

Total synthesis of naturally occurring 1-oxygenated carbazole alkaloids - clausine E, clausenapin, indizoline and formal synthesis of clausenaline D

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Abstract

Total synthesis of indizoline and clausenapin have been accomplished *via* a facile functional group transformation of the ester functionality at the C-3 carbon of the key ester intermediate derived from clausine E by employing *o*-Claisen rearrangement and Wittig homologation as the key steps. An improved synthesis of clausine E was achieved using Eaton's reagent for the annulation step. *O*-allylation of clausine E followed by *o*-Claisen rearrangement and acetylation enabled the formal synthesis of clausenaline D.

Keywords: *o*-Claisen rearrangement, Wittig reaction, clausenapin, indizoline, carbazole alkaloids

Introduction

1-Oxygenated carbazole alkaloids exhibit promising biological activities, leading to a remarkable exploitation of myriad synthetic strategies towards their synthesis.^{1,2} Moreover, there are substantial reports in the literature on the isolation of these alkaloids from various plant sources.³⁻¹³ For example, clausine E (**1**) (Fig. 1),^{3,4} isolated from the stem bark of *Clausena excavata* showed inhibition of rabbit platelet aggregation and vasocontraction.⁴ Indizoline (**2**) was isolated from various parts of *Clausena lansium*⁵⁻¹⁰ and also from the roots of *Clausena indica*.¹¹ Various constituents of *Clausena lansium* along with indizoline showed a significant concentration-dependent inhibition of nitrite production in the case of RAW264.7, a mouse macrophage cell line used to model macrophage-mediated inflammatory events *in vitro*. Further studies showed that indizoline was also effective in causing significant-concentration dependent inhibition of tumor necrosis factor- α (TNF- α) which mediates the production of many cytokines during inflammation.⁶ It also exhibited antitumor activity against human cancer cell lines.⁸ Clausenapin (**3**) was isolated from the leaves of *Clausena heptaphylla*.¹² Clausenaline D¹³ (**5**)

was isolated from *Clausena lansium*. Significant efforts have been devoted to the synthesis of carbazole alkaloids, over the past two decades.² We have earlier reported the synthesis of clausine E using Wittig reaction and annulation as the key steps.¹⁴

Argade and coworkers reported the first total synthesis of clausenaline D starting from Boc-protected 3-formylindole and dimethyl maleate featuring Wittig reaction, selective mono-alkylation, oxidative cleavage of terminal alkene and dehydrative intramolecular cyclization as the crucial steps.¹⁶ Prior to its isolation clausenapin was synthesized as Huang Minlon reduction product of indizoline.¹¹ Unsuccessful attempts were made in the past to synthesize indizoline as well as clausenapin.¹⁷ We realized that a library of 1-oxygenated carbazole alkaloids could be synthesized *via* a common intermediate **10** which could be obtained from clausine E. Continuing our interest in the application of Wittig reaction in the synthesis of naturally occurring carbazole alkaloids,¹⁴ we now wish to report the synthesis of clausenapin (**3**) and indizoline (**2**). An improved synthesis of clausine E (**1**) and the formal synthesis of clausenaline D were also accomplished.

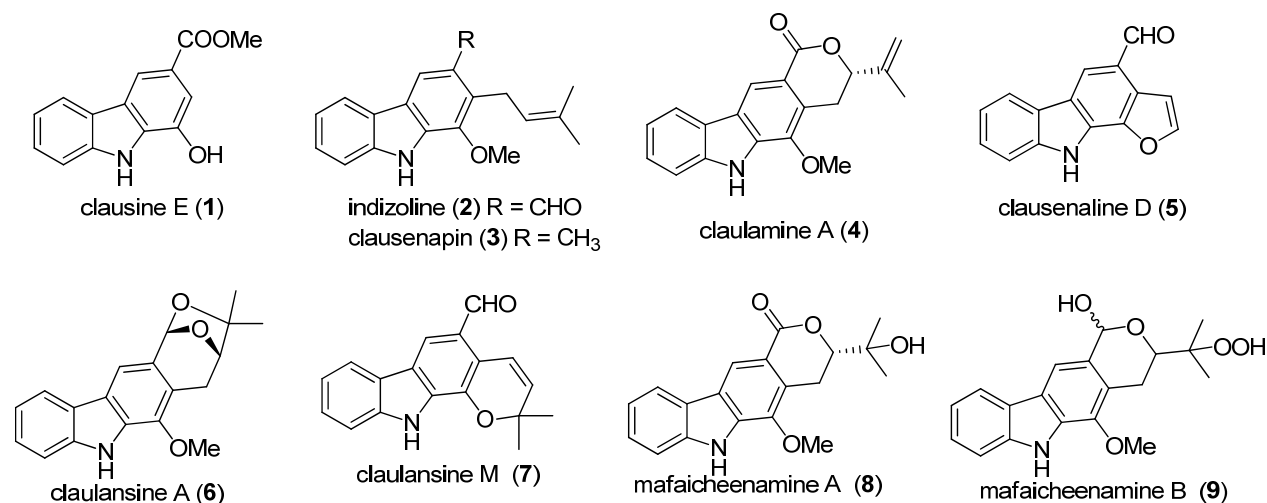


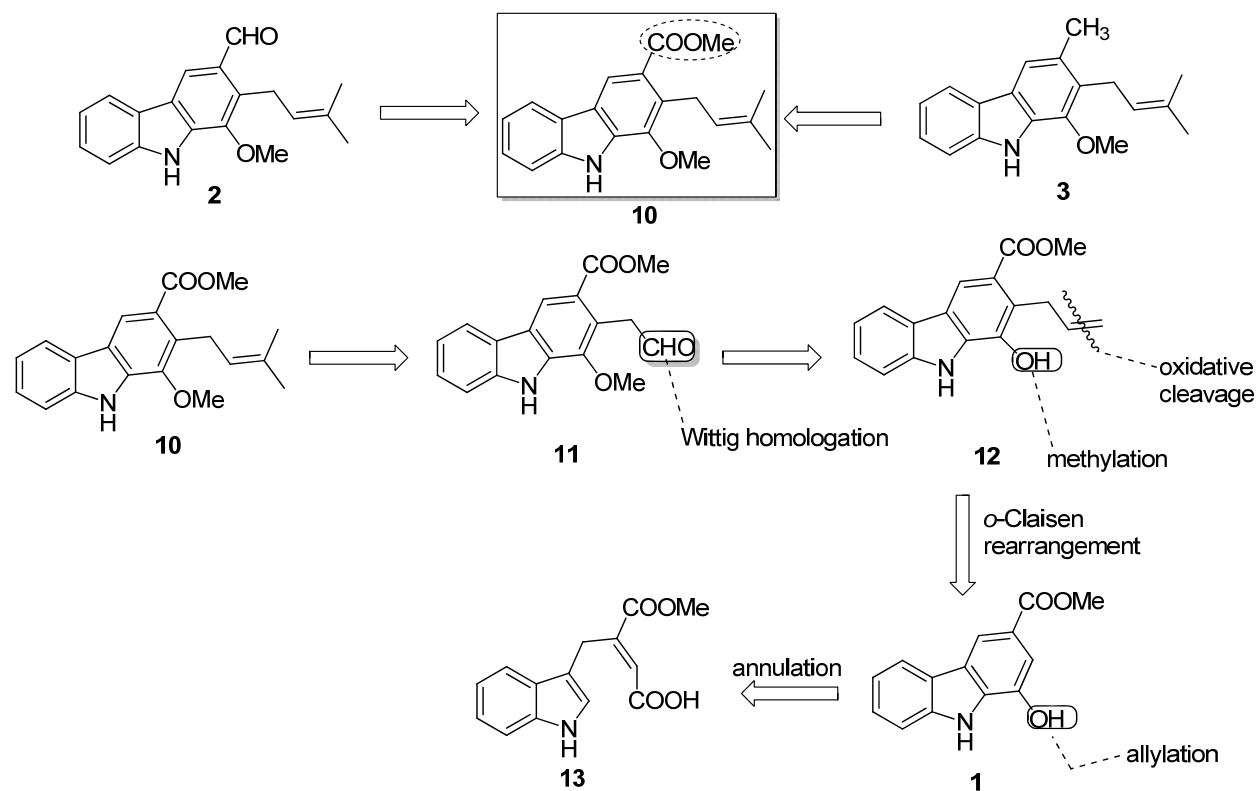
Figure 1. Structures of some naturally occurring 1-oxygenated carbazole alkaloids

We presented our preliminary work on conversion of clausine E to clausenapin *via* the intermediate **10**, at the NFCFA2015 conference.¹⁵ Subsequently, in 2016 Humne *et al.* reported synthesis of clausine E *via* Fischer-Borsche method and aromatization of 1-oxotetrahydrocarbazole using molecular iodine.¹⁸ Ullah and coworkers reported the synthesis of intermediate **10** using palladium catalysed *O*-prenylation of clausine E and subsequent microwave assisted *o*-Claisen rearrangement.¹⁹ Clausine E was synthesized using palladium acetate catalyzed cyclodehydrogenation of methyl 3-(benzyloxy)-4-(phenylamino) benzoate as the key step.¹⁹ Intermediate **10** was utilized for accomplishing the first total synthesis of (\pm)-mafaicheenamaine A. Chang and coworkers described synthesis of the intermediate 10 using *O*-allylation of

clausine E and olefin metathesis as the key steps and used it as a precursor for getting indizoline. Further they also synthesized Clausenapin, claulansine M and clausenaline D from clausine E.²⁰ Clausine E was synthesized by TFAA-mediated annulation of (*E*)-4-(1'*H*-indol-3'-yl)-3-(methoxycarbonyl)but-3-enoic acid, obtained from Stobbe condensation of indole-3-carbaldehyde,²¹ with dimethyl succinate using sodium hydride as a base.²⁰ Argade's group accomplished the synthesis of intermediate **10** by prenylation of the Wittig adduct dimethyl (*E*)-2-((1-*tert*-butoxycarbonyl)-1*H*-indol-3-yl)methylene)succinate followed by selective hydrolysis, triphosgene-induced intramolecular acylation and methylation. The intermediate **10** was then transformed to indizoline, mafaicheenamaine A, claulamine A, claulansine A and claulamine E.²²

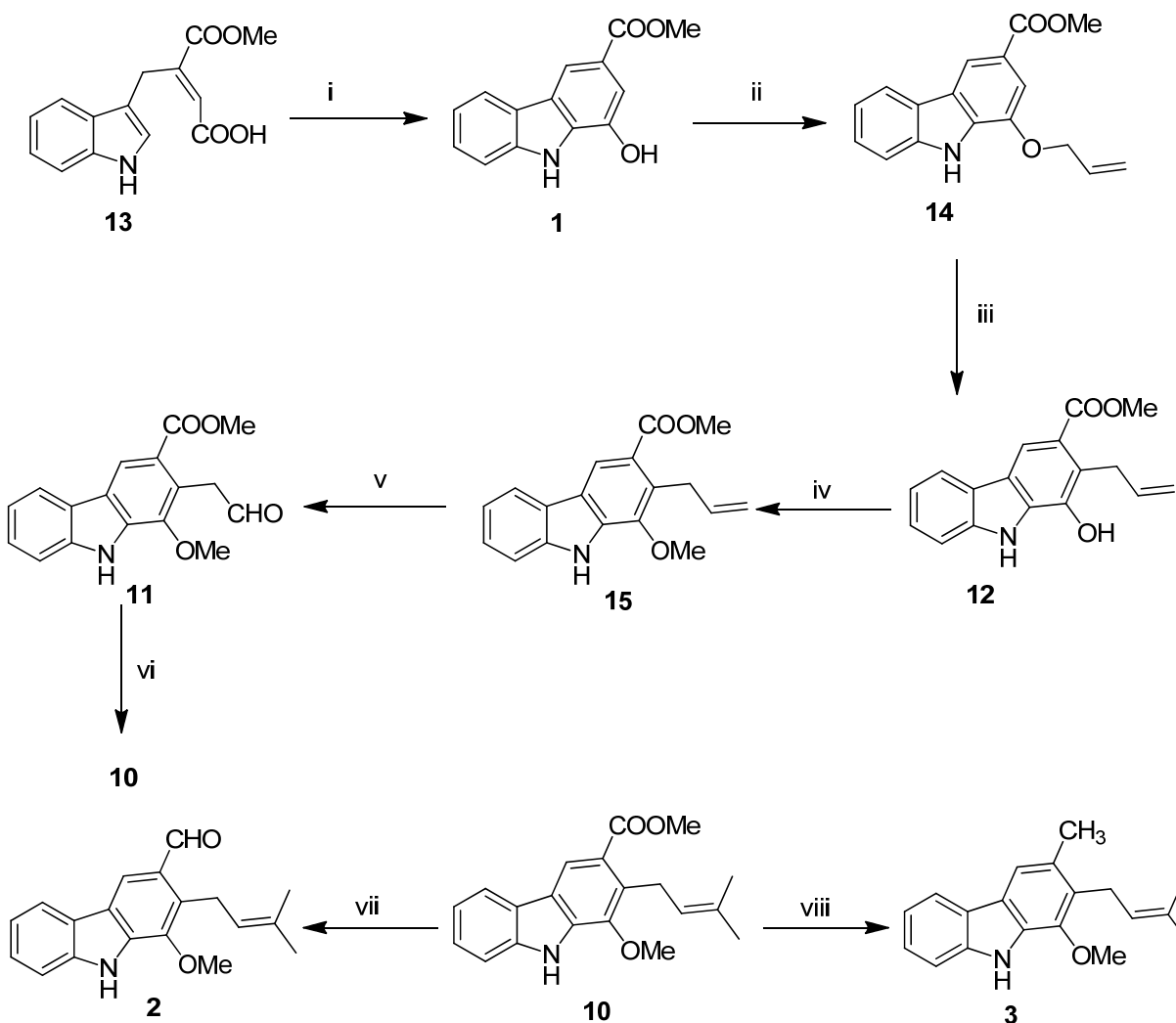
Results and Discussion

We envisioned a rapid entry into indizoline (**2**) and clausenapin (**3**) from the elaboration of a common intermediate **10** involving transformation of the ester function. A brief synthetic strategy for the synthesis of intermediate **10** is shown below (Scheme 1). Intermediate **10** would be accessible *via* Wittig homologation of the aldehyde **11**. Intermediate **11** would in turn come from carbazole **12** *via* methylation of the phenolic hydroxyl group and oxidative cleavage of the terminal double bond. Carbazole **12** would come from *O*-allylation and concomitant *o*-Claisen rearrangement of clausine E. We had earlier reported the synthesis of clausine E by carrying out annulation of acid **13**, either using sodium acetate in acetic anhydride followed by treatment with potassium carbonate in methanol or with PPA.¹⁴



Scheme 1. Retrosynthetic analysis

Our synthetic endeavours started with the acid **13**, which was subjected to annulation with the Eaton's reagent to give clausine E (**1**).²³ Use of Eaton's reagent helped in decreasing the reaction time, improved the product yield and also allowed for an easy and clean workup procedure.



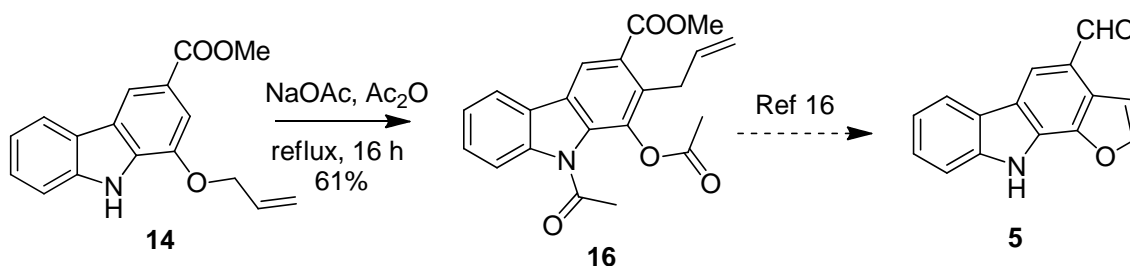
Reagents and conditions: (i) Eaton's reagent, r.t., 1.5 h, 60%; (ii) K_2CO_3 , acetone, allyl bromide, reflux, 2 h, 71%; (iii) *o*-dichlorobenzene, reflux, 45 min, 75%; (iv) K_2CO_3 , acetone, MeI, r.t., 2 h, 89%; (v) OsO_4 , $NaIO_4$, THF:H₂O (3:1), 0 °C-r.t., 10 h, 70%; (vi) triphenylphosphonium isopropyl iodide, NaHMDS, THF -10 °C-r.t., 1 h, 70%; (vii) a. LAH, THF r.t., 2 h, b. DMP, DCM, 0 °C, 0.5 h, 78%; (viii) LAH, DCM:Et₂O (1:1), r.t., 5 h, 75%.

Scheme 2. Synthesis of clausine E, indizoline and clausenapin.

Clausine E was then reacted with allyl bromide in acetone using K_2CO_3 as base to give **14** in 71% yield. The Claisen rearrangement of **14** in refluxing *o*-dichlorobenzene gave **12** in 75% yield. The initial attempts of Claisen rearrangement of intermediate **14** in *N,N*-dimethylaniline resulted in formation of the product in little lower yield (67%). The chemoselective methylation on the phenolic hydroxyl group of the Claisen rearrangement product **12** was achieved by using K_2CO_3 -MeI in acetone.²⁴ Compound **15** was transformed into the desired aliphatic aldehyde **11** by an *in situ* oxidative cleavage of the double bond, *via* the diol using OsO_4 and $NaIO_4$.¹⁶ The carbazole **11** was then treated with triphenyl(propan-2-ylidene)phosphorane,²⁵ obtained *in situ* by

the reaction of triphenylphosphonium isopropyl iodide with NaHMDS, to produce the desired intermediate **10**. The ester function in the intermediate **10** was successfully transformed into an aldehyde, on treatment with lithium aluminium hydride in THF followed by Dess-Martin periodinane mediated oxidation²⁶ resulting in the total synthesis of indizoline (**2**). The synthesis of clausenapin (**3**) was achieved from **10**, by treatment with excess LAH (Scheme 2).

We also realized that the formal synthesis of clausenaline D (**5**) could be accomplished through intermediate **16**.¹⁶ With carbazole **14** in hand, we easily synthesized intermediate **16** by refluxing **14** with sodium acetate in acetic anhydride, involving a one pot *o*-Claisen rearrangement and acetylation sequence (Scheme 3).



Scheme 3. Formal synthesis of clausenaline D.

Conclusions

In conclusion, we have accomplished the synthesis of prenylated carbazole alkaloids; indizoline, clausenapin, a formal synthesis of clausenaline D, and an improved synthesis of clausine E. Synthesis of intermediate **10** completes the formal synthesis of claulamine A, claulansine A and mafaicheenamaine A. Protection of the indole nitrogen was not vital, which helped in avoiding additional protection-deprotection steps throughout the synthesis.

Experimental Section

General. Melting points were recorded in open capillary tubes with Thiele's apparatus and are uncorrected. The IR spectra were recorded with a Shimadzu FTIR spectrophotometer. The ¹H (400 MHz) and ¹³C (100 MHz) NMR spectra were recorded on a Bruker Avance 400 instrument using CDCl₃, DMSO-d₆ or acetone-d₆ as solvent. Chemical shifts δ are expressed relative to TMS, the coupling constants *J* are given in Hz. The multiplicities of the carbon signals were obtained from DEPT-135 experiment. HRMS were recorded with a Micro-Mass ES-QTOF. Reactions were monitored by thin layer chromatography with TLC silica gel 60 F254 purchased from Merck. Column chromatography was performed on silica gel (60-120 mesh). Flash chromatography was performed on silica gel (230-400). Commercial reagents were purchased

from Sigma Aldrich or Spectrochem and used without further purification. The solvents were distilled prior to use.

Methyl 1-hydroxy-9H-carbazole-3-carboxylate (1). Freshly prepared Eaton's reagent (1 : 10 phosphorus pentoxide-methanesulfonic acid solution, 2.5 mL)²³ was added to (*E*)-3-((1H-indol-3-yl)methyl)-4-methoxy-4-oxobut-2-enoic acid (**13**) (0.8 g, 0.00308 mol) and stirred at ambient temperature for 90 minutes. To the resulting reaction mixture water (20 mL) was added and extracted with ethyl acetate (3 × 30 mL). The combined organic extracts were dried over anhydrous Na₂SO₄ and concentrated under reduced pressure. The crude residue was subjected to column chromatographic purification on silica gel using hexane :ethyl acetate (75 : 25) as an eluent to give **1** as a white solid. Yield: 0.449 g, 60%. The spectral data of **1** matched with the previously reported values.¹⁴

Methyl 1-(allyloxy)-9H-carbazole-3-carboxylate (14). To a suspension of **1** (0.6 g, 0.00248 mol) in K₂CO₃ (0.68 g, 0.00497 mol) and acetone (15 mL) was added allyl bromide (0.21 mL, 0.00248 mol) at room temperature under argon atmosphere and the reaction mixture was subjected to reflux for 2 hours. The mixture was filtered, the filtrate was concentrated under vacuum and the remaining residue was purified by flash chromatography on silica gel using hexane : ethyl acetate (90 : 10) as an eluent to give **14** as a white solid. Yield: 0.499 g, 71%, m.p. 178-180 °C; (Lit.²⁰ 179-180 °C); IR (KBr): 3369, 1687, 1311, 1217 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ: 3.99 (s, 3H), 4.79-4.81 (dt, 2H, *J* 1.2 Hz, *J* 5.6 Hz), 5.37-5.40 (dd, 1H, *J* 1.2 Hz, *J* 10.4 Hz), 5.50-5.55 (dd, 1H, *J* 1.6 Hz, *J* 17.2 Hz), 6.13-6.23 (m, 1H), 7.28-7.32 (m, 1H), 7.45-7.52 (m, 2H), 7.61 (d, 1H, *J* 1.2 Hz), 8.12 (d, *J* 8 Hz, 1H), 8.50 (s, 1H), 8.59 (br s, 1H); ¹³C NMR (100 MHz, CDCl₃) δ: 52.07 (CH₃), 69.39 (CH₂), 107.84 (CH), 111.25 (CH), 116.36 (CH), 118.42 (CH₂), 120.29 (CH), 120.76 (CH), 121.82 (Cq), 123.74 (Cq), 123.76 (Cq), 126.40 (CH), 132.84 (CH), 133.06 (Cq), 139.52 (Cq), 143.98 (Cq), 167.95 (Cq); HRMS: *m/z* [M+Na]⁺ calculated for C₁₇H₁₅NO₃Na : 304.0950, found: 304.0951.

Methyl 2-allyl-1-hydroxy-9H-carbazole-3-carboxylate (12). A solution of **14** (0.4 g, 0.00142 mol) in ortho dichlorobenzene (5 mL) under an argon atmosphere was heated at reflux for 45 minutes. The reaction mixture was purified by flash chromatography on silica gel using hexane : ethyl acetate (75 : 25) as an eluent to give **12** as a white solid. Yield: 0.3 g, 75%; m.p. 147-149 °C; IR (KBr): 3346, 3057, 1658, 1620, 1307, 1280 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ: 3.94 (s, 3H), 4.02 (d, 2H, *J* 5.6 Hz), 5.17 (d, 1H, *J* 5.2 Hz), 5.21 (s, 1H), 5.59 (s, 1H), 6.11-6.21 (m, 1H), 7.25 (m, 1H), 7.41-7.46 (m, 2H), 8.05 (d, 1H, *J* 7.6 Hz), 8.37 (s, 1H), 8.44 (br s, 1H); ¹³C NMR (100 MHz, CDCl₃) δ: 31.74 (CH₂), 52.03 (CH₃), 111.18 (CH), 116.01 (CH₂), 117.26 (CH), 120.18 (CH), 120.60 (CH), 121.81 (Cq), 122.52 (Cq), 123.69 (Cq), 126.36 (CH), 132.28 (2 × Cq), 136.93 (CH), 139.79 (Cq), 140.51 (Cq), 168.82 (Cq); HRMS: *m/z* [M+Na]⁺ calculated for C₁₇H₁₅NO₃Na : 304.0950, found: 304.0950.

Methyl 2-allyl-1-methoxy-9H-carbazole-3-carboxylate (15). To a solution of **12** (0.3 g, 0.00106 mol) in acetone (10 mL) was added K₂CO₃ (0.339 g, 0.00245 mol) and after being stirred for 5 minutes at ambient temperature, iodomethane (0.47 mL, 0.00746 mol) was added and the reaction mixture was further stirred for two hours. Acetone was evaporated under

reduced pressure. The mixture was dissolved in ethyl acetate (30 mL), washed with water (10 mL) and dried over anhydrous Na₂SO₄ and concentrated under reduced pressure. The crude residue was subjected to column chromatographic purification on silica gel using hexane : ethyl acetate (90 : 10) as an eluent to give **15** as a white solid. Yield: 0.28 g, 89%, m.p. 115-118 °C (Lit.²⁰ 112-115 °C); IR (KBr): 3346, 1681, 1606, 1346, 1263 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ: 3.95 (s, 3H), 3.98 (s, 3H), 4.03-4.06 (dt, 2H, *J* 1.6 Hz, *J* 5.6 Hz), 4.95-5.04 (m, 2H), 6.10-6.17 (m, 1H), 7.28-7.31 (m, 1H), 7.45-7.52 (m, 2H), 8.09 (d, 1H, *J* 7.6 Hz), 8.39 (br s, 1H), 8.54 (s, 1H); ¹³C NMR (100 MHz, CDCl₃) δ: 30.68 (CH₂), 51.92 (CH₃), 61.43 (CH₃), 111.13 (CH), 114.60 (CH₂), 120.29 (CH), 120.39 (CH), 120.58 (CH), 122.37 (Cq), 122.92 (Cq), 123.92 (Cq), 126.46 (CH), 131.19 (Cq), 135.51 (Cq), 138.26 (CH), 139.83 (Cq), 143.43 (Cq), 168.38 (Cq); HRMS: *m/z* [M+Na]⁺ calculated for C₁₈H₁₇NO₃Na : 318.1106, found: 318.1106.

Methyl 1-methoxy-2-(2-oxoethyl)-9H-carbazole-3-carboxylate (11). To a suspension of **15** (0.27 g, 0.000914 mol) in a mixture of THF (7.5 mL) and water (2.5 mL) was added 1% aqueous OsO₄ solution (1.6 mL, 0.00063 mol) at 0-5 °C and the reaction mixture was stirred. After 10 minutes, NaIO₄ (0.879 g, 0.00411 mol) was added to the reaction mixture and stirring was continued for 10 hours at ambient temperature. THF was evaporated under vacuum and water (15 mL) was added to the residue. The reaction mixture was extracted with ethyl acetate (3 × 15 mL) and the combined extract was washed with water, brine and dried over sodium sulfate. The organic layer was concentrated *in vacuo* and the crude residue was purified by flash chromatography on silica gel using hexane : ethyl acetate (70 : 30) as an eluent to give **11** as a white solid. Yield: 0.19 g, 70%, m.p. 170-172 °C (Lit.²⁰ 171-174 °C); IR (KBr): 3338, 1712, 1703, 1282 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ: 3.80 (s, 3H), 3.84 (s, 3H), 4.25 (s, 2H), 7.18-7.21 (m, 1H), 7.30-7.37 (m, 2H), 7.90 (d, 1H, *J* 7.6 Hz), 8.33 (d, 1H, *J* 6 Hz), 8.59 (br s, 1H), 9.87 (s, 1H); ¹³C NMR (100 MHz, CDCl₃) δ: 41.79 (CH₂), 51.96 (CH₃), 61.34 (CH₃), 111.31 (CH), 120.41 (CH), 120.51 (CH), 120.57 (CH), 120.72 (Cq), 123.62 (Cq), 123.70 (Cq), 124.76 (Cq), 126.62 (CH), 135.40 (Cq), 140.03 (Cq), 144.00 (Cq), 167.79 (Cq), 200.44 (CHO); HRMS: *m/z* [M+Na]⁺ calculated for C₁₇H₁₅NO₄Na : 320.0899, found: 320.0899.

Methyl 1-methoxy-2-(3-methylbut-2-en-1-yl)-9H-carbazole-3-carboxylate (10). A flame-dried, 2-necked round bottomed flask under argon atmosphere was charged with triphenylphosphonium isopropyl iodide (0.58 g, 0.00134 mol) and anhydrous THF (10 mL). The solution was cooled to -10 °C and 2M NaHMDS in THF (0.67 mL, 0.00134 mol) was added drop wise (till the dark red colour persisted). After ten minutes, a solution of compound **11** (0.2 g, 0.00067 mol) in THF (5 mL) was added drop wise at -10 °C to the reaction mixture. The reaction mixture was then allowed to gradually warm to room temperature and stirred for about one hour. The reaction mixture was quenched with aqueous saturated NH₄Cl solution (10 mL). THF was removed under vacuum and the crude mixture was extracted with diethyl ether (3 × 15 mL). The combined organic extract was washed with water, brine and dried over sodium sulfate. The organic layer was concentrated *in vacuo* and the crude residue was purified by flash chromatography on silica gel using hexane : ethyl acetate (80 : 20) as an eluent to give **10** as a white solid. Yield: 0.152 g, 70%, m.p. 140-142 °C (Lit.²⁰ 139-142 °C); IR (KBr): 3342, 1687,

1608, 1261 cm^{-1} ; ^1H NMR (400 MHz, acetone- d_6) δ : 1.65 (s, 3H), 1.81 (s, 3H), 3.88 (s, 3H), 3.93 (d, 2H, J 6.4 Hz), 3.95 (s, 3H), 5.22-5.25 (t, 1H, J 6.4 Hz), 7.21-7.25 (t, 1H, J 7.6 Hz), 7.41-7.45 (t, 1H, J 7.6 Hz), 7.56 (d, 1H, J 8.4 Hz), 8.15 (d, 1H, J 8 Hz), 8.48 (s, 1H), 10.73 (br s, 1H); ^{13}C NMR (100 MHz, acetone- d_6) δ : 18.19 (CH₃), 25.91 (CH₃), 26.33 (CH₂), 52.02 (CH₃), 61.25 (CH₃), 112.34 (CH), 120.37 (CH), 120.64 (CH), 121.10 (CH), 122.95 (Cq), 123.36 (Cq), 124.45 (Cq), 125.49 (CH), 127.05 (CH), 131.14 (Cq), 133.42 (Cq), 136.49 (Cq), 141.58 (Cq), 144.43 (Cq), 168.96 (Cq); HRMS: m/z $[\text{M}+\text{Na}]^+$ calculated for C₂₀H₂₁NO₃Na : 346.1419, found: 346.1419.

Methoxy-3-methyl-2-(3-methylbut-2-en-1-yl)-9H-carbazole (3). To a suspension of LAH (0.126 g, 0.0033 mol) in anhydrous CH₂Cl₂/Et₂O (6 mL, 1:1) was added drop wise compound **10** (0.09 g, 0.000278 mol) dissolved in anhydrous CH₂Cl₂/Et₂O (4 mL, 1:1) under an argon atmosphere at ambient temperature and stirring was continued for 5 hours. The reaction mixture was quenched with water (2 mL) at 0 °C and filtered. The solvent of the filtrate was removed under vacuum, the resulting residue was dissolved in chloroform (30 mL) and washed with water (10 mL). The organic layer was dried over anhydrous sodium sulphate, concentrated under vacuum and the residue was subjected to flash column chromatography on silica gel using hexane : ethyl acetate (90 : 10) as an eluent to give **3** as a pale red solid. Yield: 0.058 g, 75%, m.p. 83-85 °C (Lit.²⁰ 81-83 °C); IR (KBr): 3325, 1612, 1338, 1255 cm^{-1} ; ^1H NMR (400 MHz, acetone- d_6) δ : 1.69 (s, 3H), 1.83 (s, 3H), 2.42 (s, 3H), 3.53 (d, 2H, J 6.4 Hz), 3.92 (s, 3H), 5.11-5.15 (t, 1H, J 6.4 Hz), 7.11-7.15 (t, 1H, J 7.2 Hz), 7.31-7.35 (t, 1H, J 7.2 Hz), 7.49 (d, 1H, J 8.4 Hz), 7.66 (s, 1H), 8.01 (d, 1H, J 8 Hz), 10.27 (br s, 1H); ^{13}C NMR (100 MHz, acetone- d_6) δ : 18.07 (CH₃), 20.15 (CH₃), 25.83 (CH₃), 26.46 (CH₂), 61.09 (CH₃), 111.88 (CH), 117.50 (CH), 119.52 (CH), 120.64 (CH), 123.94 (Cq), 124.37 (Cq), 124.40 (CH), 125.99 (CH), 129.05 (Cq), 130.58 (Cq), 131.58 (Cq), 132.70 (Cq), 141.17 (Cq), 144.21 (Cq). HRMS: m/z $[\text{M}+\text{H}]^+$ calculated for C₁₉H₂₂NO: 280.1701, found: 280.1701.

1-Methoxy-2-(3-methylbut-2-en-1-yl)-9H-carbazole-3-carbaldehyde (2). To a suspension of LAH (0.025 g, 0.000649 mol) in anhydrous THF (4 mL) was added drop wise compound **10** (0.07 g, 0.000216 mol) dissolved in THF (4 mL) under an argon atmosphere. The reaction was stirred for 2 hours at ambient temperature. The reaction mixture was quenched with water (3 mL) at 0 °C and filtered. The solvent of the filtrate was removed under vacuum. The crude residue was dissolved in dry CH₂Cl₂ (5 mL) and DMP (0.0918 g, 0.000216 mol) was added to it under argon atmosphere at 0 °C. After stirring for 0.5 hours, CH₂Cl₂ was evaporated under vacuum and the reaction mixture was diluted with ether (20 mL) and washed with 10% Na₂CO₃ solution (10 mL) and water (10 mL). The organic layer was dried over anhydrous sodium sulfate and evaporated under reduced pressure. The crude residue was purified by flash chromatography on silica gel using hexane : ethyl acetate (80 : 20) as an eluent to give **2** as a white solid. Yield: 0.049 g, 78%, m.p. 168-170 °C (Lit.²⁰ 168-171 °C); IR (KBr): 3315, 1724, 1666, 1242 cm^{-1} ; ^1H NMR (400 MHz, acetone- d_6) δ : 1.67 (s, 3H), 1.85 (s, 3H), 3.98 (s, 3H), 4.00 (d, 2H, J 6.8 Hz), 5.22-5.26 (m, 1H), 7.25-7.29 (m, 1H), 7.44-7.48 (m, 1H), 7.59 (d, 1H, J 8 Hz), 8.21 (d, 1H, J 7.6 Hz), 8.47 (s, 1H), 10.26 (s, 1H), 10.93 (br s, 1H); ^{13}C NMR (100 MHz, acetone- d_6): δ 18.18

(CH₃), 24.62 (CH₂), 25.82 (CH₃), 61.51 (CH₃), 112.53 (CH), 121.05 (CH), 121.30 (CH), 122.61 (CH), 123.98 (Cq), 124.62 (Cq), 125.24 (CH), 127.33 (CH), 128.54 (Cq), 131.73 (Cq), 133.90 (Cq), 137.77 (Cq), 141.62 (Cq), 144.19 (Cq), 192.06 (CHO).

Methyl 1-acetoxy-9-acetyl-2-allyl-9H-carbazole-3-carboxylate (16). To a solution of **14** (0.11 g, 0.00039 mol) in acetic anhydride (5 mL) under argon atmosphere was added sodium acetate (0.096 g, 0.00117 mol) and the reaction mixture was refluxed for 16 hours. The reaction mixture was cooled and the solvent was removed under reduced pressure. This crude residue was dissolved in EtOAc (25 mL) washed with brine, (2 × 5 mL) and water (2 × 5 mL). The organic layer was dried over anhydrous sodium sulphate, concentrated and the residue was subjected to flash column chromatography on silica gel using hexane : ethyl acetate (90 : 10) as an eluent to give **16** as a white solid. Yield: 0.087 g, 61%, m.p. 119-121 °C (Lit.¹⁶ 120-122 °C); IR (KBr): 1755, 1732, 1697, 1607, 1427, 1224 cm⁻¹; ¹H NMR (400 MHz, acetone-d₆) δ: 2.40 (s, 3H), 2.78 (s, 3H), 3.89 (d, 2H, *J* 5.6 Hz), 3.93 (s, 3H), 4.94-5.05 (m, 2H), 5.89-5.99 (m, 1H), 7.43-7.46 (t, 1H, *J* 7.2 Hz), 7.54-7.58 (td, 1H, *J* 1.2 Hz, *J* 7.9 Hz), 8.02 (d, 1H, *J* 8.4 Hz), 8.24 (d, 1H, *J* 7.6 Hz), 8.57 (s, 1H); ¹³C NMR (100 MHz, acetone-d₆) δ: 20.63 (CH₃), 26.98 (CH₃), 32.35 (CH₂), 52.57 (CH₃), 115.14 (CH), 115.66 (CH₂), 120.87 (CH), 121.30 (CH), 124.48 (CH), 125.61 (Cq), 127.39 (Cq), 127.88 (Cq), 128.96 (CH), 133.73 (Cq), 134.18 (Cq), 137.44 (CH), 137.96 (Cq), 140.54 (Cq), 168.05 (Cq), 168.26 (Cq), 171.38 (Cq); HRMS: *m/z* [M+Na]⁺ calculated for C₂₁H₁₉NO₅Na : 388.1161, found: 388.1161.

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