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Integrated synthesis of 3,4-carbazoquinone alkaloids *N*-Me-carbazoquinocin A, B and D–F†

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Carbazole alkaloids carbazoquinocin A–F possessing a 1-alkyl-2-methyl-3,4-*ortho*-carbazoquinone framework were isolated from the microorganism *Streptomyces violaceus* 2448-SVT2 in 1995. Furthermore, they were found to exhibit strong inhibitory activity against lipid peroxidation. Herein, we report the integrated synthesis of *N*-Me-analogues of 5 members of the carbazoquinocin family of natural products, namely, *N*-Me-carbazoquinocin A, B and D–F. We employed an acid-catalyzed, intramolecular benzannulation of indole-appended Z-enoate propargylic alcohols, which was developed earlier in our laboratory, for the construction of the required carbazole framework. All five natural products were obtained in an overall yield of 50–60%, starting from a commercially available indole.

Introduction

Carbazoles (dibenzopyrroles) are nitrogen-based tricyclic frameworks, which are essential in medicinal chemistry¹ and materials chemistry.² They are also part of many structurally complex bioactive natural products and drug molecules.³ Among the many structurally diverse carbazole natural products, a 1-alkyl-2-methyl-carbazole framework has attracted special attention as it has been found to be present in different families of bioactive carbazole natural products such as lipocarbazoles, carbazoquinocins, and carazostatin.⁴ In 1995, the research group of Seto isolated six carbazoquinocins A–F, from the microorganism *Streptomyces violaceus* 2448-SVT2 (Fig. 1).⁵ These natural products share a common 1-alkyl-2-methyl-3,4-*ortho*-carbazoquinone framework and have been found to exhibit strong inhibitory activity against lipid peroxidation.⁶

Several research groups have reported the total synthesis of some of these novel alkaloids.⁷ For example, the total synthesis of carbazoquinocins A and D was achieved by the research group of Ogasawara in 1996.^{7a} Alternatively, Hibino *et al.* reported the total synthesis of carbazoquinocins B–F.^{7b} In 2003, Wulff and coworkers reported the total synthesis of carbazoquinocin C.^{7c}

Recently, in 2022, we developed an approach for the total synthesis of the 1-alkyl-2-methyl-3,4-carbazoquinone natural product *N*-Me-carbazoquinocin C together with *N*-Me-carazostatin and *N*-Me-lipocarbazole A4.^{7d} This approach employs a Brønsted acid-catalyzed intramolecular benzannulation of C-3-tethered indole-propargylic alcohols (1) (Scheme 1A)

as the key step for the construction of a carbazole unit (2). In continuation, we envisioned an opportunity to extend this methodology for the total synthesis of other members of the carbazoquinocin family, *i.e.*, carbazoquinocin A, B, and D–F (3–7) (Fig. 1). According to our retrosynthetic analysis (Scheme 1B), butyraldehyde present in 2 can be converted into the respective alkyl chain (as in 8) required for carbazoquinocin A, B and D–F through the Wittig olefination reaction using suitable phosphorous ylides. Further, oxidation at the C-3 and C-4 positions of the resultant carbazoles 8 to generate the natural products carbazoquinocin A, B and D–F (3–7) can be achieved through C-3-methylation, followed by selenium promoted oxidations.

Results and discussion

Our approach began with the synthesis of the required carbazole-butylaldehyde (2) from *N*-Me-indole, following our earlier procedure.^{7d} Initially, we aimed at the total synthesis of *N*-Me-

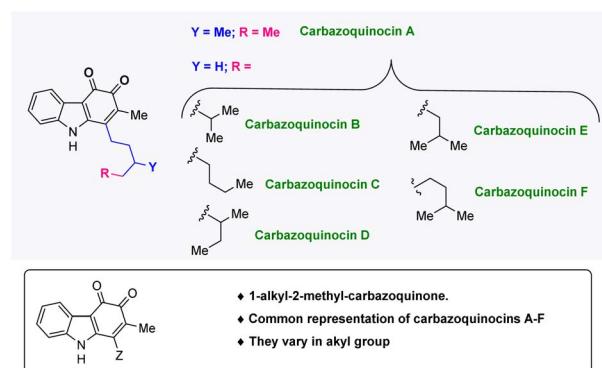
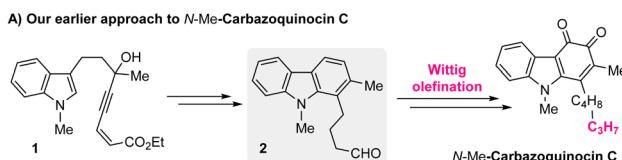


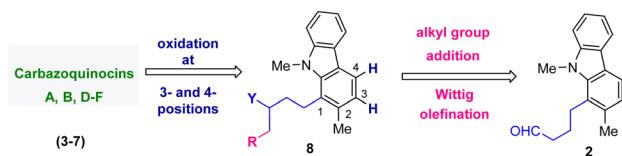
Fig. 1 Structures of carbazoquinocin A–F natural products.

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B) This work: Retrosynthetic Analysis for *N*-Me-Carbazooquinocin A, B and D-F

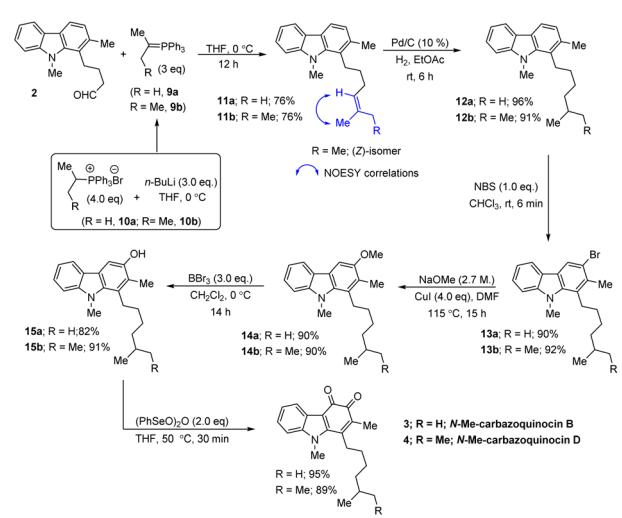


Scheme 1 (A) Our earlier work: total synthesis of *N*-Me-carbazooquinocin C. (B) Retrosynthetic analysis for the total synthesis of carbazooquinocins A, B, and D-F (3-7) from aldehyde 2.

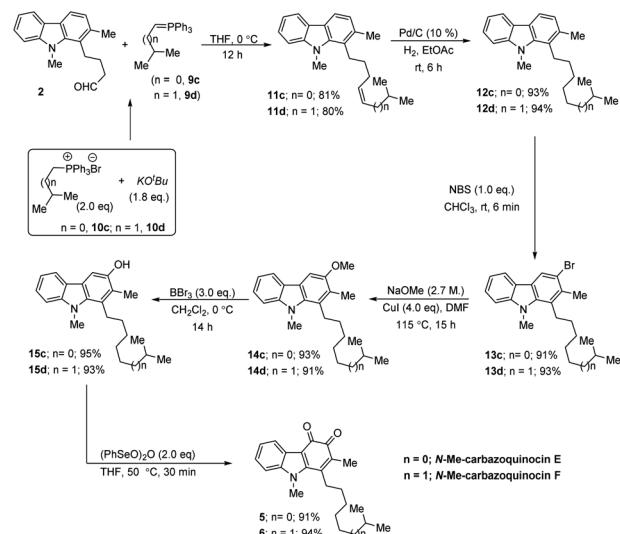
carbazooquinocin B (3) (Scheme 2, R = H). Accordingly, Wittig olefination of aldehyde 2 with phosphorous ylide (9a) at 0 °C (generated *in situ* from (iso-propyl)PPh₃Br⁸ (10a) and ⁷BuLi at 0 °C) afforded olefin (11a) in 76% yield. The hydrogenation reaction of 11a with Pd/C (10%) under an H₂ atmosphere in ethyl acetate at room temperature (rt) for 6 h gave 1-isoheptylcarbazole (12a) (96% yield).

Next, the regioselective electrophilic bromination of carbazole 12a with NBS in CHCl₃ at rt gave the corresponding 3-bromocarbazole (13a) (90%) within 6 min reaction time. Next, we employed a two-step strategy for the efficient conversion of bromine in 13a into a hydroxyl group. Initially, heating a DMF suspension of 13a, NaOMe and CuI at 115 °C gave 3-methoxy carbazole (14a) in 90% yield. Subsequent treatment of 14a with BBr₃ in CH₂Cl₂ at 0 °C to rt for 14 h resulted in the formation of the phenol product (15a) in 82% yield *via* deprotection of the methoxy group. Finally, the synthesis of *N*-Me-carbazooquinocin B 3 was achieved by employing the (PhSeO)₂O-promoted oxidation of 15a. Accordingly, heating (50 °C) a solution of 15a and (PhSeO)₂O in THF gave *N*-Me-carbazooquinocin B 3 in 95% yield. After successfully completing the total synthesis of *N*-

Me-carbazooquinocin B 3, we extended this strategy for the synthesis of *N*-Me-carbazooquinocin D 4 by employing the carbazole-butylaldehyde 2 as the starting point (Scheme 2, R = Me). According to our retrosynthetic analysis (Scheme 1B), the only difference between carbazooquinocin B 3 and carbazooquinocin D 4 is the incorporation of an iso-propyl group (R = H) vs. sec-butyl group (R = Me) in aldehyde 2. Therefore, olefination of aldehyde 2 using sec-butylidene phosphorous ylide 9b (generated from (sec-butyl)PPh₃Br⁸ 10b and ⁷BuLi at 0 °C) afforded Z-olefin 11b (R = Me) in 76% yield. The geometry of the olefin in 11b was identified by using the correlations observed between =CMe and =C-H in the corresponding NOESY NMR spectrum. Further, by operating the same sequence of five functional group transformations, as depicted in Scheme 2 (R = Me), *i.e.*, hydrogenation (12b, 91%)–bromination (13b, 92%)–methoxylation (14b, 90%)–demethylation (15b, 91%)–oxidation (4, 89%), we efficiently converted olefin 11b into *N*-Me-carbazooquinocin D 4, with an overall yield of 61% for five steps. Subsequently, we envisioned the extension of this methodology for the total synthesis of *N*-Me-carbazooquinocin E 5 and F 6 (Scheme 3). The required alkyl chains present in compounds 5 and 6 could be achieved by incorporating an iso-butyl group (n = 0) and iso-pentyl group (n = 1) in butylaldehyde 2 *via* Wittig olefination reaction, respectively. Wittig olefination of aldehyde 2 using iso-butylidene phosphorous ylide 9c (ref. 8) and iso-pentylidene phosphorous ylide 9d (ref. 8) exclusively afforded Z-olefins 11c (n = 0; 81%) and 11d (n = 1; 80%), respectively. After having both olefins 11c and 11d in hand, we employed the five-step synthetic strategy (as shown for *N*-Me-carbazooquinocin B 3 and D 4 in Scheme 2), *i.e.*, hydrogenation (12c & 12d)–bromination (13c & 13d)–methoxylation (14c & 14d)–demethylation (15c & 15d)–oxidation (5 & 6), respectively. These sequential synthetic transformations efficiently converted olefins 11c & 11d into the corresponding natural products *N*-Me-carbazooquinocin E 5 and *N*-Me-carbazooquinocin F 6, respectively (Scheme 3).

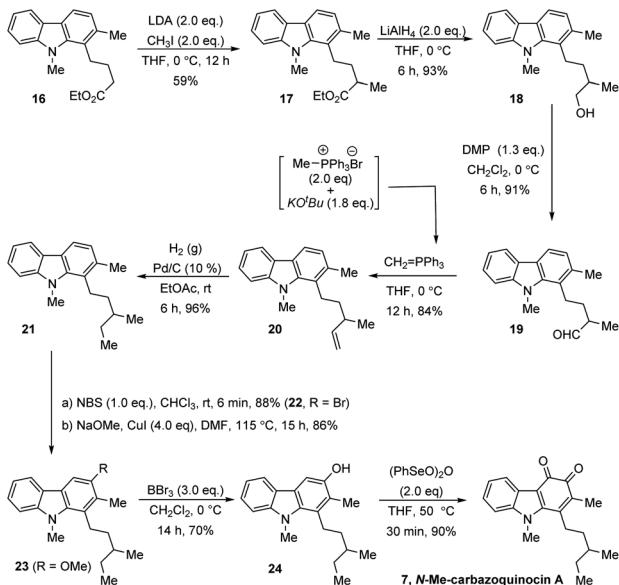


Scheme 2 Total synthesis of *N*-Me-carbazooquinocin B and D.



Scheme 3 Unified total synthesis of *N*-Me-carbazooquinocin E and F.





Scheme 4 Extension to the total synthesis of *N*-Me-carbazoquinocin A.

Finally, we also extended this strategy for the total synthesis of *N*-Me-carbazoquinocin A 7, which possesses a 3-methylpentyl alkyl chain at the C₁-position of the carbazoquinone (Scheme 4). We envisioned carbazole ester (16) as a suitable starting material for this purpose. α -Methylation (for *sec*-methyl) followed by conversion of the carboxylate group into an ethyl group generates the required 3-methylpentyl chain at the C₁-position. Accordingly, the LDA-mediated enolate formation of ester 16 followed by quenching with MeI gave α -methyl ester (17) in 59% yield. The reduction of 17 with LiAlH₄ followed by oxidation of the resultant primary alcohol (18) with Dess–Martin periodinane (DMP) gave aldehyde (19) in 84% overall yield for two steps.

Further, olefination of aldehyde 19 using methylidene-phosphorous ylide, followed by hydrogenation of the resultant olefin (20) gave 1-[3-methylpentyl] carbazole (21). Subsequently, we employed a four-step synthetic strategy, which involved bromination (22)–methoxylation (23)–demethylation (24)–oxidation (7). These sequential synthetic transformations efficiently converted carbazole 21 into *N*-Me-carbazoquinocin A 7 with an overall yield of 48% for four steps.

Conclusions

In conclusion, we developed an efficient and practically viable synthetic approach for the unified synthesis of 3,4-carbazoquinone-based carbazole alkaloids *N*-Me-carbazoquinocins A, B and D–F. Our earlier methodology, an acid-catalyzed, intramolecular benzannulation of indole-appended *Z*-enoate propargylic alcohols, was employed for the construction of the required carbazole framework. All five natural products were obtained in an overall yield of 50–60%, starting from the commercially available indole.

Experimental section

General information

All solvents were distilled prior to use and anhydrous solvents were prepared according to the standard drying procedures. All non-aqueous reactions were carried out under an atmosphere of nitrogen in flame-dried glassware. Commercially available chemicals were purchased from Sigma-Aldrich, Alfa Aesar and Spectrochem Pvt. Ltd and were used as received without further purification. Infrared (IR) spectra were recorded on a JASCO 4100 FT-IR spectrometer. ¹H NMR spectra were measured on a Bruker AVANCE 400 MHz or Bruker AVANCE 500 MHz spectrometer. Chemical shifts are reported in ppm relative to solvent signals. ¹³C NMR spectra were recorded on a Bruker AVANCE 100 MHz or Bruker AVANCE 125 MHz spectrometer with complete proton decoupling. Chemical shifts are reported in ppm from the residual solvent as an internal standard [CDCl₃ δ = 7.26 ppm for ¹H, δ = 77.16 ppm for ¹³C or calibrated to tetramethylsilane (δ = 0.00)]. The following abbreviations are used to indicate multiplicities: s-singlet; d-doublet; t-triplet; q-quartet; quint-quintet; sext-sextet; sept-septet; m-multiplet; dd-doublet of doublet; dt-doublet of triplet; dq-doublet of quartet; td-triplet of doublet; tt-triplet of triplet; dq-doublet of quartet; br-broad; *J*-coupling constant in Hz. The coupling constant *J* (Hz) was rounded to one decimal place for all compounds. When a coupling pattern can be assigned as a combination of multiplicities, the above-mentioned abbreviations were combined to describe the observed patterns (*i.e.*, dt, doublet of triplets). Mass spectra were recorded by the electrospray ionization (ESI) method on a Q-TOF Micro with a lock spray source. For thin layer chromatography (TLC) analysis throughout this work, E-Merck precoated TLC plates (silica gel 60 F254 grade, 0.25 mm) were used and visualized using a UV lamp (366 or 254 nm) or by using of one of the following visualization reagents: PMA: 1 g phosphomolybdic acid/10 mL ethanol; KMnO₄: 0.15 g potassium permanganate, 1 g K₂CO₃/20 mL water. Acme (India) silica gel (100–200 mesh) was used for column chromatography.

General procedure I (GP-I): Wittig olefination with primary alkyl-based phosphonium salts and ^tBuOK base.^{9,4d} RPPh₃X (2.0 equiv.) and anhydrous THF (8 mL) were added to a flame-dried round-bottom flask under an N₂ atmosphere. After the mixture was cooled to 0 °C, ^tBuOK (1.8 equiv.) was solubilized in anhydrous THF (4 mL) and added dropwise to the reaction mixture over a period of 5 min. The resulting mixture was warmed to room temperature and stirred for 1 h. Then the reaction mixture was cooled back 0 °C, and a solution of aldehyde 2 (1.0 equiv.) in anhydrous THF (5 mL) was added dropwise. The final reaction mixture was stirred at room temperature until the TLC showed the complete consumption of the aldehyde. After completion, saturated NH₄Cl solution (6 mL) was added to quench the reaction at 0 °C. Next, diethyl ether (5 mL) and water (5 mL) were added. The color of the mixture changed from pale-yellow to white. After separation of the layers, the residual compound from the aqueous layer was extracted with EtOAc (3 \times 5 mL). The combined organic layers were dried over anhydrous



Na_2SO_4 , filtered, and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (9 : 1 hexanes/EtOAc) provided the corresponding alkene.

General procedure II (GP-II): Wittig olefination with secondary alkyl-based phosphonium salts and *n*-BuLi base.⁹ RPPh_3X (2.0 equiv.) and anhydrous THF (8 mL) were added to a flame-dried round-bottom flask under an N_2 atmosphere. After the mixture was cooled to 0 °C, *n*-BuLi (3.0 equiv., 1.6 M in hexane) was solubilized in anhydrous THF (4 mL) and added dropwise to the reaction mixture over a period of 5 min. The resulting mixture was warmed to room temperature and stirred for 1 h. Then the reaction mixture was cooled back 0 °C, and a solution of aldehyde 2 (1.0 equiv.) in anhydrous THF (5 mL) was added dropwise. The final reaction mixture was stirred at room temperature until TLC showed the complete consumption of the aldehyde. After completion, saturated NH_4Cl solution (6 mL) was added to quench the reaction at 0 °C. Next, diethyl ether (5 mL) and water (5 mL) were added. The colour of the mixture changed from pale yellow to white. After separation of the layers, the residual compound from the aqueous layer was extracted with EtOAc (3 × 5 mL). The combined organic layers were dried over anhydrous Na_2SO_4 , filtered, and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (9 : 1 hexanes/EtOAc) provided the corresponding alkene.

General procedure III (GP-III): hydrogenation of internal alkenes using Pd/C. Pd/C (10 w/w%) was added to a well-stirred solution of internal alkenes (1.0 equiv.) in EtOAc (10 mL). The resulting reaction mixture was stirred under hydrogen (1 atm) atmosphere for an appropriate time at room temperature. After completion, the reaction mixture was filtered through a Celite® pad by using EtOAc (20 mL). The filtrate was concentrated *in vacuo*. Purification of crude product using a silica gel column chromatography (hexanes/EtOAc) provided the desired reduced product.

General procedure IV (GP-IV): bromination of carbazoles using *N*-bromosuccinimide (NBS). *N*-Bromosuccinimide (1.0 equiv.) was added a solution of carbazole derivative (1.0 equiv.) in chloroform (CHCl_3 , 10 mL) at room temperature under a nitrogen atmosphere. The reaction mixture was stirred at the same temperature for 6 min until TLC showed the complete consumption of the starting material. After completion, water (10 mL) was added. The residual compound from the aqueous layer was extracted with CH_2Cl_2 (3 × 5 mL). The combined organic layers were dried over anhydrous Na_2SO_4 , filtered, and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (hexanes/EtOAc) provided the desired 3-bromocarbazole derivative.

General procedure V (GP-V): methylation using NaOMe. A 15 mL Schlenk tube equipped with a magnetic stirrer was evacuated, and then backfilled with nitrogen gas. This process was repeated three times. Next, 2 mL of anhydrous MeOH was added, and the reaction vessel was cooled to 0 °C. To this stirring MeOH solvent, sodium metal (54 equiv.) was carefully added in portions under a positive nitrogen pressure to form an ~2.7 M solution of sodium methoxide (NaOMe) in MeOH. After the complete dissolution of Na in MeOH, the solution became

thick and light yellowish. Next, the corresponding 3-bromocarbazole (1.0 equiv.) dissolved in DMF (1.4 mL) and CuI (4.0 equiv.) were added to this freshly prepared NaOMe solution under a nitrogen atmosphere. The resulting reaction mixture was transferred to a preheated oil bath and stirred at 115 °C for 15 h. After the complete consumption of the starting material, as indicated by TLC, the crude reaction mixture was filtered through a short plug of Celite® and washed with EtOAc. The filtrate was sequentially washed with saturated NH_4Cl solution (5 mL), water (10 mL) and brine (5 mL), dried over anhydrous Na_2SO_4 and concentrated *in vacuo*. Purification of the crude product using a silica gel column chromatography (9 : 1 hexanes/EtOAc) provided the desired 3-methoxycarbazole derivatives.

General procedure VI (GP-VI): BBr_3 -mediated deprotection of the methoxy group to phenol. A 1 M solution of boron tribromide (BBr_3) in CH_2Cl_2 (3.0 equiv.) was added dropwise to a stirred solution of the corresponding 3-methoxycarbazole derivative (1.0 equiv.) in anhydrous CH_2Cl_2 (4 mL) at 0 °C. The reaction mixture was allowed to warm up to room temperature and further stirred for the appropriate time. After completion of the reaction, the reaction mixture was quenched with H_2O (5 mL). The organic layer was separated, and the residual compound from the aqueous layer was extracted with EtOAc (3 × 5 mL). The combined organic layers were washed with H_2O , brine and dried (Na_2SO_4) and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (4 : 1 hexane/EtOAc) provided the desired 3-hydroxycarbazole derivative.

General procedure VII (GP-VII): synthesis of *N*-Me carbazoquinocin natural products. $(\text{PhSeO})_2\text{O}$ (2.0 equiv.) was added to a solution of *N*-methyl-2-alkyl-3-hydroxy carbazole (1.0 equiv.) in THF (4 mL) at rt under a nitrogen atmosphere. The reaction mixture was stirred at 50 °C for 30 min. After cooling to room temperature, the mixture was quenched with water, and the residual compound from the aqueous layer was extracted with EtOAc (3 × 5 mL). The combined organic layers were washed with H_2O , brine and dried (Na_2SO_4) and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (hexane/EtOAc) provided the desired *N*-methylcarbazoquinocin natural products.

General procedure VIII (GP-VIII): preparation of required Wittig salts.⁸ Triphenylphosphine (1.0 mmol) was placed in an oven-dried Schlenk tube and dissolved in 2 mL toluene at room temperature under an N_2 -atmosphere. To this reaction mixture, the required alkyl bromide (1.1 mmol) was added. A white precipitate was formed after stirring and refluxing at 110 °C for 48 h using an oil bath. The reaction mixture was cooled to room temperature, and the phosphonium salt was recrystallized from petroleum ether and ethyl acetate solvent mixture. The salt was separated, washed with Et_2O , and dried *in vacuo* to give the corresponding phosphonium bromide.

2,9-Dimethyl-1-(5-methylhex-4-en-1-yl)-9*H*-carbazole (11a). Following GP-II. *n*-BuLi (0.22 mL, 0.36 mmol, 3.0 equiv., 1.6 M in hexane) was added to a solution of isopropyltriphenylphosphonium bromide **10a** (208 mg, 0.48 mmol, 4.0 equiv.) in anhydrous THF (5 mL), followed by aldehyde 2 (32 mg, 0.12 mmol, 1.0 equiv.) and stirred for 12 h.



The corresponding alkene product **11a** was isolated as a colourless gummy compound (26 mg, 0.08 mmol, 76%) using hexane/EtOAc (9 : 1) as the eluent.

¹H NMR (400 MHz, CDCl₃) δ 8.02 (d, *J* = 7.7 Hz, 1H), 7.84 (d, *J* = 7.8 Hz, 1H), 7.44 (dd, *J*₁ = 7.6 Hz and *J*₂ = 7.2 Hz, 1H), 7.36 (d, *J* = 8.2 Hz, 1H), 7.20 (dd, *J*₁ = 7.6 Hz and *J*₂ = 7.2 Hz, 1H), 7.06 (d, *J* = 7.8 Hz, 1H), 5.26 (t, *J* = 7.0 Hz, 1H), 4.07 (s, 3H), 3.17–3.07 (m, 2H), 2.51 (s, 3H), 2.22 (q, *J* = 7.1 Hz, 2H), 1.75 (s, 3H), 1.72 (d, *J* = 8.6 Hz, 1H), 1.68 (s, 3H) and 1.61–1.57 (m, 1H) ppm.

¹³C[¹H] NMR (100 MHz, CDCl₃) δ 142.2, 140.0, 134.6, 132.5, 125.3, 124.1, 124.1, 123.2, 122.7, 122.4, 119.6, 118.9, 117.6, 108.7, 32.6, 31.7, 28.4, 28.1, 25.9, 20.2 and 17.9 ppm.

IR (ATR): 2956, 1423, 1421, 1389, 1265 and 749 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₁H₂₆N 292.2060; found 292.2063 (1.0 ppm).

TLC: *R*_f = 0.4 (19 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(5-methylhexyl)-9H-carbazole (12a). Following GP-III. Pd/C (3 mg, 10 w/w%) was added to a stirred solution of olefin **11a** (24 mg, 0.08 mmol, 1.0 equiv.) in EtOAc (4 mL). The resulting reaction mixture was stirred under a hydrogen (1 atm) atmosphere for 6 h at room temperature. This reaction mixture was filtered and purified using silica gel column chromatography (19 : 1 hexanes/EtOAc) to provide the desired hydrogenation product **12a** (23 mg, 0.07 mmol, 96%) as a gummy compound.

¹H NMR (500 MHz, CDCl₃) δ 7.89 (d, *J* = 7.7 Hz, 1H), 7.71 (d, *J* = 7.8 Hz, 1H), 7.30 (dd, *J*₁ = 7.0 Hz and *J*₂ = 8.0 Hz, 1H), 7.21 (d, *J* = 8.2 Hz, 1H), 7.08 (dd, *J*₁ = 7.0 Hz and *J*₂ = 7.5 Hz, 1H), 6.93 (d, *J* = 7.8 Hz, 1H), 3.90 (s, 3H), 3.00–2.93 (m, 2H), 2.39 (s, 3H), 1.54–1.52 (m, 3H), 1.40–1.38 (m, 2H), 1.25 (q, *J* = 7.8 Hz, 2H) and 0.81 (d, *J* = 6.6 Hz, 6H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 142.3, 140.0, 134.5, 125.3, 124.3, 123.2, 122.7, 122.5, 119.6, 118.9, 117.6, 108.7, 39.0, 32.6, 32.0, 28.5, 28.2, 27.9, 22.8 and 20.2 ppm.

IR (ATR): 3451, 2928, 2854, 1666, 1589, 1453, 1443, 1362, 1324, 1243, 1096, 826 and 738 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₁H₂₈N 294.2216; found 294.2224 (3.0 ppm).

TLC: *R*_f = 0.5 (hexane).

3-Bromo-2,9-dimethyl-1-(5-methylhexyl)-9H-carbazole (13a). Following GP-IV. N-Bromosuccinimide (26 mg, 0.14 mmol, 1.0 equiv.) was added to a solution of carbazole **12a** (42 mg, 0.08 mmol, 1.0 equiv.) in chloroform (CHCl₃) (6 mL) under a nitrogen atmosphere. The reaction mixture was stirred for 6 min. After completion, water (10 mL) was added and extracted with CH₂Cl₂ (3 × 5 mL). The combined organic layers were dried and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (19 : 1 hexanes/EtOAc) provided the desired product **13a** (48 mg, 0.12 mmol, 90%) as a white solid.

¹H NMR (400 MHz, CDCl₃) δ 8.14 (s, 1H), 7.97 (d, *J* = 7.7 Hz, 1H), 7.46 (t, *J* = 7.6 Hz, 1H), 7.35 (d, *J* = 8.2 Hz, 1H), 7.22 (t, *J* = 7.3 Hz, 1H), 4.01 (s, 3H), 3.18–3.07 (m, 2H), 2.59 (s, 3H), 1.68–1.59 (m, 3H), 1.56–1.49 (m, 2H), 1.30 (d, *J* = 8.2 Hz, 2H) and 0.94 (d, *J* = 6.6 Hz, 6H) ppm.

¹³C[¹H] NMR (100 MHz, CDCl₃) δ 142.5, 139.0, 132.9, 126.0, 123.7, 122.1, 121.6, 119.8, 119.3, 116.8, 108.9, 38.8, 32.7, 32.0, 29.6, 28.1, 27.8, 22.8 and 19.8 ppm.

IR (ATR): 2964, 2846, 2365, 1456, 1413, 1271, 1144, 1016, 828, 773 and 736 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₁H₂₇NBr 372.1321; found 372.1326 (1.3 ppm).

TLC: *R*_f = 0.55 (hexane).

M.P.: 173–175 °C.

3-Methoxy-2,9-dimethyl-1-(5-methylhexyl)-9H-carbazole (14a).

Following GP-V. A freshly prepared NaOMe solution (~2.7 M in MeOH) was added to a DMF (1.6 mL) solution of 3-bromocarbazole **13a** (38 mg, 0.1 mmol, 1.0 equiv.), followed by CuI (77 mg, 0.39 mmol, 4.0 equiv.), and the reaction mixture was stirred at 115 °C for 15 h. The reaction mixture was filtered, and the filtrate was sequentially washed with saturated NH₄Cl solution (5 mL), water (10 mL) and brine (5 mL), dried concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (9 : 1 hexanes/EtOAc) provided the desired 3-methoxycarbazole **14a** (29 mg, 0.09 mmol, 90%) as a colorless solid.

¹H NMR (500 MHz, CDCl₃) δ 7.99 (d, *J* = 7.7 Hz, 1H), 7.44–7.39 (m, 2H), 7.36 (d, *J* = 8.2 Hz, 1H), 7.17 (t, *J* = 7.3 Hz, 1H), 4.05 (s, 3H), 3.95 (s, 3H), 3.16–3.10 (m, 2H), 2.38 (s, 3H), 1.70–1.63 (m, 2H), 1.60 (dd, *J* = 13.3, 6.7 Hz, 1H), 1.52–1.31 (m, 2H), 1.29–1.26 (m, 2H) and 0.92 (d, *J* = 6.6 Hz, 6H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 152.4, 142.5, 134.9, 125.8, 125.1, 124.8, 123.3, 121.9, 119.5, 118.5, 108.8, 99.3, 56.3, 39.0, 32.8, 31.9, 28.8, 28.2, 27.9, 22.8 and 12.1 ppm.

IR (ATR): 2945, 2856, 2392, 1471, 1414, 1288, 1213, 1154, 1126, 845 and 742 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₂H₃₀NO 324.2322; found 324.2342 (6.1 ppm).

TLC: *R*_f = 0.2 (hexane).

M.P.: 162–164 °C.

2,9-Dimethyl-1-(5-methylhexyl)-9H-carbazol-3-ol (15a).

Following GP-VI. A 1 M solution of boron tribromide (BBr₃, in CH₂Cl₂; 0.33 mL, 0.33 mmol, 4.0 equiv.) was added a stirring solution of 3-methoxycarbazole **14a** (27 mg, 0.08 mmol, 1.0 equiv.) in anhydrous CH₂Cl₂ (4 mL) at 0 °C and further stirred for 11 h. The reaction was quenched with H₂O (5 mL) and extracted with CH₂Cl₂ (3 × 5 mL). The combined organic layers were dried over (Na₂SO₄) and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (4 : 1 hexanes/EtOAc) provided phenol **15a** (21 mg, 0.07 mmol, 82% yield) as a pale-yellow liquid.

¹H NMR (400 MHz, CDCl₃) δ 7.91 (d, *J* = 7.7 Hz, 1H), 7.42 (t, *J* = 7.6 Hz, 1H), 7.34 (d, *J* = 7.8 Hz, 2H), 7.15 (t, *J* = 7.3 Hz, 1H), 4.03 (s, 3H), 3.16–3.08 (m, 2H), 2.40 (s, 3H), 1.70–1.58 (m, 3H), 1.53 (dd, *J* = 15.0, 7.2 Hz, 2H), 1.32–1.26 (m, 2H) and 0.92 (d, *J* = 6.6 Hz, 6H) ppm.

¹³C[¹H] NMR (100 MHz, CDCl₃) δ 147.7, 142.6, 135.1, 125.7, 125.4, 122.8, 122.4, 122.2, 119.7, 118.4, 108.7, 103.1, 38.9, 32.8, 32.0, 28.8, 28.2, 27.9, 22.8 and 12.1 ppm.

IR (ATR): 3325, 2926, 2854, 2363, 2351, 1485, 1451, 1264, 1232, 902, 782 and 731 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₁H₂₈NO 310.2165; found 310.2174 (3.0 ppm).

TLC: *R*_f = 0.25 (9 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(5-methylhexyl)-3H-carbazole-3,4(9H)-dione [N-methylcarbazoquinolin B] (3).

Following GP-VII. A solution



of carbazol-3-ol **15a** (20 mg, 0.06 mmol) and $(\text{PhSeO})_2\text{O}$ (47.0 mg, 0.13 mmol, 2.0 equiv.) in THF (4 mL) was stirred at 50 °C for 30 min. The mixture was quenched with water and the residual compound from the aqueous layer was extracted with EtOAc (3×5 mL). The combined organic layers were dried (Na_2SO_4) and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (3 : 1 hexanes/EtOAc) provided the desired *N*-methyl carbazoquinocin B **3** (20 mg, 0.06 mmol 95% yield) as a dark-brown glittering solid.

^1H NMR (400 MHz, CDCl_3) δ 8.04 (s, 1H), 7.18 (s, 3H), 3.80 (s, 3H), 2.68–2.55 (m, 2H), 1.82 (s, 3H), 1.49 (d, $J = 5.4$ Hz, 3H), 1.42 (d, $J = 6.7$ Hz, 2H), 1.18 (s, 2H) and 0.83 (d, $J = 6.5$ Hz, 6H) ppm.

$^{13}\text{C}[^1\text{H}]$ NMR (100 MHz, CDCl_3) δ 183.2, 173.6, 144.9, 142.6, 139.6, 134.5, 125.7, 124.9, 121.8, 113.7, 110.9, 38.7, 33.1, 29.9, 28.6, 28.1, 27.8, 22.7 and 11.9 ppm.

IR (ATR): 2961, 2942, 2863, 1665, 1652, 1632, 1499, 1481, 1459, 1369, 1234 and 781 cm^{-1} .

HRMS (ESI) m/z : [M + H]⁺ calcd for $\text{C}_{21}\text{H}_{26}\text{NO}_2$ 324.1958; found 324.1935 (7.0 ppm).

TLC: $R_f = 0.34$ (4 : 1 hexanes/EtOAc).

M.P.: 152–154 °C.

2,9-Dimethyl-1-(5-methylhept-4-en-1-yl)-9*H*-carbazole (11b).

Following GP-II. *n*-BuLi (0.19 mL, 0.31 mmol, 3.0 equiv., 1.6 M in hexane) was added to a solution of *sec*-butyl-triphenylphosphonium bromide **10b** (144 mg, 0.42 mmol, 4.0 equiv.) in anhydrous THF (5 mL), followed by aldehyde **2** (28 mg, 0.10 mmol, 1.0 equiv.) and stirred for 12 h. The corresponding alkene product **11b** was isolated as a colorless liquid (23.3 mg, 0.07 mmol, 76%) using hexanes/EtOAc (9 : 1) as the eluent.

^1H NMR (400 MHz, CDCl_3) major isomer: δ 7.93 (d, $J = 7.6$ Hz, 1H), 7.74 (d, $J = 7.7$ Hz, 1H), 7.34 (dd, $J_1 = 7.6$ Hz and $J_2 = 8.0$ Hz, 1H), 7.27 (d, $J = 8.1$ Hz, 1H), 7.11 (t, $J = 7.3$ Hz, 1H), 6.96 (d, $J = 7.7$ Hz, 1H), 5.22–5.10 (m, 1H), 3.97 (s, 3H), 3.09–2.98 (m, 2H), 2.42 (s, 3H), 2.14 (d, $J = 5.3$ Hz, 2H), 2.06–1.90 (m, 2H), 1.65 (s, 3H), 1.59 (s, 2H) and 1.00–0.88 (m, 3H) ppm.

^1H NMR (400 MHz, CDCl_3) minor isomer: δ 7.93 (d, $J = 7.6$ Hz, 1H), 7.74 (d, $J = 7.7$ Hz, 1H), 7.34 (dd, $J_1 = 7.6$ Hz and $J_2 = 8.0$ Hz, 1H), 7.27 (d, $J = 8.1$ Hz, 1H), 7.11 (t, $J = 7.3$ Hz, 1H), 6.96 (d, $J = 7.7$ Hz, 1H), 5.04–4.94 (m, 1H), 3.97 (s, 3H), 3.09–2.98 (m, 2H), 2.42 (s, 3H), 2.14 (d, $J = 5.3$ Hz, 2H), 2.06–1.90 (m, 2H), 1.65 (s, 3H), 1.59 (s, 2H) and 1.00–0.88 (m, 3H) ppm.

$^{13}\text{C}[^1\text{H}]$ NMR (100 MHz, CDCl_3) δ 142.2, 139.9, 138.2, 138.1, 134.6, 125.3, 124.1, 124.1, 123.6, 123.1, 122.6, 122.4, 119.6, 118.9, 117.6, 108.7, 32.6, 32.5, 31.9, 31.7, 28.2, 28.1, 28.0, 24.9, 23.0, 20.2, 16.2 and 13.0 ppm.

IR (ATR): 2966, 1445, 1438, 1390, 1268, 1146, 1121 and 748 cm^{-1} .

HRMS (ESI) m/z : [M + NH₄]⁺ calcd for $\text{C}_{22}\text{H}_{31}\text{N}_2$ 323.2482; found 323.2496 (4.3 ppm).

TLC: $R_f = 0.8$ (9 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(5-methylheptyl)-9*H*-carbazole (12b).

Following GP-III. Pd/C (4 mg, 10 wt%) was added to a stirring solution of olefin **11b** (37 mg, 0.12 mmol, 1.0 equiv.) in EtOAc (4 mL). The resulting reaction mixture was stirred under a hydrogen (1 atm) atmosphere for 5 h at room temperature. This reaction mixture was filtered and purified using silica gel

column chromatography (19 : 1 hexanes/EtOAc) to provide the desired hydrogenation product **12b** (34 mg, 0.11 mmol, 91%) as a colorless liquid.

^1H NMR (400 MHz, CDCl_3) δ 7.93 (d, $J = 7.5$ Hz, 1H), 7.75 (d, $J = 7.7$ Hz, 1H), 7.35 (t, $J = 7.5$ Hz, 1H), 7.27 (d, $J = 8.1$ Hz, 1H), 7.11 (t, $J = 7.2$ Hz, 1H), 6.97 (d, $J = 7.7$ Hz, 1H), 3.98 (s, 3H), 3.08–2.96 (m, 2H), 2.43 (s, 3H), 1.47 (m, 5H), 1.32 (m, 3H), 1.10–1.09 (m, 1H) and 0.81 (d, $J = 5.8$ Hz, 6H) ppm.

$^{13}\text{C}[^1\text{H}]$ NMR (100 MHz, CDCl_3) δ 142.2, 139.9, 134.5, 125.3, 124.3, 123.2, 122.7, 122.4, 119.6, 118.9, 117.6, 108.7, 36.6, 34.5, 32.6, 32.0, 29.7, 28.6, 27.7, 20.3, 19.4, 11.6 ppm.

IR (ATR): 3421, 2925, 2862, 1642, 1586, 1482, 1452, 1431, 1374, 1361, 1285, 1046, 923 and 722 cm^{-1} .

HRMS (ESI) m/z : [M + H]⁺ calcd for $\text{C}_{22}\text{H}_{30}\text{N}$ 308.2373; found 308.2382 (3.0 ppm).

TLC: $R_f = 0.45$ (hexane).

3-Bromo-2,9-dimethyl-1-(5-methylheptyl)-9*H*-carbazole (13b).

Following GP-IV. *N*-Bromo-succinimide (18 mg, 0.09 mmol, 1.0 equiv.) was added to a solution of carbazole **12b** (30 mg, 0.09 mmol, 1.0 equiv.) in chloroform (CHCl_3) (6 mL) under a nitrogen atmosphere. The reaction mixture was stirred for 6 min. After completion, water (10 mL) was added and extracted with CH_2Cl_2 (3×5 mL). The combined organic layers were dried and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (19 : 1 hexanes/EA) provided the desired product **13b** (34 mg, 0.08 mmol, 92%) as a gummy compound.

^1H NMR (400 MHz, CDCl_3) δ 8.08 (s, 1H), 7.90 (d, $J = 7.7$ Hz, 1H), 7.40 (t, $J = 7.6$ Hz, 1H), 7.29 (d, $J = 8.2$ Hz, 1H), 7.15 (t, $J = 7.1$ Hz, 1H), 3.97 (s, 3H), 3.14–3.02 (m, 2H), 2.52 (s, 3H), 1.60–1.45 (m, 5H), 1.32–1.31 (m, 3H), 1.20–1.15 (m, 1H) and 0.84 (d, $J = 5.7$ Hz, 6H) ppm.

$^{13}\text{C}[^1\text{H}]$ NMR (100 MHz, CDCl_3) δ 142.5, 139.1, 132.9, 126.0, 123.7, 122.1, 121.6, 119.8, 119.3, 116.8, 108.9, 36.5, 34.5, 32.8, 32.0, 29.6, 27.5, 19.8, 19.3, 11.6 ppm.

IR (ATR): 2934, 2868, 2376, 1468, 1402, 1282, 1149, 1016, 812, 762 and 742 cm^{-1} .

HRMS (ESI) m/z : [M + H]⁺ calcd for $\text{C}_{22}\text{H}_{29}\text{NBr}$ 386.1478; found 386.1460 (5 ppm).

TLC: $R_f = 0.5$ (hexane).

3-Methoxy-2,9-dimethyl-1-(5-methylheptyl)-9*H*-carbazole (14b).

Following GP-V. A freshly prepared NaOMe solution (~2.7 M in MeOH) was added to a DMF (1.6 mL) solution of 3-bromocarbazole **13b** (29 mg, 0.07 mmol, 1.0 equiv.), followed by CuI (59 mg, 0.23 mmol, 4.0 equiv.), and the reaction mixture stirred at 115 °C for 15 h. The reaction mixture was filtered, and the filtrate was sequentially washed with saturated NH_4Cl solution (5 mL), water (10 mL) and brine (5 mL), dried and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (9 : 1 hexanes/EtOAc) provided the desired 3-methoxycarbazole **14b** (23 mg, 0.06 mmol, 90%) as a colourless liquid.

^1H NMR (500 MHz, CDCl_3) δ 7.91 (d, $J = 7.7$ Hz, 1H), 7.36–7.30 (m, 2H), 7.27 (d, $J = 8.2$ Hz, 1H), 7.08 (t, $J = 7.3$ Hz, 1H), 3.96 (s, 3H), 3.86 (s, 3H), 3.07–3.01 (m, 2H), 2.29 (s, 3H), 1.62–1.42 (m, 5H), 1.36–1.25 (m, 3H), 1.11–0.85 (m, 1H) and 0.84–0.78 (m, 6H) ppm.



$^{13}\text{C}[\text{H}]$ NMR (125 MHz, CDCl_3) δ 152.3, 142.5, 134.8, 125.8, 125.1, 124.8, 123.3, 121.8, 119.5, 118.5, 108.8, 99.2, 56.3, 36.6, 34.6, 32.8, 32.0, 29.7, 28.8, 27.6, 19.4, 12.1 and 11.6 ppm.

IR (ATR): 2929, 2857, 2389, 1471, 1416, 1273, 1220, 1152, 1104, 838 and 741 cm^{-1} .

HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ calcd for $\text{C}_{23}\text{H}_{32}\text{NO}$ 338.2478; found 338.2469 (3.0 ppm).

TLC: R_f = 0.18 (hexane).

2,9-Dimethyl-1-(5-methylheptyl)-9H-carbazol-3-ol (15b).

Following GP-VI. To a stirring solution of 3-methoxycarbazole **14b** (21 mg, 0.06 mmol, 1.0 equiv.) in anhydrous CH_2Cl_2 (4 mL) at 0 $^\circ\text{C}$, a 1 M solution of boron tribromide (BBr_3) in CH_2Cl_2 (0.25 mL, 0.25 mmol, 4.0 equiv.) was added dropwise and further stirred for 8 h. The reaction was quenched with H_2O (5 mL). The organic layer was extracted with CH_2Cl_2 (3×5 mL). The combined organic layers were dried over (Na_2SO_4) and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (4 : 1 hexanes/EtOAc) provided the desired phenol **15b** (18.2 mg, 0.05 mmol, 91% yield) as a gummy compound.

^1H NMR (400 MHz, CDCl_3) δ 7.84 (d, J = 7.6 Hz, 1H), 7.33 (d, J = 7.4 Hz, 1H), 7.26 (s, 2H), 7.07 (t, J = 7.2 Hz, 1H), 3.95 (s, 3H), 3.08–2.96 (m, 2H), 2.31 (s, 3H), 1.63–1.40 (m, 5H), 1.28 (s, 2H) and 1.16–1.08 (m, 2H), 0.81–0.80 (m, 6H) ppm.

$^{13}\text{C}[\text{H}]$ NMR (100 MHz, CDCl_3) δ 148.1, 141.3, 135.4, 127.9, 126.0, 124.6, 123.5, 122.3, 121.2, 111.2, 110.3, 103.1, 39.2, 32.9, 31.7, 30.3, 28.7, 28.1, 27.4, 22.8, 12.0 and 11.2 ppm.

IR (ATR): 3326, 2922, 2862, 2396, 1469, 1296, 1245, 1077 and 743 cm^{-1} .

HRMS (ESI) m/z : $[\text{M} + \text{NH}_4]^+$ calcd for $\text{C}_{22}\text{H}_{33}\text{N}_2\text{O}$ 341.2587; found 341.2633 (13.4 ppm).

TLC: R_f = 0.4 (9 : 1, hexanes/EtOAc).

2,9-Dimethyl-1-(5-methylheptyl)-3H-carbazole-3,4(9H)-

dione-[N-methylcarbazoquinocin D] 4. Following GP-VII. A solution of carbazol-3-ol **15b** (15 mg, 0.05 mmol) and (PhSeO_2O)_{0.4} mg, 0.09 mmol, 2.0 equiv.) in THF (4 mL) was stirred at 50 $^\circ\text{C}$ for 30 min, the mixture was quenched with water, and the residual compound from the aqueous layer was extracted with EtOAc (3×4 mL). The combined organic layers were dried (Na_2SO_4) and concentrated *in vacuo*. Purification of crude product *via* silica gel column chromatography (3 : 1 hexanes/EtOAc) provided the desired *N*-methyl carbazoquinocin D 4 (14.2 mg, 0.04 mmol, 89% yield) as a dark-brown glittering solid.

^1H NMR (500 MHz, CDCl_3) δ 8.12–7.99 (m, 1H), 7.21–7.14 (m, 3H), 3.82 (s, 2H), 2.63 (t, J = 8.1 Hz, 2H), 1.84 (s, 3H), 1.57–1.41 (m, 4H), 1.33–1.24 (m, 3H), 1.15–1.05 (m, 2H) and 0.88–0.74 (m, 6H) ppm.

$^{13}\text{C}[\text{H}]$ NMR (125 MHz, CDCl_3) δ 183.3, 173.7, 144.9, 142.6, 139.6, 134.5, 125.8, 124.9, 121.9, 113.8, 110.8, 36.4, 34.5, 33.1, 29.9, 29.6, 28.7, 27.5, 19.3, 11.9 and 11.5 ppm.

IR (ATR): 2971, 2934, 2893, 1688, 1652, 1632, 1494, 1441, 1414, 1398, 1242 and 765 cm^{-1} .

HRMS (ESI) m/z : $[\text{M} + \text{Na}]^+$ calcd for $\text{C}_{22}\text{H}_{28}\text{NO}_2$ 338.2115; found 338.2116 (0.3 ppm).

TLC: R_f = 0.3 (8 : 1 hexanes/EtOAc).

M.P.: 148–150 $^\circ\text{C}$.

(Z)-2,9-Dimethyl-1-(6-methylhept-4-en-1-yl)-9H-carbazole (11c).

Following GP-II. $^t\text{BuOK}$ (41 mg, 0.33 mmol, 1.8 equiv.) was added to a solution of isobutyl (triphenyl phosphonium) bromide **10c** (128 mg, 0.38 mmol, 2.0 equiv.) in anhydrous THF (5 mL), followed by aldehyde **2** (50 mg, 0.18 mmol, 1.0 equiv.) and stirred for 12 h. The corresponding alkene product **11c** was isolated as a colorless gummy compound (46 mg, 0.15 mmol, 81%) using hexanes/EtOAc (9 : 1) as the eluent.

^1H NMR (500 MHz, CDCl_3) δ 7.91 (d, J = 7.5 Hz, 1H), 7.73 (d, J = 8.0 Hz, 1H), 7.32 (dd, J_1 = 7.0 Hz and J_2 = 8.0 Hz, 1H), 7.24 (d, J = 8.0 Hz, 1H), 7.08 (dd, J_1 = 7.5 Hz and J_2 = 1.0 Hz, 1H), 6.94 (d, J = 8.0 Hz, 1H), 5.27–5.18 (m, 2H), 3.94 (s, 3H), 3.03–3.00 (m, 2H), 2.60–2.58 (m, 1H), 2.41 (s, 3H), 2.17 (q, J = 7.5 Hz, 2H), 1.65–1.62 (m, 2H) and 0.89 (q, J = 2.5 Hz, 6H) ppm.

$^{13}\text{C}[\text{H}]$ NMR (125 MHz, CDCl_3) δ 142.3, 140.0, 138.7, 134.6, 126.5, 125.3, 124.0, 123.2, 122.8, 122.5, 119.7, 119.0, 117.7, 108.7, 32.7, 31.7, 28.1, 27.7, 26.7, 23.3 and 20.2 ppm.

IR (ATR): 2917, 1486, 1485, 1325, 1235 and 756 cm^{-1} .

HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ calcd for $\text{C}_{22}\text{H}_{28}\text{N}$ 306.2216; found 306.2212 (1.3 ppm).

TLC: R_f = 0.4 (19 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(6-methylheptyl)-9H-carbazole (12c). Following GP-III. Pd/C (4 mg, 10 wt%) was added to a stirred solution of olefin **11c** (40 mg, 0.13 mmol, 1.0 equiv.) in EtOAc (4 mL). The resulting reaction mixture was stirred under a hydrogen (1 atm) atmosphere for 4 h at room temperature. The reaction mixture was filtered and purification of the crude product *via* silica gel column chromatography (19 : 1 hexanes/EtOAc) provided the desired hydrogenation product **12c** (37 mg, 0.12 mmol, 93%) as a colorless liquid.

^1H NMR (500 MHz, CDCl_3) δ 8.04 (d, J = 7.5 Hz, 1H), 7.85 (dt, J_1 = 7.5 Hz, J_2 = 2.0 Hz, 1H), 7.45 (t, 1H), 7.37 (d, J = 8.0 Hz, 1H), 7.23–7.20 (m, 1H), 7.07 (d, J = 7.5 Hz, 1H), 4.08 (s, 3H), 3.14–3.13 (m, 2H), 2.53 (s, 3H), 1.72–1.69 (m, 2H), 1.56–1.53 (m, 1H), 1.52–1.51 (m, 2H), 1.43–1.41 (m, 2H) and 1.27–1.24 (m, 2H), 0.92–0.91 (m, 6H) ppm.

$^{13}\text{C}[\text{H}]$ NMR (125 MHz, CDCl_3) δ 142.3, 140.0, 134.5, 125.3, 124.3, 123.3, 122.8, 122.5, 119.6, 118.9, 117.6, 108.7, 39.2, 32.6, 31.7, 30.4, 28.5, 28.1, 27.4, 22.8 and 20.2 ppm.

IR (ATR): 3426, 2986, 2843, 1642, 1531, 1479, 1413, 1372, 1324, 1253, 1026, 866 and 732 cm^{-1} .

HRMS (ESI) m/z : $[\text{M} + \text{H}]^+$ calcd for $\text{C}_{22}\text{H}_{30}\text{N}$ 308.2373; found 308.2324 (16.0 ppm).

TLC: R_f = 0.4 (hexane).

3-Bromo-2,9-dimethyl-1-(6-methylheptyl)-9H-carbazole (13c).

Following GP-IV. *N*-Bromo-succinimide (20 mg, 0.11 mmol, 1.0 equiv.) was added to a solution of carbazole **12c** (35 mg, 0.11 mmol, 1.0 equiv.) in chloroform (CHCl_3) (6 mL) under a nitrogen atmosphere. The reaction mixture was stirred for 6 min. After completion, water (10 mL) was added and extracted with CH_2Cl_2 (3×5 mL). The combined organic layers were dried and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (19 : 1 hexanes/EA) provided the desired product **13c** (38 mg, 0.10 mmol, 91%) as a gummy compound.



¹H NMR (500 MHz, CDCl₃) δ 8.14 (s, 1H), 7.96 (d, *J* = 7.5 Hz, 1H), 7.47 (dd, *J*₁ = 7.5 Hz and *J*₂ = 1.0 Hz, 1H), 7.34 (d, *J* = 8.0 Hz, 1H), 7.21 (dd, *J*₁ = 8.0 Hz and *J*₂ = 7.0 Hz, 1H), 4.00 (s, 3H), 3.14–3.11 (m, 2H), 2.58 (s, 3H), 1.67–1.65 (m, 2H), 1.58–1.52 (m, 1H), 1.51–1.49 (m, 2H), 1.42–1.38 (m, 2H) and 1.26–1.23 (m, 2H), 0.91 (d, *J* = 6.5 Hz, 6H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 142.6, 139.1, 133.0, 126.0, 126.1, 123.8, 122.2, 121.6, 119.8, 119.3, 116.9, 108.9, 39.1, 32.8, 31.7, 30.3, 29.6, 28.1, 27.4, 22.8 and 19.8 ppm.

IR (ATR): 2923, 2857, 2396, 1462, 1406, 1274, 1134, 1056, 829, 762 and 746 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₂H₂₉NBr 386.1478; found 386.1432 (12 ppm).

TLC: *R*_f = 0.5 (hexane).

3-Methoxy-2,9-dimethyl-1-(6-methylheptyl)-9*H*-carbazole (14c).

Following GP-V. A freshly prepared NaOMe solution (~2.7 M in MeOH) was added to a DMF (1.6 mL) solution of 3-bromocarbazole **13c** (35 mg, 0.09 mmol, 1.0 equiv.), followed by CuI (72 mg, 0.36 mmol, 4.0 equiv.), and the reaction mixture stirred at 115 °C for 15 h. The reaction mixture was filtered, and the filtrate was sequentially washed with saturated NH₄Cl solution (5 mL), water (10 mL) and brine (5 mL), dried and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (9 : 1 hexanes/EtOAc) provided the desired 3-methoxycarbazole **14c** (31 mg, 0.09 mmol, 93%) as a yellowish liquid.

¹H NMR (500 MHz, CDCl₃) δ 7.97 (d, *J* = 7.5 Hz, 1H), 7.4–7.40 (m, 2H), 7.35 (d, *J* = 8 Hz, 1H), 7.16 (dd, *J*₁ = 7 Hz and *J*₂ = 7.5 Hz, 1H), 4.04 (s, 3H), 3.94 (s, 3H), 3.12 (t, 2H), 2.37 (s, 3H), 1.55–1.51 (m, 2H), 1.52–1.50 (m, 3H), 1.39–1.31 (m, 2H), 1.26–1.21 (m, 2H) and 0.88 (d, *J* = 6.5 Hz, 6H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 152.4, 142.5, 134.9, 125.8, 125.1, 124.8, 123.3, 121.8, 119.5, 118.5, 108.8, 99.3, 56.3, 39.2, 32.8, 31.7, 30.4, 28.8, 28.1, 27.4, 22.8 and 12.1 ppm.

IR (ATR): 2922, 2879, 2362, 1471, 1424, 1288, 1212, 1144, 1109, 834 and 732 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₃H₃₂NO 338.2478; found 338.2443 (10.3 ppm).

TLC: *R*_f = 0.2 (hexane).

2,9-Dimethyl-1-(6-methylheptyl)-9*H*-carbazol-3-ol (15c).

Following GP-VI. To a stirring solution of 3-methoxycarbazole **14c** (28 mg, 0.08 mmol, 1.0 equiv.) in anhydrous CH₂Cl₂ (4 mL) at 0 °C, a 1 M solution of boron tribromide (BBr₃) in CH₂Cl₂ (0.33 mL, 0.33 mmol, 4.0 equiv.) was added dropwise, and further stirred for 12 h. The reaction was quenched with H₂O (5 mL). The organic layer was extracted with CH₂Cl₂ (3 × 5 mL). The combined organic layers were dried (Na₂SO₄) and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (4 : 1 hexanes/EtOAc) provided the desired phenol **15c** (25.5 mg, 0.08 mmol, 95% yield) as a white gummy compound.

¹H NMR (500 MHz, CDCl₃) δ 7.91 (d, *J* = 8.0 Hz, 1H), 7.41 (dd, *J*₁ = 8.0 Hz and *J*₂ = 7.0 Hz, 1H), 7.34–7.33 (m, 2H), 7.14 (dd, *J*₁ = 7.5 Hz and *J*₂ = 7.0 Hz, 1H), 4.03 (s, 3H), 3.12 (t, 2H), 2.39 (s, 3H), 1.67–1.64 (m, 2H), 1.58–1.57 (m, 1H), 1.54–1.50 (m, 2H), 1.39–1.37 (m, 2H), 1.26–1.21 (m, 2H) and 0.88 (d, *J* = 6.5 Hz, 6H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 148.1, 141.3, 135.4, 127.9, 126.0, 124.6, 123.5, 122.3, 121.2, 111.2, 110.3, 103.1, 39.2, 32.9, 31.7, 30.3, 28.7, 28.1, 27.4, 22.8 and 12.2 ppm.

IR (ATR): 3345, 2934, 2859, 2359, 2346, 1485, 1449, 1277, 1236, 929, 782 and 734 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₂H₃₀NO 324.2322; found 324.2278 (13.5 ppm).

TLC: *R*_f = 0.2 (9 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(6-methylheptyl)-3*H*-carbazole-3,4(*9H*)dione-[*N*-methyl carbazoquinocin E] 5. Following GP-VII. A solution of carbazol-3-ol **15c** (7 mg, 0.02 mmol) and (PhSeO)₂O (16.0 mg, 0.04 mmol, 2.0 equiv.) in THF (4 mL) was stirred at 50 °C for 30 min. The mixture was quenched with water, and the residual compound from the aqueous layer was extracted with EtOAc (3 × 4 mL). The combined organic layers were dried (Na₂SO₄) and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (3 : 1 hexanes/EtOAc) provided the desired *N*-methyl carbazoquinocin E 5 (6.6 mg, 0.02 mmol, 91% yield) as a dark-brown glittering solid.

¹H NMR (500 MHz, CDCl₃) δ 8.09–8.08 (m, 1H), 7.20–7.20 (m, 3H), 3.84 (s, 3H), 2.66–2.64 (m, 2H), 1.88 (s, 3H), 1.58–1.52 (m, 3H), 1.47 (d, *J* = 6.6 Hz, 1H), 1.43–1.37 (m, 2H), 1.33–1.28 (m, 2H), 1.17–1.12 (m, 2H) and 0.81 (d, *J* = 6.5 Hz, 6H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 183.4, 173.8, 145.0, 142.6, 139.7, 134.6, 125.9, 124.9, 122.0, 113.9, 110.7, 39.0, 33.1, 30.2, 30.0, 28.5, 28.1, 27.2, 22.7 and 12.0 ppm.

IR (ATR): 2952, 2934, 2863, 1658, 1647, 1635, 1482, 1463, 1434, 1372, 1234 and 762 cm⁻¹.

HRMS (ESI) *m/z*: [M + Na]⁺ calcd for C₂₂H₂₇NNaO₂ 360.1934; found 360.1964 (8.3 ppm).

TLC: *R*_f = 0.4 (4 : 1 hexane/EtOAc).

M.P.: 156–158 °C.

2,9-Dimethyl-1-(7-methyloct-4-en-1-yl)-9*H*-carbazole (11d).

Following GP-II. ³BuOK (18 mg, 0.14 mmol, 1.8 equiv.) was added to a solution of (4-methylpentyl) triphenyl phosphonium bromide **10d** (80 mg, 0.18 mmol, 2.0 equiv.) in anhydrous THF (5 mL), followed by aldehyde **2** (25 mg, 0.09 mmol, 1.0 equiv.) and stirred for 12 h. The corresponding alkene product **11d** was isolated as a pale-yellow gummy compound (23 mg, 0.07 mmol, 80%) using hexane/EtOAc (9 : 1) as the eluent.

¹H NMR (500 MHz, CDCl₃) δ 7.91 (d, *J* = 7.5 Hz, 1H), 7.73 (d, *J* = 8.0 Hz, 1H), 7.33 (dd, *J*₁ = 8.5 Hz and *J*₂ = 7.0 Hz, 1H), 7.24–7.23 (m, 1H), 7.09 (dd, *J*₁ = 7.5 Hz and *J*₂ = 7.0 Hz, 1H), 6.95 (d, *J* = 8.0 Hz, 1H), 5.45–5.38 (m, 2H), 3.96 (s, 3H), 3.02 (dd, *J*₁ = 8.5 Hz and *J*₂ = 9.0 Hz, 2H), 2.41 (s, 3H), 2.16 (dd, *J*₁ = 7.0 Hz and *J*₂ = 7.0 Hz, 2H), 1.89 (t, *J* = 6.5 Hz, 2H), 1.70–1.61 (m, 2H), 1.60–1.51 (m, 1H) and 0.83 (d, *J* = 6.5 Hz, 6H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 142.2, 139.9, 134.6, 129.8, 129.6, 125.3, 124.0, 123.2, 122.7, 122.4, 119.6, 118.9, 117.7, 108.7, 36.6, 32.7, 31.5, 28.8, 28.1, 27.7, 22.6 and 20.2 ppm.

IR (ATR): 2917, 1465, 12463, 1330, 1245, 1096, 826 and 736 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₃H₃₀N 320.2373; found 320.2341 (10.0 ppm).

TLC: *R*_f = 0.4 (19 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(7-methyloctyl)-9*H*-carbazole (12d). Following GP-III. To a stirring solution of olefin **11d** (48 mg, 0.15 mmol, 1.0



equiv.) in EtOAc (4 mL), Pd/C (5 mg, 10 wt%) was added. The resulting reaction mixture was stirred under a hydrogen (1 atm) atmosphere for 6 h at room temperature. The reaction mixture was filtered and purification of the crude product *via* silica gel column chromatography (19 : 1 hexane/EtOAc) gave the desired hydrogenation product **12d** (46 mg, 0.14 mmol, 95.4%) as a colorless gummy compound.

¹H NMR (500 MHz, CDCl₃) δ 8.01 (d, *J* = 7.5 Hz, 1H), 7.82 (d, *J* = 8.0 Hz, 1H), 7.43 (dd, *J*₁ = 8.0 Hz and *J*₂ = 7.5 Hz, 1H), 7.36 (d, *J* = 8.5, 1H), 7.19 (dd, *J*₁ = 7.0 Hz and *J*₂ = 7.0 Hz, 1H), 7.05 (d, *J* = 8.0 Hz, 1H), 4.07 (s, 3H), 3.11 (dd, *J*₁ = 8.0 Hz and *J*₂ = 8.5 Hz, 2H), 2.5 (s, 3H), 1.72–1.65 (m, 2H), 1.51 (t, *J* = 6.7 Hz, 2H), 1.41–1.36 (m, 2H), 1.36–1.30 (m, 3H), 1.22–1.16 (m, 2H) and 0.86 (d, *J* = 6.5, 6H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 142.3, 140.0, 134.5, 125.3, 124.3, 123.3, 122.8, 122.5, 119.6, 119.0, 117.6, 108.7, 39.2, 32.7, 31.7, 30.2, 29.9, 28.5, 28.1, 27.5, 22.8 and 20.2 ppm.

IR (ATR): 3413, 2928, 2855, 1648, 1595, 1429, 1413, 1372, 1324, 1256, 1049, 862 and 734 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₃H₃₂N 322.2529; found 322.2499 (9.3 ppm).

TLC: *R*_f = 0.45 (hexane).

3-Bromo-2,9-dimethyl-1-(7-methyloctyl)-9H-carbazole (13d).

Following GP-IV. N-Bromo-succinimide (25.5 mg, 0.14 mmol, 1.0 equiv.) was added to a solution of carbazole **12d** (44 mg, 0.14 mmol, 1.0 equiv.) in chloroform (CHCl₃) (6 mL) under a nitrogen atmosphere. The reaction mixture was stirred for 6 min. After completion, water (10 mL) was added and extracted with CH₂Cl₂ (3 × 5 mL). The combined organic layers were dried and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (19 : 1 hexanes/EA) provided the desired product **13d** (53 mg, 0.13 mmol, 93%) as a gummy compound.

¹H NMR (500 MHz, CDCl₃) δ 8.14 (s, 1H), 7.96 (d, *J* = 8.0 Hz, 1H), 7.46 (dd, *J*₁ = 8.0 Hz and *J*₂ = 7.0 Hz, 1H), 7.34 (d, *J* = 8.0 Hz, 1H), 7.22 (t, *J* = 7.5 Hz, 1H), 3.9 (s, 3H), 3.11 (dd, *J*₁ = 8.0 Hz and *J*₂ = 8.5 Hz, 2H), 2.58 (s, 3H), 1.66 (dd, *J*₁ = 10.5 Hz and *J*₂ = 6.0 Hz, 2H), 1.55 (dt, *J*₁ = 14.9 Hz and *J*₂ = 7.0 Hz, 3H), 1.40 (dd, *J*₁ = 9.9 Hz and *J*₂ = 4.5 Hz, 2H), 1.37–1.33 (m, 2H), 1.26–1.19 (m, 2H) and 0.91 (d, *J* = 6.5 Hz, 6H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 142.5, 139.1, 132.9, 126.0, 126.0, 123.7, 122.1, 121.6, 119.8, 119.3, 116.2, 108.9, 39.2, 32.7, 31.7, 30.0, 29.9, 29.6, 28.1, 27.5, 22.8 and 19.8 ppm.

IR (ATR): 2961, 2856, 2377, 1462, 1402, 1276, 1132, 1012, 819, 774 and 735 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₃H₃₁BrN 400.1634; found 400.1636 (0.5 ppm).

TLC: *R*_f = 0.5 (hexane).

3-Methoxy-2,9-dimethyl-1-(6-methylheptyl)-9H-carbazole

(14d). Following GP-V. A freshly prepared NaOMe solution (~2.7 M in MeOH) was added to a DMF (1.6 mL) solution of 3-bromocarbazole **13d** (40 mg, 0.10 mmol, 1.0 equiv.), followed by CuI (79 mg, 0.40 mmol, 4.0 equiv.), and the reaction mixture stirred at 115 °C for 15 h. The reaction mixture was filtered, and the filtrate was sequentially washed with saturated NH₄Cl solution (5 mL), water (10 mL) and brine (5 mL), dried and concentrated *in vacuo*. Purification of the crude product *via*

silica gel column chromatography (9 : 1 hexanes/EtOAc) provided the desired 3-methoxycarbazole **14d** (32 mg, 0.09 mmol, 91%) as a colorless liquid.

¹H NMR (400 MHz, CDCl₃) δ 8.0 (d, *J* = 7.6 Hz, 1H), 7.44–7.41 (m, 2H), 7.37–7.36 (m, 1H), 7.17 (dd, *J*₁ = 7.2 Hz and *J*₂ = 7.6 Hz, 1H), 4.0 (s, 3H), 3.95 (s, 3H), 3.13 (dd, *J*₁ = 8.0 Hz and *J*₂ = 8.0 Hz, 2H), 2.83 (s, 3H), 1.68 (q, *J* = 8.2 Hz, 2H), 1.57–1.49 (m, 3H), 1.36 (q, *J* = 8.3 Hz, 4H), 1.21 (t, *J* = 7.1 Hz, 2H) and 0.89 (d, *J* = 6.4, 6H) ppm.

¹³C[¹H] NMR (100 MHz, CDCl₃) δ 152.2, 142.4, 134.7, 125.8, 125.1, 124.7, 123.2, 121.8, 119.5, 118.4, 108.8, 99.1, 56.2, 39.2, 32.8, 31.7, 30.1, 29.9, 28.8, 28.1, 27.5, 22.8 and 12.1 ppm.

IR (ATR): 2935, 2853, 2384, 1466, 1419, 1293, 1220, 1149, 1129, 844 and 765 cm⁻¹.

HRMS (ESI) *m/z*: [M + Na]⁺ calcd for C₂₄H₃₃NNaO 374.2454; found 374.2486 (8.5 ppm).

TLC: *R*_f = 0.4 (19 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(7-methyloctyl)-9H-carbazol-3-ol (15d). Following GP-VI. To a stirring solution of 3-methoxycarbazole **14d** (28 mg, 0.08 mmol, 1.0 equiv.) in anhydrous CH₂Cl₂ (4 mL) at 0 °C, a 1 M solution of boron tribromide (BBr₃) in CH₂Cl₂ (0.33 mL, 0.33 mmol, 4.0 equiv.) was added dropwise, and further stirred for 11 h. The reaction was quenched with H₂O (5 mL). The organic layer was extracted with CH₂Cl₂ (3 × 5 mL). The combined organic layers were dried (Na₂SO₄) and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (4 : 1 hexanes/EtOAc) provided the desired phenol **15d** (26.2 mg, 0.07 mmol, 93% yield) as a gummy compound.

¹H NMR (400 MHz, CDCl₃) δ 7.91 (d, *J* = 7.6 Hz, 1H), 7.42 (t, *J* = 7.6 Hz, 1H), 7.34 (s, 2H), 7.14 (dd, *J*₁ = 7.2 Hz and *J*₂ = 7.6 Hz, 1H), 4.03 (s, 3H), 3.11 (dd, *J*₁ = 8.0 Hz and *J*₂ = 8.4 Hz, 2H), 2.39 (s, 3H), 1.67 (t, *J* = 7.8 Hz, 2H), 1.56–1.49 (m, 3H), 1.35 (dt, *J*₁ = 16.1 Hz and *J*₂ = 7.4 Hz, 4H), 1.22–1.17 (m, 2H) and 0.88 (d, *J* = 6.4, 6H) ppm.

¹³C[¹H] NMR (100 MHz, CDCl₃) δ 146.6, 141.5, 133.9, 124.6, 124.2, 121.6, 121.3, 121.0, 118.5, 117.3, 107.6, 101.9, 38.0, 31.6, 30.6, 29.0, 28.7, 27.6, 26.9, 26.4, 21.6 and 11.0 ppm.

IR (ATR): 3342, 2934, 2863, 2372, 2351, 1492, 1469, 1297, 1270, 902, 776 and 734 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₃H₃₂NO 338.2478; found 338.2485 (2.0 ppm).

TLC: *R*_f = 0.3 (9 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(6-methylheptyl)-3H-carbazole-3,4(9H)-dione-[N-methyl carbazoquinolin-F] (6). Following GP-VII. A solution of carbazol-3-ol **15d** (21 mg, 0.06 mmol) and (PhSeO)₂O (45.0 mg, 0.12 mmol, 2.0 equiv.) in THF (4 mL) was stirred at 50 °C for 30 min. The reaction mixture was quenched with water, and the residual compound from the aqueous layer was extracted with EtOAc (3 × 4 mL). The combined organic layers were dried (Na₂SO₄) and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (3 : 1 hexanes/EtOAc) provided the desired *N*-methyl carbazoquinolin F **6** (20.6 mg, 0.06 mmol, 94% yield) as a dark-brown glittering solid.

¹H NMR (400 MHz, CDCl₃) δ 8.06–8.05 (m, 1H), 7.18 (s, 3H), 3.82 (s, 3H), 2.62 (dd, *J*₁ = 6.8 Hz and *J*₂ = 8.0 Hz, 2H), 1.84 (s,



3H), 1.54–1.48 (m, 2H), 1.47–1.38 (m, 3H), 1.27 (s, 4H), 1.14–1.07 (m, 2H) and 0.80 (d, J = 6.4 Hz, 6H) ppm.

$^{13}\text{C}[\text{H}]$ NMR (100 MHz, CDCl_3) δ 183.3, 173.6, 144.9, 142.7, 139.6, 134.5, 125.8, 124.9, 121.9, 113.8, 110.8, 77.5, 77.2, 76.8, 39.0, 33.1, 30.0, 29.9, 29.7, 28.4, 28.3, 28.1, 27.4, 22.8 and 11.9 ppm.

IR (ATR): 2961, 2934, 2846, 1661, 1655, 1642, 1489, 1472, 1424, 1368, 1262 and 742 cm^{-1} .

HRMS (ESI) m/z : [M + H]⁺ calcd for $\text{C}_{23}\text{H}_{30}\text{NO}_2$ 352.2271; found 352.2248 (6.5 ppm).

TLC: R_f = 0.4 (4 : 1 hexane/EtOAc).

M.P.: 169–171 °C.

Ethyl 4-(2,9-dimethyl-9*H*-carbazol-1-yl)-2-methylbutanoate (17).

Following the reported procedure,¹⁰ under an N_2 atmosphere, a flame-dried round-bottom flask was charged with a solution of ethyl 4-(2,9-dimethyl-9*H*-carbazol-1-yl)butanoate **16** (50 mg, 0.16 mmol, 1.0 equiv.) in anhydrous THF (5 mL) and LDA (37 mg, 0.32 mmol, 2.0 equiv.) was added at 0 °C portion-wise and stirred for 30 min. This was followed by the dropwise addition of methyl iodide (46 mg, 0.32 mmol, 2.0 equiv.) at 0 °C, and then the reaction was stirred for 6 h at room temperature until TLC showed the complete consumption of the ester. After completion, a saturated NH_4Cl solution (6 mL) was added to quench the reaction at 0 °C. Then ethyl acetate (5 mL) and water (5 mL) were added. After separation of the layers, the residual compound from the aqueous layer was extracted with EtOAc (3 × 5 mL). The combined organic layers were dried over anhydrous Na_2SO_4 , filtered, and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (hexane/EA) provided the corresponding ethyl 4-(2,9-dimethyl-9*H*-carbazol-1-yl)-2-methylbutanoate **17** (31 mg, 0.09 mmol, 59% yield) as a pale-yellow color liquid.

^1H NMR (400 MHz, CDCl_3) δ 7.94 (d, J = 7.7 Hz, 1H), 7.77 (d, J = 7.8 Hz, 1H), 7.36 (t, J = 7.6 Hz, 1H), 7.29 (d, J = 8.2 Hz, 1H), 7.12 (t, J = 7.4 Hz, 1H), 6.98 (d, J = 7.8 Hz, 1H), 4.13 (q, J = 7.1 Hz, 2H), 4.00 (s, 3H), 3.16–2.97 (m, 2H), 2.59 (dd, J_1 = 13.4 Hz and J_2 = 7.1 Hz, 1H), 2.43 (s, 3H), 1.96–1.94 (m, 1H), 1.76–1.66 (m, 1H) and 1.28–1.18 (m, 6H) ppm.

$^{13}\text{C}[\text{H}]$ NMR (100 MHz, CDCl_3) δ 176.5, 142.2, 139.9, 134.7, 125.4, 123.1, 123.1, 122.8, 122.5, 119.7, 119.0, 117.9, 108.7, 60.6, 40.2, 35.3, 32.7, 26.2, 20.1, 17.5 and 14.5 ppm.

IR (ATR): 2944, 1738, 1580, 1453, 1218, 1199, 1153, 1127, 1026, 925, 827 and 749 cm^{-1} .

HRMS (ESI) m/z : [M + H]⁺ calcd for $\text{C}_{23}\text{H}_{32}\text{N}$ 324.1958; found 324.1988 (9.2 ppm).

TLC: R_f = 0.6 (9 : 1 hexane/EtOAc).

4-(2,9-Dimethyl-9*H*-carbazol-1-yl)-2-methylbutan-1-ol (18).

Under an N_2 atmosphere, a flame-dried round-bottom flask was charged with a solution of 4-(2,9-dimethyl-9*H*-carbazol-1-yl)-2-methylbutanoate **17** (62 mg, 0.19 mmol, 1.0 equiv.) in anhydrous THF (5 mL) and LAH (15 mg, 0.38 mmol, 2.0 equiv.) was added at 0 °C portion-wise and stirred for 4 h at room temperature until TLC showed the complete consumption of the ester. After completion, a saturated NH_4Cl solution (6 mL) was added to quench the reaction at 0 °C. EtOAc (5 mL) and water (5 mL) were added. After separation of the layers, the residual compound from the aqueous layer was extracted with

EtOAc (3 × 5 mL). The combined organic layers were dried over anhydrous Na_2SO_4 , filtered, and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (hexanes/EA) provided the corresponding ethyl 4-(2,9-dimethyl-9*H*-carbazol-1-yl)-2-methylbutan-1-ol **18** (50 mg, 0.18 mmol, 93% yield) as a colourless gummy compound.

^1H NMR (500 MHz, CDCl_3) δ 7.92 (d, J = 7.7 Hz, 1H), 7.74 (d, J = 7.9 Hz, 1H), 7.34 (t, J = 7.8 Hz, 1H), 7.25 (d, J = 8.3 Hz, 1H), 7.11 (d, J = 7.5 Hz, 1H), 6.96 (d, J = 7.9 Hz, 1H), 3.95 (s, 3H), 3.53–3.41 (m, 2H), 3.12–2.94 (m, 2H), 2.41 (s, 3H), 1.76–1.56 (m, 2H), 1.52 (s, 1H), 1.42 (dt, J_1 = 13.3 Hz and J_2 = 5.8 Hz, 1H) and 1.00 (d, J = 6.6 Hz, 3H) ppm.

$^{13}\text{C}[\text{H}]$ NMR (125 MHz, CDCl_3) δ 142.2, 139.9, 134.4, 125.3, 123.9, 123.1, 122.8, 122.5, 119.6, 118.9, 117.7, 108.7, 68.0, 36.5, 34.9, 32.6, 26.0 and 20.1, 16.7 ppm.

IR (ATR): 3360, 2944, 1738, 1580, 1453, 1218, 1199, 1153, 1127, 1026, 925, 827 and 749 cm^{-1} .

HRMS (ESI) m/z : [M + H]⁺ calcd for $\text{C}_{19}\text{H}_{23}\text{NO}$ 281.1780; found 281.1792 (4.2 ppm).

TLC: R_f = 0.2 (9 : 1 hexane/EtOAc).

4-(2,9-Dimethyl-9*H*-carbazol-1-yl)-2-methylbutanal (19).

Under an N_2 atmosphere, a flame-dried round-bottom flask was charged with a solution of ethyl 4-(2,9-dimethyl-9*H*-carbazol-1-yl)-2-methylbutan-1-ol **18** (42 mg, 0.14 mmol, 1.0 equiv.) in anhydrous dichloromethane (5 mL) and Dess–Martin periodate (82 mg, 0.19 mmol, 1.3 equiv.) was added at 0 °C portion-wise and stirred for 6 h at room temperature until TLC showed the complete consumption of the alcohol. After completion, the reaction was filtered and washed with dichloromethane and dried over anhydrous Na_2SO_4 and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (hexanes/EtOAc) provided the corresponding ethyl 4-(2,9-dimethyl-9*H*-carbazol-1-yl)-2-methylbutanal **19** (38 mg, 0.13 mmol, 91% yield) as a colorless liquid.

^1H NMR (400 MHz, CDCl_3) δ 9.57 (s, 1H), 7.89 (d, J = 7.6 Hz, 1H), 7.72 (d, J = 7.8 Hz, 1H), 7.32 (t, J = 7.5 Hz, 1H), 7.21 (d, J = 8.2 Hz, 1H), 7.09 (t, J = 7.2 Hz, 1H), 6.93 (d, J = 7.8 Hz, 1H), 3.87 (s, 3H), 3.07–2.90 (m, 2H), 2.37 (s, 4H), 1.88 (dd, J_1 = 13.1 Hz and J_2 = 5.6 Hz, 1H), 1.64–1.50 (m, 1H) and 1.12 (d, J = 7.1 Hz, 3H) ppm.

$^{13}\text{C}[\text{H}]$ NMR (100 MHz, CDCl_3) δ 204.4, 142.2, 139.8, 134.5, 125.4, 123.0, 122.8, 122.5, 119.6, 119.0, 117.9, 108.7, 46.7, 32.6, 31.9, 25.7, 20.1 and 13.6 ppm.

IR (ATR): 2976, 2941, 1725, 1673, 1622, 1468, 1444, 1433, 1266, 842, 823 and 746 cm^{-1} .

HRMS (ESI) m/z : [M + H]⁺ calcd for $\text{C}_{19}\text{H}_{22}\text{NO}$ 380.1696; found 380.1725 (8.0 ppm).

TLC: R_f = 0.4 (19 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(3-methylpent-4-en-1-yl)-9*H*-carbazole (20).

Following GP-II. $^3\text{BuOK}$ (37 mg, 0.30 mmol, 1.8 equiv.) was added to a solution of methyl(triphenyl phosphonium) bromide (120 mg, 0.33 mmol, 2.0 equiv.) in anhydrous THF (5 mL), followed by aldehyde **19** (50 mg, 0.16 mmol, 1.0 equiv.) and stirred for 12 h. The corresponding desired alkene product **20** was isolated as a colorless gummy compound (39 mg, 0.14 mmol, 84%) using hexane/EtOAc (9 : 1) as the eluent.



¹H NMR (400 MHz, CDCl₃) δ 7.88 (d, *J* = 7.6 Hz, 1H), 7.70 (d, *J* = 7.8 Hz, 1H), 7.30 (t, *J* = 7.1 Hz, 1H), 7.19 (d, *J* = 8.2 Hz, 1H), 7.11–7.03 (m, 1H), 6.92 (d, *J* = 7.6 Hz, 1H), 5.71 (dt, *J*₁ = 17.0 Hz and *J*₂ = 8.5 Hz, 1H), 4.96–4.62 (m, 2H), 3.86 (s, 3H), 3.04–2.82 (m, 2H), 2.36 (s, 3H), 2.29–2.16 (m, 1H), 1.53 (d, *J* = 6.6 Hz, 2H) and 1.01 (dd, *J*₁ = 3.7 Hz and *J*₂ = 2.9 Hz, 3H) ppm.

¹³C[¹H] NMR (100 MHz, CDCl₃) δ 144.0, 142.2, 139.9, 134.5, 125.3, 123.9, 123.1, 122.7, 122.5, 119.6, 118.9, 117.6, 113.6, 108.6, 38.7, 38.2, 32.7, 26.2, 20.3 and 20.2 ppm.

IR (ATR): 2917, 1496, 1475, 1325, 1308, 1235, 842 and 738 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₀H₂₆N 278.1903; found 278.1894 (3.2 ppm).

TLC: *R*_f = 0.4 (19 : 1 hexane/EtOAc).

2,9-Dimethyl-1-(3-methylpentyl)-9*H*-carbazole (21). Following GP-III. Pd/C (4.4 mg, 10 wt%) was added to a stirred solution of olefin **20** (44 mg, 0.16 mmol, 1.0 equiv.) in EtOAc (4 mL). The resulting reaction mixture was stirred under a hydrogen (1 atm) atmosphere for 3 h at room temperature. The reaction mixture was filtered and purification of the crude product *via* silica gel column chromatography (19 : 1 hexanes/EtOAc) provided the desired hydrogenation product **21** (44.5 mg, 0.16 mmol, 96%) as a colorless liquid.

¹H NMR (400 MHz, CDCl₃) δ 7.90 (d, *J* = 7.4 Hz, 1H), 7.72 (dd, *J*₁ = 7.8 Hz and *J*₂ = 1.6 Hz, 1H), 7.31 (t, *J* = 7.6 Hz, 1H), 7.22 (d, *J* = 8.2 Hz, 1H), 7.08 (t, *J* = 6.4 Hz, 1H), 6.94 (d, *J* = 7.7 Hz, 1H), 3.91 (s, 3H), 3.08–2.87 (m, 2H), 2.39 (s, 3H), 1.64–1.53 (m, 1H), 1.51–1.31 (m, 3H), 1.23–1.11 (m, 1H), 0.94 (dd, *J*₁ = 6.3 Hz and *J*₂ = 1.6 Hz, 3H) and 0.84–0.81 (m, 3H) ppm.

¹³C[¹H] NMR (100 MHz, CDCl₃) δ 142.2, 139.9, 134.4, 125.2, 124.4, 123.2, 122.7, 122.5, 119.6, 118.9, 117.5, 108.6, 38.5, 35.5, 32.6, 29.6, 26.3, 20.1, 19.3 and 11.7 ppm.

IR (ATR): 3423, 2932, 2889, 1651, 1532, 1449, 1431, 1372, 1334, 1243, 1089, 834 and 748 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₀H₂₆N 280.2060; found 280.2063 (1.0 ppm).

TLC: *R*_f = 0.4 (hexane).

3-Bromo-2,9-dimethyl-1-(3-methylpentyl)-9*H*-carbazole (22). Following GP-IV. N-Bromo-succinimide (25.4 mg, 0.14 mmol, 1.0 equiv.) was added a solution of carbazole **21** (40 mg, 0.14 mmol, 1.0 equiv.) in chloroform (CHCl₃) (6 mL) under a nitrogen atmosphere. The reaction mixture was stirred for 6 min. After completion, water (10 mL) was added and extracted with CH₂Cl₂ (3 × 5 mL). The combined organic layers were dried and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (19 : 1 hexanes/EtOAc) provided the corresponding 3-bromocarbazole **22** (45 mg, 0.12 mmol, 88%) as a colorless liquid.

¹H NMR (400 MHz, CDCl₃) δ 7.99 (s, 1H), 7.82 (s, 1H), 7.31 (t, *J* = 7.6 Hz, 1H), 7.17 (d, *J* = 8.2 Hz, 1H), 7.07 (t, *J* = 7.4 Hz, 1H), 3.80 (s, 3H), 3.03–2.83 (m, 2H), 2.42 (s, 3H), 1.58–1.47 (m, 1H), 1.43 (dd, *J*₁ = 12.2 Hz and *J*₂ = 6.0 Hz, 1H), 1.37–1.27 (m, 2H), 1.15 (dt, *J*₁ = 14.2 Hz and *J*₂ = 7.3 Hz, 1H), 0.92 (d, *J* = 6.4 Hz, 3H) and 0.83 (t, *J* = 7.3 Hz, 3H) ppm.

¹³C[¹H] NMR (100 MHz, CDCl₃) δ 142.4, 139.0, 132.8, 126.0, 126.0, 123.6, 122.1, 121.5, 119.8, 119.2, 116.8, 108.8, 38.4, 35.5, 32.7, 29.6, 27.3, 19.7, 19.2 and 11.7 ppm.

IR (ATR): 2969, 2869, 2374, 1463, 1416, 1291, 1186, 1055, 832, 768 and 749 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₀H₂₅NBr 358.1165; found 358.1159 (2.0 ppm).

TLC: *R*_f = 0.5 (hexane).

3-Methoxy-2,9-dimethyl-1-(3-methylpentyl)-9*H*-carbazole

(23). Following GP-V. A freshly prepared NaOMe solution (~2.7 M in MeOH) was added to a DMF (1.6 mL) solution of 3-bromocarbazole **22** (40 mg, 0.11 mmol, 1.0 equiv.), followed by CuI (93 mg, 0.46 mmol, 4.0 equiv.), and the reaction mixture stirred at 115 °C for 15 h. The reaction mixture was filtered, and the filtrate was sequentially washed with saturated NH₄Cl solution (5 mL), water (10 mL) and brine (5 mL), dried and concentrated *in vacuo*. Purification of the crude product *via* silica gel column chromatography (9 : 1 hexanes/EtOAc) provided the desired 3-methoxycarbazole **23** (29.6 mg, 0.09 mmol, 86%) as a pale-yellow liquid.

¹H NMR (400 MHz, CDCl₃) δ 7.90 (d, *J* = 7.7 Hz, 1H), 7.32 (t, *J* = 7.4 Hz, 2H), 7.25 (d, *J* = 8.2 Hz, 1H), 7.09 (d, *J* = 7.3 Hz, 1H), 3.95 (s, 3H), 3.85 (s, 3H), 3.14–2.93 (m, 2H), 2.28 (s, 3H), 1.66–1.56 (m, 1H), 1.53–1.44 (m, 2H), 1.42–1.37 (m, 1H), 1.20 (dd, *J*₁ = 13.7 Hz and *J*₂ = 6.7 Hz, 1H), 0.97 (d, *J* = 6.4 Hz, 3H) and 0.86 (t, *J* = 7.3 Hz, 3H) ppm.

¹³C[¹H] NMR (100 MHz, CDCl₃) δ 152.3, 142.4, 134.8, 125.9, 125.1, 124.6, 123.2, 121.8, 119.5, 118.4, 108.8, 99.1, 56.2, 38.4, 35.6, 32.8, 29.7, 26.6, 19.3, 11.9, and 11.7 ppm.

IR (ATR): 2935, 2862, 2388, 1476, 1413, 1278, 1229, 1164, 1122, 861 and 736 cm⁻¹.

HRMS (ESI) *m/z*: [M + H]⁺ calcd for C₂₁H₂₈NO 310.2165; found 310.2143 (7.0 ppm).

TLC: *R*_f = 0.2 (hexane).

2,9-Dimethyl-1-(3-methylpentyl)-9*H*-carbazol-3-ol (24).

Following GP-VI. To a stirring solution of 3-methoxycarbazole **23** (27 mg, 0.09 mmol, 1.0 equiv.) in anhydrous CH₂Cl₂ (4 mL) at 0 °C, a 1 M solution of boron tribromide (BBr₃, in CH₂Cl₂; 0.35 mL, 0.34 mmol, 4.0 equiv.) was added and further stirred for 11 h. The reaction was quenched with H₂O (5 mL) and extracted with CH₂Cl₂ (3 × 5 mL). The combined organic layers were dried (Na₂SO₄) and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (4 : 1 hexanes/EtOAc) provided phenol **24** (18 mg, 0.06 mmol, 70% yield) as a light-yellow gummy compound.

¹H NMR (500 MHz, CDCl₃) δ 7.92 (d, *J* = 7.6 Hz, 1H), 7.42 (t, *J* = 7.9 Hz, 1H), 7.39–7.31 (m, 2H), 7.16 (d, *J* = 6.5 Hz, 1H), 4.69 (s, 1H), 4.04 (s, 3H), 3.13–3.01 (m, 2H), 2.40 (s, 3H), 1.69 (dd, *J*₁ = 10.7 Hz and *J*₂ = 5.6 Hz, 1H), 1.63–1.56 (m, 1H), 1.55–1.43 (m, 2H), 1.33–1.25 (m, 1H), 1.07 (dd, *J*₁ = 6.7 Hz and *J*₂ = 2.7 Hz, 3H) and 0.96 (t, *J* = 7.4 Hz, 3H) ppm.

¹³C[¹H] NMR (125 MHz, CDCl₃) δ 147.8, 142.7, 135.1, 125.8, 125.4, 122.8, 122.4, 122.2, 119.7, 118.4, 108.7, 103.1, 38.5, 35.5, 32.8, 29.7, 26.6, 19.3, 12.0 and 11.7 ppm.

IR (ATR): 3325, 2933, 2854, 2362, 2346, 1485, 1448, 1272, 1236, 921, 776 and 731 cm⁻¹.

HRMS (ESI) *m/z*: [M + NH₄]⁺ calcd for C₂₀H₂₉N₂O 313.2274; found 313.2246 (9.0 ppm).

TLC: *R*_f = 0.3 (9 : 1 hexane/EtOAc).



2,9-Dimethyl-1-(3-methylpentyl)-3*H*-carbazole-3,4(*9H*)-dione [N-methylcarbazoquinocin A] (7). Following GP-VII. A solution of carbazol-3-ol **24** (15 mg, 0.05 mmol) and $(\text{PhSeO})_2\text{O}$ (36.5 mg, 0.10 mmol, 2.0 equiv.) in THF (4 mL) was stirred at 50 °C for 30 min. The mixture was quenched with water, and the residual compound from the aqueous layer was extracted with EtOAc (3 × 5 mL). The combined organic layers were dried (Na_2SO_4) and concentrated *in vacuo*. Purification of the crude product using silica gel column chromatography (3 : 1 hexanes/EtOAc) provided the desired *N*-methyl carbazoquinocin A **7** (14 mg, 0.04 mmol 90% yield) as a dark-brown glittering solid.

^1H NMR (400 MHz, CDCl_3) δ 8.13–8.11 (m, 1H), 7.25 (s, 3H), 3.88 (s, 3H), 2.81–2.57 (m, 2H), 1.90 (s, 3H), 1.50 (td, $J_1 = 13.2$ Hz and $J_2 = 6.5$ Hz, 2H), 1.43 (d, $J = 5.2$ Hz, 1H), 1.29–1.23 (m, 2H), 1.02 (d, $J = 6.2$ Hz, 3H) and 0.94 (t, $J = 7.3$ Hz, 3H) ppm.

$^{13}\text{C}[^1\text{H}]$ NMR (100 MHz, CDCl_3) δ 183.2, 173.7, 144.9, 143.0, 139.6, 134.5, 125.8, 124.9, 121.9, 113.8, 110.8, 35.4, 35.0, 33.1, 29.5, 27.9, 19.0, 11.8 and 11.6 ppm.

IR (ATR): 2966, 2957, 2853, 1662, 1658, 1636, 1492, 1471, 1454, 1379, 1235 and 776 cm^{-1} .

HRMS (ESI) *m/z*: [M + H]⁺ calcd for $\text{C}_{20}\text{H}_{24}\text{NO}_2$ 310.1802; found 310.1807 (2.0 ppm).

TLG: $R_f = 0.35$ (4 : 1 hexane/EtOAc).

M.P.: 149–151 °C.

Synthesis of isopropyltriphenylphosphonium bromide (10a).

Following GP-VIII. A solution of triphenylphosphine (1.0 g, 3.7 mmol) and isopropyl bromide (0.5 g, 4.12 mmol) was stirred at 110 °C for 48 h. The reaction was filtered and washed with ether to afford 1.1 g (76%) of product **10a** as a white solid.

Synthesis of sec-butyltriphenylphosphonium bromide (10b).

Following GP-VIII. A solution of triphenylphosphine (1.1 g, 4.19 mmol) and sec-butyl bromide (0.63 g, 4.61 mmol) was stirred at 110 °C for 48 h. The reaction was filtered and washed with ether to afford 540 mg (33%) of product **10b** as an off-white solid.

Synthesis of isobutyltriphenylphosphonium bromide (10c).

Following GP-VIII. A solution of triphenylphosphine (1.1 g, 4.19 mmol) and 1-bromo-2-methylpropane (0.63 g, 4.61 mmol) was stirred at 110 °C for 48 h. The reaction was filtered and washed with ether to afford 1.2 g (72%) of product **10c** as a white solid.

Synthesis of isopentyltriphenylphosphonium bromide (10d).

Following GP-VIII. A solution of triphenylphosphine (1.1 g, 4.19 mmol) and 1-bromo-3-methylbutane (0.69 g, 4.61 mmol) was stirred at 110 °C for 48 h. The reaction was filtered and washed with ether to afford 1.5 g (86%) of product **10d** as a white solid.

Synthesis of methyltriphenylphosphonium bromide.

Following GP-VIII. A solution of triphenylphosphine (1.1 g, 4.19 mmol) and methyl bromide (0.65 g, 4.61 mmol) was stirred at 110 °C for 48 h. The reaction was filtered and washed with ether to afford 1.4 g (93%) of the product as a white solid.

Abbreviations

Pd/C	Palladium on carbon
wt%	Weight%
atm	Atmosphere
NBS	<i>N</i> -Bromosuccinimide

DIBAL-H	Diisobutylaluminium hydride
<i>n</i> -BuLi	<i>n</i> -Butyllithium
equiv.	Equivalents
Cat.	Catalytic amount
M	Molar
anhyd.	Anhydrous
mL	Microliter
rt	Room temperature
TLC	Thin layer chromatography
R_f	Retardation or retention factor
NaHCO_3	Sodium bicarbonate
Na_2SO_4	Sodium sulfate
dr	Diastereomeric ratio
NMR	Nuclear magnetic resonance spectroscopy
IR	Infrared spectroscopy
ATR	Attenuated total reflection
HRMS (ESI)	High-resolution electrospray ionization mass spectrometry
Calcd	Calculated
M.P.	Melting point
ppm	Parts per million
^nBu	<i>n</i> -Butyl group
atm	Atmosphere

Data availability

The data underlying this study are available in the published article and its ESI.[†]

Conflicts of interest

The authors declare no competing financial interest.

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