3.86–3.80 (m, 2H), 3.60 (s, 3H), 3.23 (dd, *J* = 15.7, 4.5 Hz, 1H), 2.97 (dd, *J* = 15.7, 7.0 Hz, 1H), 1.71–1.61 (m, 2H), 1.00 (t, *J* = 7.3 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 174.2, 139.6, 136.1, 134.6, 128.9,

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128.3, 127.2, 127.1, 121.6, 119.5, 118.3, 110.8, 106.5, 59.5, 59.1, 58.4, 51.7, 27.0, 20.4, 11.1; HRMS (ESI) calcd for $C_{22}H_{25}N_2O_2\ m/z$ 349.1911 [M + H]⁺, found 349.1915.

General procedure for TCCA mediated oxidative rearrangement reaction. To a stirred homogeneous solution of THBCs (1.0 equiv.) in THF/H₂O (1:1), TCCA (0.35 equiv.) was added at room temperature in one batch. The resulting reaction mixture was stirred at room temperature for 30 min. After the reaction was completed, as monitored by TLC analysis (EtOAc: hexane or MeOH: CH₂Cl₂ solvent systems), it was quenched by addition of saturated NaHCO₃ and extracted with EtOAc three times. The combined organic phases were washed with water, dried over Na₂SO₄, filtered, and evaporated under reduced pressure. The resulting crude product was purified by flash column chromatography (EtOAc/hexane = 1:10 to 1:1) to provide the pure spirooxindole. Note: This reaction was performed at 0 °C in the case of THBCs 5c and 5t.

1'-Tosylspiro[indoline-3,3'-pyrrolidin]-2-one (6a).^{13f} White solid (98 mg, 99% yield), mp: 158–160 °C; ¹H NMR (400 MHz, CDCl₃): δ 8.63 (s, 1H), 7.77 (d, J = 8.2 Hz, 2H), 7.37 (d, J = 8.1 Hz, 2H), 7.21 (t, J = 7.7 Hz, 1H), 7.03 (d, J = 7.0 Hz, 1H), 6.96 (t, J = 7.4 Hz, 1H), 6.88 (d, J = 7.7 Hz, 1H), 3.74–3.68 (m, 1H), 3.57 (d, J = 9.6 Hz, 1H), 3.58–3.50 (m, 1H) 3.47 (d, J = 9.8 Hz, 1H), 2.46 (s, 3H), 2.35–2.24 (m, 1H), 2.10–2.00 (m, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 179.3, 144.0, 139.9, 133.4, 132.8, 129.9, 128.6, 127.8, 123.2, 123.2, 110.1, 56.1, 53.0, 47.4, 36.4, 21.7; HRMS (ESI) calcd for C₁₈H₁₉N₂O₃S m/z 343.1111 [M + H]⁺, found 343.1123.

1'-(Methylsulfonyl)spiro[indoline-3,3'-pyrrolidin]-2-one (6b). White solid (90 mg, 94% yield), mp: 217–219 °C; ¹H NMR (500 MHz, DMSO- d_6): δ 10.54 (s, 1H), 7.38 (d, J = 7.4 Hz, 1H), 7.23 (t, J = 7.7, 1H), 7.01 (t, J = 7.5 Hz, 1H), 6.88 (d, J = 7.7 Hz, 1H), 3.74–3.65 (m, 1H), 3.63–3.59 (m, 1H), 3.56 (d, J = 10.3 Hz, 1H), 3.44 (d, J = 10.3 Hz, 1H), 3.03 (s, 3H), 2.28–2.23 (m, 1H), 2.20–2.14 (m, 1H); 13 C NMR (125 MHz, DMSO- d_6): δ 179.1, 141.5, 131.4, 128.4, 123.0, 121.87, 109.4, 54.9, 52.3, 47.0, 35.9, 34.3; HRMS (ESI) calcd for $C_{12}H_{15}N_2O_3S$ m/z 267.0798 [M + H] $^+$, found 267.0846.

1'-Acetylspiro[indoline-3,3'-pyrrolidin]-2-one (6e). This reaction was performed at 0 °C. Pale yellow solid (101 mg, 82% yield, found as 50 : 50 rotamers CDCl₃), mp: 75–78 °C; Th NMR (500 MHz, CDCl₃): δ 9.38 and 9.22 (s, 1H), 7.29–7.20 (m, 1H), 7.19–7.10 (s, 1H), 7.09–7.00 (s, 1H), 6.95 (ddd, J=21.6, 14.1, 8.0 Hz, 1H), 4.05–3.95 (m, 1H), 3.93–3.80 (m, 2H), 3.68 (dd, J=65.0, 15.0 Hz, 1H), 2.54–2.40 (m, 1H), 2.29–2.21 (m, 1H), 2.19 and 2.10 (s, 3H); The NMR (125 MHz, CDCl₃): δ 180.5, 179.4, 169.7, 169.5, 140.7, 140.4, 132.5, 131.7, 128.8, 128.7, 123.1, 122.9, 122.7, 110.4, 110.3, 55.5, 53.9, 53.6, 51.9, 46.7, 45.2, 36.5, 35.1, 22.6, 22.4; HRMS (ESI) calcd for C₁₅H₁₅N₂O₂ m/z 231.1128 [M + H]⁺, found 231.1108.

1'-Benzoylspiro[indoline-3,3'-pyrrolidin]-2-one (6d). White solid (95 mg, 89% yield, found as 62 : 38 rotamers in CDCl₃), mp: 83–85 °C; ¹H NMR (500 MHz, CDCl₃): δ 8.07 and 8.03 (s, 1H), 7.63 (d, J=7.0 Hz, 1H), 7.52 (d, J=7.5 Hz, 1H), 7.47–7.43 (m, 1H), 7.41–7.32 (m, 2H), 7.27–7.20 (s, 2H), 7.17–7.01 (m, 1H), 6.91 (dd, J=22.2, 7.7 Hz, 1H), 4.25–4.02 (m, 2H), 3.99–3.56 (m, 2H), 2.66–2.31 (m, 1H), 2.27–2.10 (m, 1H); ¹³C NMR (125 MHz, CDCl₃): δ 180.6, 178.8, 170.2, 170.1, 140.6, 140.1, 136.4, 132.4,

130.3, 128.8, 128.7, 128.5, 127.3, 123.2, 122.7, 110.3, 57.6, 54.2, 53.6, 51.9, 48.7, 45.6, 36.9, 35.0; HRMS (ESI) calcd for $C_{18}H_{17}N_2O_2S_2$ m/z 293.1285 $\left[M+H\right]^+$, found 293.1253.

Ethyl 2-oxospiro[indoline-3,3'-pyrrolidine]-1'-carboxylate (6e). Colorless oil (85 mg, 83% yield, found as 56 : 44 rotamers in CDCl₃); ¹H NMR (500 MHz, CDCl₃): δ 9.11 and 9.04 (s, 1H), 7.27–7.22 (m, 1H), 7.17 (t, J = 5.0 Hz, 1H), 7.05 (t, J = 7.4 Hz, 1H), 6.97–6.90 (m, 1H), 4.23–4.12 (m, 2H), 3.95–3.87 (m, 1H), 3.85–3.74 (m, 2H), 3.64 (dd, J = 18.0, 9.1 Hz, 1H), 2.48–2.40 (m, 1H), 2.16–2.07 (m, 1H), 1.32 (t, J = 7.0 Hz, 1.3H), 1.24 (t, J = 7.0 Hz, 1.7H); ¹³C NMR (125 MHz, CDCl₃): δ 180.2, 179.9, 155.2, 155.1, 140.3, 140.2, 132.9, 132.5, 128.6, 123.4, 123.1, 123.1, 122.8, 111.2, 110.2, 61.6, 61.5, 54.3, 54.1, 53.4, 52.4, 45.7, 45.3, 36.5, 35.6, 14.9, 14.8; HRMS (ESI) calcd for C₁₄H₁₇N₂O₃ m/z 261.1234 [M + H]⁺, found 269.1246.

tert-Butyl 2-oxospiro[indoline-3,3′-pyrrolidine]-1′-carboxylate (6f). ^{13f} White solid (102 mg, 88% yield, found as 51 : 49 rotamers in DMSO- d_6), mp: 123–125 °C; ¹H NMR (500 MHz, DMSO- d_6): δ 10.50 (s, 1H), 7.25–7.19 (m, 2H), 6.97 (t, J = 6.5 Hz, 1H), 6.86 (d, J = 7.6 Hz, 1H), 3.67–3.57 (m, 2H), 3.51–3.43 (m, 2H), 2.18–2.09 (m, 2H), 1.44 and 1.39 (s, 9H); ¹³C NMR (125 MHz, DMSO- d_6): δ 179.3, 153.5, 141.7, 131.8, 128.2, 122.8, 121.5, 109.4, 78.6, 53.8, 53.5, 52.2, 51.3, 45.0, 44.9, 35.5, 34.6, 28.1, 28.0; HRMS (ESI) calcd for C₁₆H₂₀N₂O₃Na m/z 311.1366 [M + Na]⁺, found 311.1416.

Benzyl 2-oxospiro[indoline-3,3'-pyrrolidine]-1'-carboxylate (6g). Colourless oil (75 mg, 88% yield, found as 52 : 48 rotamers in CDCl₃) ¹H NMR (500 MHz, CDCl₃): δ 8.65 and 8.59 (s, 1H), 7.43–7.37 (m, 2H), 7.33–7.32 (m, 3H), 7.24 (t, J = 12.8 Hz, 1H), 7.15 (dd, J = 15.2, 7.4 Hz, 1H), 7.05–7.03 (m, 1H), 6.93 (d, J = 7.8 Hz, 1H), 5.21 (d, J = 2.8 Hz, 1H), 5.16 (s, 1H), 3.97–3.90 (m, 1H), 3.87–3.78 (m, 2H), 3.69 (dd, J = 23.0, 11.1 Hz, 1H), 2.47–2.41 (dt, J = 12.7, 8.1 Hz, 1H), 2.16–2.08 (m, 1H); ¹³C NMR (125 MHz, CDCl₃): δ 179.7, 154.8, 140.1, 136.9, 136.7, 132.7, 132.3, 128.6, 128.6, 128.1, 128.1, 128.0, 127.9, 123.1, 122.9, 110.1, 67.2, 54.5, 54.2, 53.3, 52.4, 45.8, 45.4, 36.5, 35.6; HRMS (ESI) calcd for C₁₉H₁₉N₂O₃ m/z 323.1390 [M + H]⁺, found 323.1349.

(±)-Coerulescine (1). ^{13e,13f} Off-white solid (1.21 g, 93% yield), mp: 110–114 °C; ¹H NMR (400 MHz, CDCl₃): δ 9.24 (s, 1H), 7.42 (d, J = 8.0 Hz, 1H), 7.22 (t, J = 8.0 Hz, 1H), 7.06 (t, J = 8.0 Hz, 1H), 6.95 (d, J = 8.0 Hz, 1H), 3.06–3.00 (m, 1H), 2.92 (dd, J = 25.1, 9.3 Hz, 2H), 2.85 (dd, J = 15.1, 6.8 Hz, 1H), 2.50 (s, 3H), 2.49–2.41 (m, 1H), 2.19–2.09 (m, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 183.5, 140.4, 136.3, 127.8, 123.3, 122.8, 109.8, 66.4, 56.9, 53.8, 41.9, 38.0; HRMS (ESI) calcd for $C_{12}H_{15}N_2O$ m/z 203.1179 [M + H]⁺, found 203.1175.

1-Benzyl-1'-tosylspiro[indoline-3,3'-pyrrolidin]-2-one (6h). ^{23a} Colorless gummy oil (78 mg, 93% yield); ¹H NMR (400 MHz, CDCl₃): δ 7.78 (d, J = 8.2 Hz, 2H), 7.37 (d, J = 8.0 Hz, 2H), 7.33–

7.25 (m, 3H), 7.24–7.08 (m, 4H), 6.97 (td, J=7.6, 0.8 Hz, 1H), 6.72 (d, J=7.8 Hz, 1H), 4.87 (d, J=1.3 Hz, 2H), 3.81–3.72 (m, 1H), 3.63–3.47 (m, 3H), 2.47 (s, 3H), 2.39–2.30 (m, 1H), 2.13–2.01 (m, 1H); 13 C NMR (100 MHz, CDCl₃): δ 177.0, 144.0, 141.8, 135.6, 133.4, 132.5, 130.0, 128.9, 128.6, 127.8, 127.2, 123.3, 123.0, 109.3, 56.3, 52.6, 47.4, 44.0, 36.4, 21.7; HRMS (ESI) calcd for $C_{25}H_{25}N_2O_3$ S m/z 433.1580 [M + H]⁺, found 433.1588.

1-Methyl-1'-tosylspiro[indoline-3,3'-pyrrolidin]-2-one (6i). ^{24a} Off-white solid (76 mg, 95% yield), 147–151 °C; ¹H NMR (400 MHz, CDCl₃): δ 7.76 (d, J = 8.2 Hz, 2H), 7.36 (d, J = 8.1 Hz, 2H), 7.32–7.26 (m, 1H), 7.10 (d, J = 7.4 Hz, 1H), 7.01 (t, J = 7.5 Hz, 1H), 6.84 (d, J = 7.8 Hz, 1H), 3.78–3.70 (m, 1H), 3.57–3.47 (m, 2H), 3.42 (d, J = 9.6 Hz, 1H), 3.18 (s, 3H), 2.47 (s, 3H), 2.33–2.23 (m, 1H), 2.05–1.97 (m, 1H); ¹³C NMR (100 MHz, CDCl₃): δ 176.8, 143.9, 142.7, 133.4, 132.6, 129.9, 128.6, 127.8, 123.3, 122.9, 108.3, 56.2, 52.6, 47.4, 36.3, 26.5, 21.7; HRMS (ESI) calcd for C₁₉H₂₁N₂O₃S m/z 357.1267 [M + H]⁺, found 357.1273.

Ethyl 2-oxo-1-tosylspiro[indoline-3,3'-pyrrolidine]-1'-carboxylate (6j). Pale yellow oil (63 mg, 78% yield); ¹H NMR (400 MHz, CDCl₃ + DMSO- d_6): δ 7.94 (d, J = 7.6 Hz, 2H), 7.44 (d, J = 8.4 Hz, 1H), 7.30 (d, J = 7.4 Hz, 1H), 7.24 (d, J = 7.7 Hz, 3H), 7.02 (t, J = 7.4 Hz, 1H), 4.20–4.10 (m, 3H), 3.98–3.51 (m, 2H), 3.03–2.99 (m, 1H), 2.37 (s, 3H), 2.27–1.99 (m, 2H), 1.30–1.24 (m, 3H); ¹³C NMR (100 MHz, CDCl₃ + DMSO- d_6): δ 173.0, 155.5, 143.2, 137.2, 129.7, 128.9, 127.7, 123.7, 123.0, 122.1, 113.2, 95.4, 75.7, 61.0, 60.8, 48.4, 32.4, 21.1, 14.3; HRMS (ESI) calcd for C₂₁H₂₂N₂O₅S m/z 414.1249 [M]⁺, found 414.1255.

tert-Butyl 2-oxo-1'-tosylspiro[indoline-3,3'-pyrrolidine]-1-carboxylate (6k). Colorless oil (84 mg, 82% yield); ¹H NMR (400 MHz, CDCl₃): δ 7.62 (d, J = 8.2 Hz, 2H), 7.53 (d, J = 8.0 Hz, 1H), 7.31–7.23 (m, 4H), 7.04 (t, J = 7.5 Hz, 1H), 3.92 (d, J = 13.3 Hz, 1H), 3.51–3.46 (m, 1H), 3.30 (d, J = 13.3 Hz, 1H), 2.80–2.72 (m, 1H), 2.40 (s, 3H), 2.26–2.14 (m, 2H), 1.63 (s, 9H); ¹³C NMR (100 MHz, CDCl₃): δ 174.0, 152.9, 143.6, 140.3, 135.0, 130.6, 130.3, 129.8, 127.4, 123.7, 123.3, 115.3, 91.1, 83.8, 75.7, 50.0, 42.0, 32.2, 28.5, 21.6; HRMS (ESI) calcd for C₂₃H₂₇N₂O₅S m/ z 443.1635 [M + H]⁺, found 443.1643.

2'-Methyl-1'-(methylsulfonyl)spiro[indoline-3,3'-pyrrolidin]-2-one (6l). White solid (58 mg, 87% yield, dr 1 : 1.6), mp: 140–142 °C; ¹H NMR (500 MHz, DMSO- d_6): major isomer: δ 10.52 (s, 1H), 7.43 (d, J = 7.4 Hz, 1H), 7.22 (t, J = 8.3 Hz, 1H), 7.01 (t, J = 7.5 Hz, 1H), 6.84 (d, J = 7.7 Hz, 1H), 3.89–3.75 (m, 2H), 3.69–3.62 (m, 1H), 3.08 (s, 3H), 2.24–2.16 (m, 1H), 2.10 (dd, J = 6.4, 2.1 Hz, 1H), 1.07 (d, J = 6.4 Hz, 3H); minor isomer: δ 10.58 (s, 1H), 7.33 (d, J = 7.4 Hz, 1H), 7.22 (t, J = 8.3 Hz, 1H), 7.01 (t, J = 7.5 Hz, 1H), 6.88 (d, J = 7.7 Hz, 1H), 3.77–3.70 (m, 2H), 3.60 (dd, J = 7.0, 3.1 Hz, 1H), 3.01 (s, 3H), 2.14 (dd, J = 12.9, 6.1 Hz, 1H), 2.07 (dd, J = 6.3, 1.9 Hz, 1H), 1.02 (d, J = 6.5 Hz, 3H); ¹³C NMR (125 MHz, CDCl₃): δ 180.3, 178.9, 140.7, 130.0, 129.0, 128.7, 125.5, 123.2, 123.0, 122.8, 110.3, 110.0, 63.1, 61.4, 57.8, 57.4, 47.3, 47.2, 40.3, 35.1, 34.8, 34.3, 20.0, 15.8; HRMS (ESI) calcd for $C_{13}H_{17}N_2O_3S$ m/z 281.0954 $[M+H]^+$, found 281.0968.

2'-Methyl-1'-tosylspiro[indoline-3,3'-pyrrolidin]-2-one (6m). White solid (92 mg, 89% yield, dr 1 : 4), mp: 195–199 °C; 1 H NMR (400 MHz, CDCl₃): major isomer: δ 8.58 (s, 1H), 7.84 (d, J = 8.2 Hz, 2H), 7.40 (d, J = 8.0 Hz, 2H), 7.19 (t, J = 7.7 Hz, 1H), 6.95–6.85 (m, 2H), 6.57 (d, J = 7.4 Hz, 1H), 4.08–4.00 (m, 1H), 3.93–

3.85 (m, 1H), 3.80–3.68 (m, 1H), 2.49 (s, 3H), 2.21–2.13 (m, 1H), 1.80–1.74 (m, 1H), 1.33 (d, J=6.5 Hz, 3H); minor isomer: δ 8.44 (s, 1H), 7.79 (d, J=8.2 Hz, 2H), 7.40 (d, J=8.0 Hz, 2H), 7.09 (t, J=7.6 Hz, 1H), 6.95–6.85 (m, 2H), 6.57 (d, J=7.4 Hz, 1H), 4.08–4.00 (m, 1H), 3.93–3.85 (m, 1H), 3.80–3.68 (m, 1H), 2.46 (s, 3H), 2.21–2.13 (m, 1H), 1.80–1.74 (m, 1H), 1.14 (d, J=6.4 Hz, 3H); 13 C NMR (100 MHz, CDCl₃): δ 178.9, 178.5, 143.9, 143.8, 140.6, 140.2, 135.7, 133.5, 130.6, 129.9, 128.8, 127.9, 127.6, 125.2, 123.0, 122.9, 122.6, 110.2, 110.0, 63.7, 62.3, 58.4, 57.8, 48.4, 48.0, 34.8, 33.8, 23.9, 21.7, 17.5, 17.3; HRMS (ESI) calcd for $C_{19}H_{21}N_2O_3S$ m/z 357.1267 [M + H]⁺, found 357.1269.

2′-Ethyl-1′-(methylsulfonyl)spiro[indoline-3,3′-pyrrolidin]-2-one (6n). White solid (82 mg, 85% yield), mp: 158–160 °C; $^1\mathrm{H}$ NMR (500 MHz, DMSO- d_6): δ 10.58 (s, 1H), 7.46 (d, J=7.4 Hz, 1H), 7.19 (d, J=7.7 Hz, 1H), 7.00 (t, J=7.5 Hz, 1H), 6.83 (d, J=7.7 Hz, 1H), 3.83–3.76 (m, 2H), 3.69–3.63 (m, 1H), 3.12 (s, 3H), 2.18–2.09 (m, 1H), 2.09–2.02 (m, 1H), 2.00–1.88 (m, 1H), 1.83–1.72 (m, 1H), 0.39 (t, J=7.6 Hz, 4H); $^{13}\mathrm{C}$ NMR (125 MHz, DMSO- d_6): δ 178.1, 141.2, 131.5, 128.2, 122.9, 121.8, 109.3, 67.7, 56.6, 47.4, 36.8, 34.8, 25.7, 10.1; HRMS (ESI) calcd for $\mathrm{C}_{14}\mathrm{H}_{19}\mathrm{N}_2\mathrm{O}_3\mathrm{S}$ m/z 295.1111 [M + H] $^+$, found 295.1099.

2'-Ethyl-1'-tosylspiro[indoline-3,3'-pyrrolidin]-2-one (6ο). White solid (74 mg, 87% yield), mp: 208–210 °C; 1 H NMR (500 MHz, CDCl₃): δ 8.07 (s, 1H), 7.85 (d, J = 8.2 Hz, 2H), 7.41 (d, J = 8.3 Hz, 2H), 7.17 (t, J = 8.1 Hz, 1H), 6.89 (t, J = 7.6 Hz, 1H), 6.83 (d, J = 7.8 Hz, 1H), 6.56 (d, J = 7.4 Hz, 1H), 3.97 (ddd, J = 11.1, 10.1, 5.8 Hz, 1H), 3.86 (dd, J = 9.9, 4.4 Hz, 1H), 3.79–3.72 (m, 1H), 2.49 (s, 3H), 2.12–2.05 (m, 2H), 2.04–1.95 (m, 1H), 1.73–1.66 (s, 1H), 0.60 (t, J = 7.5 Hz, 3H); 13 C NMR (125 MHz, CDCl₃): δ 178.8, 143.9, 139.8, 135.9, 132.4, 130.0, 128.5, 127.8, 123.0, 122.6, 109.8, 69.4, 57.0, 48.1, 37.4, 26.3, 21.7, 10.8; HRMS (ESI) calcd for C₁₅H₁₈NOS₂ m/z 371.1424 [M + H]⁺, found 371.1439.

2'-Isopropyl-1'-(methylsulfonyl)spiro[indoline-3,3'-pyrrolidin]-2-one (6p). Colourless oil (83 mg, 80% yield); $^1\mathrm{H}$ NMR (500 MHz, CDCl₃): δ 7.62 and 7.60 (d, J = 9.4 Hz, 1H), 7.41 (d, J = 7.5 Hz, 1H), 7.22 (t, J = 7.7 Hz, 1H), 7.09 (t, J = 7.6 Hz, 1H), 6.85 (d, J = 7.7 Hz, 1H), 4.21 (d, J = 9.3 Hz, 1H), 3.91–3.84 (m, 1H), 3.81–3.73 (m, 1H), 3.08 (s, 3H), 3.92–3.84 (m, 1H), 2.35–2.21 (m, 2H), 1.02 (d, J = 6.8 Hz, 3H), 0.69 (d, J = 6.6 Hz, 3H); $^{13}\mathrm{C}$ NMR (125 MHz, CDCl₃): δ 179.3, 138.9, 133.7, 128.2, 123.4, 123.1, 109.4, 73.2, 56.4, 48.4, 41.0, 40.8, 30.4, 20.9, 19.7; HRMS (ESI) calcd for $\mathrm{C_{15}H_{21}N_2O_3S}$ m/z 309.1267 [M + H]⁺, found 309.1231.

2′-**Isopropyl-1**′-**tosylspiro**[**indoline-3,3**′-**pyrrolidin**]-**2-one** (**6q**). White solid (86 mg, 85% yield), mp: 216–219 °C; ¹H NMR (500 MHz, CDCl₃): δ 7.63 (d, J = 8.3 Hz, 2H), 7.55 (d, J = 7.7 Hz, 1H), 7.37 (t, J = 7.6 Hz, 1H), 7.29 (d, J = 6.8 Hz, 1H), 7.22 (t, J = 7.4 Hz, 1H), 7.07 (d, J = 8.4 Hz, 2H), 4.72 (d, J = 11.3 Hz, 1H), 3.95 (dd, J = 15.4, 4.2 Hz, 1H), 3.73 (ddd, J = 15.1, 12.8, 2.1 Hz, 1H), 2.85–2.75 (m, 1H), 2.31 (d, J = 14.7 Hz, 1H), 2.20 (s, 3H), 1.25–1.16 (m, 1H), 1.15 (d, J = 6.6 Hz, 3H), 1.02 (d, J = 6.6 Hz, 3H); ¹³C NMR (125 MHz, CDCl₃): δ 174.6, 151.3, 143.4, 139.7, 137.3, 130.0, 129.4, 127.3, 127.0, 122.1, 121.3, 67.6, 65.7, 39.0, 37.7, 28.0, 21.2, 20.2, 19.3; HRMS (ESI) calcd for C₂₁H₂₅N₂O₃S m/ z 385.1580 [M + H]⁺, found 385.1587.

2'-Phenyl-1'-tosylspiro[indoline-3,3'-pyrrolidin]-2-one (6r). This reaction was performed at 0 °C. White solid (52 mg, 45%)

419.1428.

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yield, dr 1 : 7.9), mp: 169–173 °C; ¹H NMR (400 MHz, DMSO- d_6): major isomer: δ 10.43 (s, 1H), 7.72 (d, J = 8.0 Hz, 2H), 7.47 (d, J = 7.9 Hz, 2H), 7.23–6.92 (m, 6H), 6.66 (dd, J = 14.1, 7.4 Hz, 2H), 6.55 (d, J = 7.4 Hz, 1H), 4.81 (s, 1H), 4.00–3.92 (m, 1H), 3.90–3.82 (m, 1H), 2.45 (s, 3H), 2.07–1.96 (m, 1H), 1.92–1.80 (m, 1H); minor isomer: δ 10.55 (s, 1H), 7.84 (d, J = 8.0 Hz, 2H), 7.57 (d, J

6.55 (d, J = 7.4 Hz, 1H), 4.81 (s, 1H), 4.00–3.92 (m, 1H), 3.90–3.82 (m, 1H), 2.45 (s, 3H), 2.07–1.96 (m, 1H), 1.92–1.80 (m, 1H); minor isomer: δ 10.55 (s, 1H), 7.84 (d, J = 8.0 Hz, 2H), 7.57 (d, J = 7.9 Hz, 2H), 7.23–6.92 (m, 6H), 6.66 (dd, J = 14.1, 7.4 Hz, 2H), 6.55 (d, J = 7.4 Hz, 1H), 4.83 (s, 1H), 4.28–4.00 (m, 2H), 2.45 (s, 3H), 2.19–2.08 (m, 2H); 13 C NMR (100 MHz, DMSO- d_6): δ 177.8, 143.5, 141.3, 138.7, 133.7, 129.7, 128.4, 128.1, 127.7, 127.4, 127.2, 126.8, 125.0, 121.0, 109.2, 69.1, 58.5, 47.9, 34.5, 21.1; HRMS (ESI) calcd for C₂₄H₂₃N₂O₃S m/z 419.1424 [M + H]⁺, found

N-(2-(2-Benzoyl-1H-indol-3-yl)ethyl)-4-

methylbenzenesulfonamide (6r'). ^{24c} Yellow oil (110 mg, 50% yield); ¹H NMR (400 MHz, DMSO- d_6): δ 11.45 (s, 1H), 7.75–7.56 (m, 9H), 7.43 (d, J=8.1 Hz, 1H), 7.33–7.25 (m, 3H), 7.09 (t, J=7.2 Hz, 1H), 2.96 (s, 4H), 2.34 (s, 3H); ¹³C NMR (100 MHz, DMSO- d_6): δ 188.4, 142.4, 138.7, 137.6, 136.7, 132.3, 131.4, 129.5, 128.9, 128.6, 127.2, 126.4, 125.3, 120.3, 120.0, 119.9, 112.8, 43.3, 25.3, 20.9; HRMS (ESI) calcd for C₂₄H₂₃N₂O₃S m/z 419.1424 [M + H]⁺, found 419.1432.

Methyl 2-oxo-1'-tosylspiro[indoline-3,3'-pyrrolidine]-5'-carboxylate (6s). Light brown oil (87 mg, 95% yield, dr 1 : 1.9); 1 H NMR (400 MHz, CDCl₃): δ 8.62 and 8.40 (s, 1H), 7.81 (d, J = 8.0 Hz, 2H), 7.53 (d, J = 7.4 Hz, 0.5H), 7.34 (d, J = 8.0 Hz, 2H), 7.26–7.18 (m, 1H), 7.06 (t, J = 7.6 Hz, 0.5H), 6.92–6.85 (m, 1.5H), 6.77 (d, J = 7.5 Hz, 0.5H), 4.79 (t, J = 8.1 Hz, 0.66H), 4.57 (t, J = 7.9 Hz, 0.34H), 3.83 and 3.72 (s, 3H), 3.82–3.57 (m, 2H), 2.68–2.60 (m, 1H), 2.46 and 2.43 (s, 3H), 2.42–2.34 (m, 1H); 13 C NMR (100 MHz, CDCl₃): δ 179.4, 177.6, 172.0, 171.5, 144.2, 144.0, 140.2, 139.8, 136.1, 133.9, 132.3, 130.5, 129.8, 129.8, 129.0, 128.8, 128.2, 127.6, 124.1, 123.4, 123.2, 123.0, 110.3, 110.1, 60.9, 60.5, 56.8, 56.5, 53.0, 52.8, 52.7, 52.1, 41.5, 40.6, 21.7, 21.7; HRMS (ESI) calcd for C₂₀H₂₁N₂O₅S m/z 401.1166 [M + H]⁺, found 401.1172.

1'-(tert-Butyl) 5'-methyl 2-oxospiro[indoline-3,3'-pyrrolidine]-1',5'-dicarboxylate (6t).^{23b} Colorless oil (98 mg, 93% yield, dr 1 : 3); 1 H NMR (400 MHz, CDCl₃) Major isomer: δ 8.71 (m, 1H), 7.35–7.21 (m, 1H), 7.16–7.02 (m, 2H), 6.99–6.88 (m, 1H), 4.88–4.71 (m, 1H), 3.94–3.68 (m, 2H), 3.79 (s, 3H), 2.74–2.48 (m, 2H), 1.46 (s, 9H); minor isomer: δ 7.63 (m, 1H), 7.35–7.21 (m, 1H), 7.16–7.02 (m, 2H), 6.99–6.88 (m, 1H), 4.66–4.59 (m, 1H), 3.94–3.68 (m, 2H), 3.80 (s, 3H), 2.45–2.26 (m, 2H), 1.48 (s, 9H); 13 C NMR (100 MHz, CDCl₃): δ 178.0, 173.4, 172.5, 172.2, 154.4, 153.5, 140.8, 139.8, 133.3, 133.1, 129.4, 129.0, 128.8, 123.4, 123.2, 123.1, 122.5, 110.4, 110.1, 81.0, 80.7, 59.1, 58.7, 55.4, 54.9, 54.5, 52.4, 52.3, 41.0, 40.6, 40.3, 39.7, 28.4, 28.3; HRMS (ESI) calcd for C₁₈H₂₃N₂O₅ m/z 347.1601 [M + H] $^+$, found 347.1613.

Methyl 1'-benzyl-2'-methyl-2-oxospiro[indoline-3,3'-pyrrolidine]-5'-carboxylate (6u). Pale yellow oil (93 mg, 95% yield, dr 1.7 : 1); ¹H NMR (400 MHz, CDCl₃): δ 8.96–8.73 (m, 1H), 7.80 (d, J = 7.4 Hz, 1H), 7.41–7.29 (m, 4H), 7.28–7.18 (m, 2H), 7.14–7.03 (m, 1H), 6.95–6.88 (m, 1H), 4.03–3.80 (m, 1H), 3.79–3.60 (m, 3H), 3.50 (s, 2H), 3.17 (dd, J = 12.1, 6.0 Hz, 1H), 2.76–2.66 (m, 0.6H), 2.51–2.45 (m, 0.4H), 2.30 (dd, J = 13.3, 9.2 Hz, 0.4H), 2.11 (dd, J = 13.4, 6.0 Hz, 0.6H), 0.75 and 0.73 (d, J = 10.3 Hz, 3H);

¹³C NMR (100 MHz, CDCl₃): δ 180.9, 180.3, 174.3, 174.0, 140.6, 140.3, 139.2, 137.4, 132.8, 132.5, 129.5, 128.7, 128.4, 128.2, 128.0, 127.9, 127.2, 126.3, 125.0, 122.8, 122.5, 109.8, 109.6, 67.0, 64.5, 64.2, 60.4, 57.9, 57.3, 55.8, 51.8, 51.5, 50.9, 38.6, 38.2, 14.5, 14.3; HRMS (ESI) calcd for C₂₁H₂₃N₂O₃ m/z 351.1703 [M + H]⁺, found 351.1711.

Methyl 1'-benzyl-2'-methyl-2-oxospiro[indoline-3,3'-pyrrolidine]-5'-carboxylate (6v). Pale yellow oil (87 mg, 94% yield, dr 1 : 8.3); ¹H NMR (400 MHz, CDCl₃): δ 8.28 (s, 1H), 7.51 (d, J = 7.4 Hz, 1H), 7.44 (d, J = 7.3 Hz, 2H), 7.37–7.31 (m, 1H), 7.29 (d, J = 7.6 Hz, 1H), 7.25–7.16 (m, 2H), 7.06 (td, J = 7.6, 0.9 Hz, 1H), 6.87 (d, J = 7.7 Hz, 1H), 4.06–3.98 (m, 2H), 3.78–3.70 (m, 2H), 3.69 (s, 3H), 2.76 (dd, J = 13.7, 9.3 Hz, 1H), 2.17 (dd, J = 13.7, 3.6 Hz, 1H), 1.01 (d, J = 6.5 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 180.8, 175.0, 140.6, 139.3, 133.3, 128.5, 128.3, 128.1, 127.0, 124.2, 122.9, 109.3, 66.1, 62.3, 57.3, 51.8, 51.5, 38.3, 12.9; HRMS (ESI) calcd for C₂₁H₂₃N₂O₃ m/z 351.1703 [M + H]⁺, found 351.1711.

Methyl 1'-benzyl-2'-ethyl-2-oxospiro[indoline-3,3'-pyrrolidine]-5'-carboxylate (6w). Yellow oil (96 mg, 89% yield, dr 4.5 : 1); 1 H NMR (400 MHz, CDCl₃): δ 8.58 and 8.41 (s, 1H), 7.48–7.42 (m, 3H), 7.31 (t, J=7.4 Hz, 2H), 7.26–7.15 (m, 2H), 7.05 (td, J=7.6, 0.8 Hz, 1H), 6.86 (d, J=7.6 Hz, 1H), 4.09 (d, J=14.2 Hz, 1H), 3.99 (dd, J=9.1, 3.3 Hz, 1H), 3.88 (d, J=14.2 Hz, 1H), 3.68 (s, 3H), 2.75 (dd, J=13.6, 9.1 Hz, 1H), 2.16 (dd, J=13.6, 3.3 Hz, 1H), 1.78–1.65 (m, 1H), 1.63–1.50 (m, 2H), 0.59 (t, J=7.5 Hz, 3H); 13 C NMR (100 MHz, CDCl₃): δ 181.3, 175.1, 140.4, 139.8, 134.4, 128.4, 128.3, 127.9, 126.9, 124.0, 122.9, 109.4, 73.0, 63.1, 57.1, 52.0, 51.5, 40.0, 21.2, 11.1; HRMS (ESI) calcd for $C_{22}H_{25}N_2O_3$ m/z 365.1860 [M + H]⁺, found 365.1866.

Methyl 1'-benzyl-2'-ethyl-2-oxospiro[indoline-3,3'-pyrrolidine]-5'-carboxylate (6x). Pale yellow oil (98 mg, 91% yield, dr 1: 6.2); 1 H NMR (400 MHz, CDCl₃): δ 8.74 (d, J = 48.5 Hz, 1H), 7.80 (d, J = 7.4 Hz, 1H), 7.39 (d, J = 7.2 Hz, 2H), 7.32 (t, J = 10.1 Hz, 2H), 7.27–7.19 (m, 2H), 7.12–7.06 (m, 1H), 6.91 (d, J = 7.7 Hz, 1H), 4.01 (d, J = 14.3 Hz, 1H), 3.80 (d, J = 14.3 Hz, 1H), 3.69 (dd, J = 10.2, 6.3 Hz, 1H), 3.47 (s, 3H), 3.11 (dd, J = 10.1, 3.7 Hz, 1H), 2.66 (dd, J = 13.3, 10.5 Hz, 1H), 2.09 (dd, J = 13.3, 6.0 Hz, 1H), 1.69–1.54 (m, 2H), 0.51 (t, J = 7.5 Hz, 3H); 13 C NMR (100 MHz, CDCl₃): δ 181.6, 174.0, 140.3, 138.0, 132.3, 129.3, 128.2, 127.9, 127.1, 126.5, 122.8, 109.7, 73.0, 64.8, 56.6, 56.1, 51.8, 40.1, 23.3, 9.8; HRMS (ESI) calcd for C₂₂H₂₅N₂O₃ m/z 365.1860 [M + H] $^+$, found 365.1866.

1-Methyl-4,9-dihydro-3*H***-pyrido**[3,4-*b*]**indole** (8a). ^{13e} Yellow solid (22 mg, 35% yield), mp: 181–184 °C; ¹H NMR (400 MHz, CDCl₃): δ 9.38 (s, 1H), 7.51 (d, J = 8.0 Hz, 1H), 7.37 (d, J = 8.3 Hz, 1H), 7.20 (t, J = 7.5 Hz, 1H), 7.07 (t, J = 7.5 Hz, 1H), 3.77 (t, J = 8.3 Hz, 2H), 2.82 (t, J = 8.4 Hz, 2H), 2.38 (s, 3H); ¹³C NMR (100 MHz, CDCl₃): δ 159.2, 137.5, 128.6, 125.1, 125.0, 120.4, 120.1, 117.4, 112.2, 47.0, 21.4, 19.2; HRMS (ESI) calcd for $C_{12}H_{13}N_2$ m/z 185.1073 [M + H]⁺, found 185.1077.

1-Phenyl-4,9-dihydro-3*H***-pyrido**[3,4-*b*]indole (8b). ^{13e} Pale yellow solid (28 mg, 30%), mp: 219–223 °C;

¹H NMR (400 MHz, CDCl₃): δ 8.35 (br s, 1H), 7.85–7.60 (m, 3H), 7.57–7.45 (m, 3H), 7.43–7.15 (m, 3H), 4.06 (s, 2H), 3.01 (s, 2H);

¹³C NMR (100 MHz, CDCl₃): δ 159.4, 137.7, 136.6, 130.0, 128.9, 127.9, 127.9, 125.6, 124.6, 120.5, 120.1, 117.9, 112.1, 48.9, 19.3; HRMS (ESI) calcd for $C_{17}H_{15}N_2$ m/z 247.1230 [M + H]⁺, found 247.1226.

There are no conflicts to declare.

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