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2025

Seankongsuk, P., Sae-Lao, P., He, S., Pereira, V. & Bates, R. W. (2025). Synthesis of dihydrogambirtannine by alkyne [2 + 2 + 2]-cyclotrimerisation. *Organic & Biomolecular Chemistry*. <https://dx.doi.org/10.1039/D5OB00162E>

<https://hdl.handle.net/10356/182986>

<https://doi.org/10.1039/D5OB00162E>

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Cite this: DOI: 10.1039/d5ob00162e

Received 28th January 2025,
Accepted 20th February 2025

DOI: 10.1039/d5ob00162e

rs.c.li/obc

Synthesis of dihydrogambirtannine by alkyne [2 + 2 + 2]-cyclotrimerisation†

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The first asymmetric synthesis of (–)-dihydrogambirtannine has been achieved using nucleophilic addition to a chiral sulfinyl imine to establish the stereogenic centre and [2 + 2 + 2] alkyne cyclotrimerisation to construct the aromatic E ring.

Introduction

Dihydrogambirtannine **1** is one of the yohimbine alkaloids,¹ distinguished by the fact that the E ring is aromatic. It was isolated from *Uncaria gambier*,² a plant that was formerly of economic importance in South East Asia as its extracts were used in dyeing and tanning. It has also been isolated from *Ochrosia borbonica*, a tree native to Mauritius and Réunion.³ The enantiomer of natural dihydrogambirtannine had earlier been prepared by degradation of deserpideine.⁴ (±)-Dihydrogambirtannine has been synthesized by Beisler,⁵ and by Wenkert *et al.*⁶ Each of these syntheses employed a late stage formation of the C-ring. (–)-Dihydrogambirtannine has been obtained as a minor product during transformations of secologanin.⁷ The analog lacking the methoxycarbonyl group has also been synthesized in racemic form.^{8,9} To the best of our knowledge, there has been no asymmetric synthesis of this alkaloid.

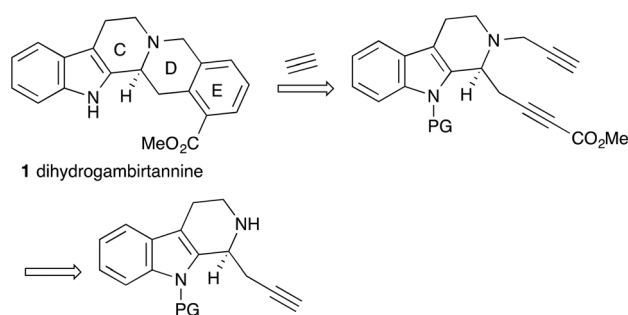
Results and discussion

A challenge in the synthesis of dihydrogambirtannine **1** is the fact that the E-ring is a 1,2,3-trisubstituted benzene ring. One effective approach to the synthesis of this motif is by the [2 + 2 + 2]-cyclotrimerisation of alkynes, in which one partner is acetylene (Scheme 1).^{10,11} The other partner would be a suitably substituted diyne, derived, in turn, from a functionalised indole. We wish to report the successful application of this strategy to the asymmetric synthesis of this alkaloid. We note that the Knölker synthesis of the desmethoxycarbonyl analog⁹

does follow this strategy, but in a stepwise fashion and *via* an organoiron intermediate.

One way to introduce the required stereogenic centre would be by asymmetric propargylation of the known imine **3** (Scheme 2). The imine was prepared from tryptamine **2** by the reported method,¹² and subjected to asymmetric propargylation using the asymmetric catalyst reported by Snyder *et al.*¹³ In our hands, the results were capricious and poorly reproducible. In addition, we note that this reaction can also require a very high loading of a complex ligand.¹⁴

We, therefore, turned to a more practical method (Scheme 3) using an imine bearing a sulfinyl chiral auxiliary,¹⁵ applying the propargylation reaction reported by Subba Reddy *et al.*¹⁶ Alcohol **6** was prepared by reduction of indole-3-acetic acid **5** and *N*-protected with a *p*-methoxybenzyl group using the conditions developed by Ley and Heaney.¹⁷ Alcohol **7** was then subjected to Vilsmeier–Haack formylation to give aldehyde **8** in modest yield. Aldehyde **8** was then converted to imine **9** under the reported conditions.¹⁶ Propargylation of imine **9** under carefully controlled conditions¹⁸ delivered a 9 : 1 mixture of diastereoisomers which could be separated by careful chromatography. Use of dichloromethane as the solvent for this reaction at –43 °C was essential to achieving

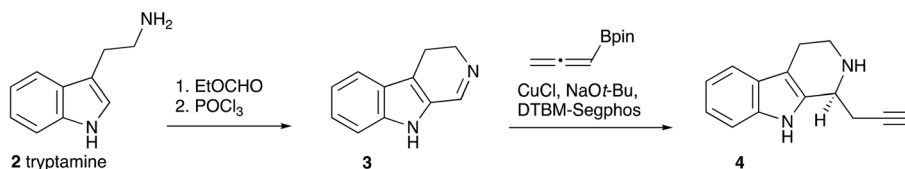


Scheme 1 Dihydrogambirtannine retrosynthesis.

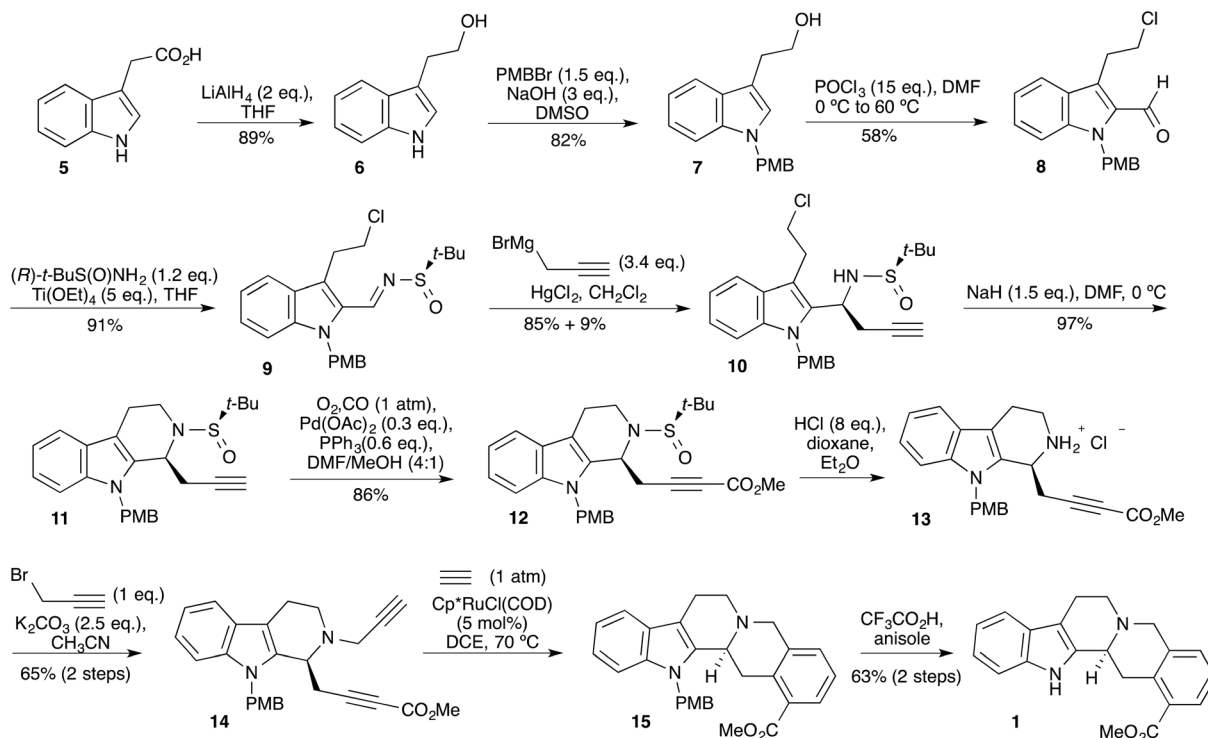
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† Electronic supplementary information (ESI) available: ¹H NMR spectra for compounds **6**–**8**; ¹H and ¹³C NMR spectra for compounds **1** and **9**–**15**. See DOI: <https://doi.org/10.1039/d5ob00162e>





Scheme 2 Asymmetric propargylation.



Scheme 3 Synthesis of (-)-dihydrogambirtannine.

this outcome. Use of either THF or toluene as the solvent resulted in the other diastereoisomer being the major product, but with ratios of just 1:2 and 1:4, respectively. The major diastereoisomer **10** was then treated with base to close the C-ring with formation of tricycle **11**. Attempts to then introduce the methoxycarbonyl group using a combination of a strong base, *n*-butyllithium or LiHMDS, and methyl chloroformate or dimethyl carbonate were fruitless, yielding mixtures. However, palladium catalyzed methoxycarbonylation¹⁹ under carefully controlled conditions delivered the desired ester **12**. The *t*-butylsulfinyl group was then removed under acidic conditions. The deprotected compound was isolated as its hydrochloride salt **13**. Cleavage of the *t*-butylsulfinyl group under these conditions gives *t*-butylsulfinyl chloride as a by-product.²⁰ It is essential that this material is fully removed otherwise sulfinamide **12** can subsequently re-form. The second alkyne moiety was introduced by *N*-alkylation with propargyl bromide to provide diyne **14** for the key [2 + 2 + 2]-cyclo-trimerisation reaction, which would simultaneously create the D and E rings. While use of Wilkinson's catalyst in either

ethanol or toluene resulted in little or no product formation, it was found that this reaction proceeded well using Cp**Ru*(COD)Cl as the catalyst^{21,22} in 1,2-dichloroethane with slow addition of the substrate by syringe pump.¹¹ Pentacycle **15** was formed cleanly and used directly in the next step. The synthesis was completed by removal of the methoxybenzyl group under acidic conditions in the presence of anisole as a carbocation scavenger to give (-)-dihydrogambirtannine **1**.²³ The spectroscopic data were in good agreement with that reported³ for the natural product.

Conclusion

The first asymmetric synthesis of (-)-dihydrogambirtannine has been completed in 11 steps from acid **5** and in 11% overall yield, demonstrating again the power of the [2 + 2 + 2]-cyclo-trimerisation of alkynes in the synthesis of densely substituted benzene rings.



Experimental

General experimental

All reagents were obtained from commercial suppliers and used without further purification unless otherwise indicated. Reactions requiring anhydrous or air free conditions were performed under an atmosphere of nitrogen. Glassware was oven dried at 120 °C and cooled under vacuum. Anhydrous THF and ether were distilled from sodium metal and benzophenone under nitrogen. Anhydrous toluene was distilled from sodium and anhydrous dichloromethane was distilled from calcium hydride. Anhydrous methanol was distilled from magnesium. ¹H and ¹³C NMR spectra were recorded on a Bruker Avance III 400 spectrometer, a 500 MHz ECZ Luminous (JNM-ECZL series) or a JEOL ECA400 UltraShield in CDCl₃. Chemical shifts are given in parts per million (ppm) with residual protic solvent as the internal standard. Coupling constants are given in Hertz. High resolution mass spectra were recorded on a Waters Xevo G2-X2 MS in either positive or negative mode equipped with Waters Acquity UPLC. Analytical thin layer chromatography was performed on Merck DC pre coated TLC plates with 0.25 mm Kieselgel 60 F254. The plates were visualised with a 254 nm UV lamp, or by staining with ammonium molybdate or potassium permanganate. Flash chromatography was performed on silica gel 230–400 mesh.

2-(1*H*-Indol-3-yl)ethan-1-ol (6)

Indole-3-acetic acid (1.75 g, 9.98 mmol) in anhydrous THF (15 mL) was added dropwise to a stirred suspension of lithium aluminium hydride (0.76 g, 20 mmol) in anhydrous THF (30 mL) at 0 °C. The mixture was stirred at 0 °C for 1 h, allowed to warm to room temperature and stirred for 18 h. The mixture was cooled to 0 °C and quenched with saturated Na₂SO₄ solution (20 mL). A white precipitate formed. The mixture was diluted with EtOAc (50 mL) and filtered through Celite, washing with EtOAc. The filtrate was dried (Na₂SO₄) and concentrated under reduced pressure to give alcohol (6) as a light brown solid (1.43 g, 89% yield) which was used directly without further purification. ¹H NMR (400 MHz, CDCl₃) δ 8.03 (brs, 1H), 7.63–7.61 (d, *J* = 7.6 Hz, 1H), 7.38–7.36 (d, *J* = 8.4 Hz, 1H), 7.25–7.08 (m, 2H), 7.04 (d, *J* = 2.4 Hz, 1H), 3.91 (t, *J* = 6.4 Hz, 2H), 3.05 (t, *J* = 6.4 Hz, 2H), 1.55 (brs, 1H). The spectroscopic data were in good agreement with literature values.²⁴

2-(1-(4-Methoxybenzyl)-1*H*-indol-3-yl)ethan-1-ol (7)

1-(Bromomethyl)-4-methoxybenzene (5.2 mL, 37.5 mmol) was added to a stirred mixture of sodium hydroxide (3 g, 75 mmol) and 2-(1*H*-indol-3-yl)ethan-1-ol (4 g, 25 mmol) in DMSO (60 mL) at 0 °C. The mixture was warmed to room temperature and stirred for 3 h. The mixture was diluted with EtOAc (100 mL), extracted with H₂O (3 × 200 mL) and the organic layer was washed with brine (100 mL), dried (Na₂SO₄) and concentrated under reduced pressure. The residue was purified by flash chromatography (silica gel, 20% EtOAc/Hexanes) to give alcohol (7) as a yellow oil (5.75 g, 82% yield).

¹H NMR (400 MHz, CDCl₃) δ 7.64 (d, *J* = 6.4 Hz, 1H), 7.31 (d, *J* = 6.4 Hz, 1H), 7.20–7.10 (m, 2H), 7.08 (d, *J* = 6.8 Hz, 2H), 6.99 (s, 1H), 6.85 (d, *J* = 6.8 Hz, 2H), 5.21 (s, 2H), 3.90 (t, *J* = 5.2 Hz, 2H), 3.78 (s, 3H), 3.03 (t, *J* = 5.2 Hz, 2H), 1.57 (brs, 1H). The spectroscopic data were in good agreement with literature values.²⁴

3-(2-Chloroethyl)-1-(4-methoxybenzyl)-1*H*-indole-2-carbaldehyde (8)

Phosphoryl chloride (21 mL, 0.22 mol) was added dropwise to a stirred solution of 2-(1-(4-methoxybenzyl)-1*H*-indol-3-yl)ethan-1-ol (7) (4 g, 14.2 mmol) in DMF (28 mL) at 0 °C. The mixture was heated at 60 °C for 12 h. The mixture was poured into an ice water (400 mL) and the pH of the solution was slowly adjusted to pH 10 by addition of aq. NaOH solution (2 M, 150 mL). The solution was warmed to room temperature, and stirred for 1 h. The mixture was extracted with EtOAc (3 × 50 mL) and the combined organic layers were washed with brine (50 mL), dried (Na₂SO₄) and concentrated under reduced pressure. The residue was purified by flash chromatography (silica gel, 15% EtOAc/Hexanes) to give aldehyde (8) as an off-white solid (2.69 g, 58% yield). ¹H NMR (400 MHz, CDCl₃) δ 10.15 (s, 1H), 7.74 (d, *J* = 8.4, 1H), 7.40–7.39 (m, 2H), 7.21–7.17 (m, 1H), 7.04 (d, *J* = 8.8 Hz, 2H), 6.78 (d, *J* = 8.4 Hz, 2H), 5.74 (s, 2H), 3.81 (t, *J* = 6.8 Hz, 2H), 3.74 (s, 3H), 3.57 (t, *J* = 7.2 Hz, 2H). The spectroscopic data were in good agreement with literature values.²⁵

(*R,E*)-*N*-((3-(2-Chloroethyl)-1-(4-methoxybenzyl)-1*H*-indol-2-yl)methylene)-2-methylpropane-2-sulfinamide (9)

Ti(OEt)₄ (1.3 mL, 5.65 mmol) was added to a solution of aldehyde (8) (0.37 g, 1.13 mmol) and (*R*)-*t*-butanesulfinamide (164 mg, 1.36 mmol) in anhydrous THF (20 mL). The mixture was heated at reflux for 24 h. The mixture was cooled to room temperature and poured into water (50 mL). The resulting slurry was stirred for 30 min and filtered through Celite. The organic layer was washed with brine (30 mL), dried (Na₂SO₄) and concentrated under reduced pressure. The residue was triturated with hexane. The desired product was isolated by filtration, washed with 20% EtOAc/Hexanes (2 × 10 mL) to give sulfinyl imine (9) as an off-white solid (441 mg, 91% yield); Mp 104–105 °C.

¹H NMR (400 MHz, CDCl₃) δ 8.79 (s, 1H), 7.72 (d, *J* = 6.8 Hz, 1H), 7.33–7.32 (m, 2H), 7.18–7.15 (m, 1H), 6.88–6.86 (m, 2H), 6.77–6.75 (m, 2H), 5.93 (d, *J* = 13.2 Hz, 1H), 5.70 (d, *J* = 13.2 Hz, 1H), 3.79–3.76 (m, 2H), 3.73 (s, 3H), 3.58–3.53 (m, 2H), 1.10 (s, 9H).

¹³C NMR (100 MHz, CDCl₃) δ 158.7, 151.7, 139.3, 130.1, 129.7, 127.0, 126.8, 126.2, 122.1, 120.7, 120.3, 114.0, 110.5, 57.5, 55.2, 47.3, 44.4, 27.9, 22.2.

$[\alpha]_D^{22} = -67.02$ (*c* = 0.57, CHCl₃).

FTIR (AT-IR) ν_{\max} 2962, 1581, 1510, 1243, 1077, 877, 751 cm⁻¹.

HRMS (ESI-Quadrupole) *m/z* calc'd for [C₂₃H₂₇N₂O₂S³⁵Cl + H]⁺ 431.1560, found 431.1562.



(R)-N-((S)-1-(3-(2-Chloroethyl)-1-(4-methoxybenzyl)-1H-indol-2-yl)but-3-yn-1-yl)-2-methylpropane-2-sulfonamide (10)

A 1 M solution of propargyl magnesium bromide was prepared using magnesium turnings (1.2 g, 49.4 mmol), 2 iodine balls and mercury chloride (120 mg, 0.89 mol%) in diethyl ether (25 mL). Propargyl bromide (80% w/w in toluene, 3.6 mL, 32.1 mmol) was added slowly. The mixture was stirred and heated to 35 °C for 2 h. A solution of propargyl magnesium bromide solution in Et₂O (1 M, 8 mL, 8 mmol) was added dropwise to a solution of sulfinyl imine (**9**) (1.01 g, 2.35 mmol) in dichloromethane (25 mL) at -43 °C (ref. 26) over 15–20 min. The mixture was stirred at -43 °C for 16 h. The mixture was quenched with saturated aq. NH₄Cl solution (10 mL) at -43 °C, warmed to room temperature and extracted with EtOAc (3 × 20 mL), and the combined organic layers were washed with brine (20 mL), dried (Na₂SO₄) and concentrated under reduced pressure. The residue was purified by flash chromatography (silica gel, 30% EtOAc/Hexanes) to give the sulfonamide (**10**) diastereomer A as an off-white amorphous solid (0.93 g, 85% yield) and its diastereomer as an off-white amorphous solid (0.16 g, 9% yield).

Sulfonamide (**10**) (more polar, major diastereoisomer): ¹H NMR (400 MHz, CDCl₃) δ 7.64–7.62 (m, 1H), 7.19–7.11 (m, 3H), 6.88 (d, *J* = 8.8 Hz, 2H), 6.82 (d, *J* = 9.2 Hz, 2H), 5.70 (d, *J* = 17.2 Hz, 1H), 5.39 (d, *J* = 17.2 Hz, 1H), 5.01–4.99 (m, 1H), 4.17 (d, *J* = 2.4 Hz, 1H), 3.94–3.87 (m, 1H), 3.81–3.75 (m, 4H), 3.46–3.33 (m, 2H), 2.78 (ddd, *J* = 17.2, 10.0, 2.8 Hz, 1H), 2.60 (ddd, *J* = 16.8, 5.2, 2.8 Hz, 1H), 2.13 (t, *J* = 2.8 Hz, 1H), 1.21 (s, 9H).

¹³C NMR (100 MHz, CDCl₃) δ 159.6, 138.3, 132.9, 130.2, 128.0, 127.7, 123.6, 120.6, 119.6, 115.1, 114.8, 113.5, 111.0, 80.4, 73.2, 56.4, 56.0, 49.7, 47.8, 45.2, 29.2, 26.7, 23.4, 23.2.

$[\alpha]_D^{22} = -80.62$ (*c* = 0.61, CHCl₃).

FTIR (AT-IR) ν_{\max} 3292, 2959, 2302, 1613, 1512, 1466, 1245, 1059, 734 cm⁻¹.

HRMS (ESI-Quadrupole) *m/z* calc'd for [C₂₆H₃₁N₂O₂S³⁵Cl + H]⁺ 471.1873, found 471.1870.

Minor Diastereomer (less polar): ¹H NMR (400 MHz, CDCl₃) δ 7.62–7.60 (m, 1H), 7.20–7.12 (m, 3H), 6.86–6.77 (m, 4H), 5.54 (d, *J* = 17.2 Hz, 1H), 5.45 (d, *J* = 17.2 Hz, 1H), 5.04–5.00 (m, 1H), 3.83 (td, *J* = 8.4, 1.2 Hz, 2H), 3.74 (s, 3H), 3.62 (d, *J* = 2.0 Hz, 1H), 3.46–3.32 (m, 2H), 2.93 (ddd, *J* = 17.2, 10.0, 2.8 Hz, 1H), 2.78 (ddd, *J* = 16.8, 5.2, 2.8 Hz, 1H), 1.91 (t, *J* = 2.8 Hz, 1H), 1.11 (s, 9H).

¹³C NMR (100 MHz, CDCl₃) δ 159.3, 137.7, 134.5, 129.9, 127.6, 127.2, 123.3, 120.4, 119.3, 114.8, 111.9, 110.6, 80.2, 77.7, 71.9, 56.4, 55.7, 50.2, 47.5, 44.8, 29.0, 25.5, 22.8

$[\alpha]_D^{22} = -57.26$ (*c* = 1.14, CHCl₃).

FTIR (AT-IR) ν_{\max} 3290, 2958, 2355, 1612, 1512, 1464, 1245, 1175, 1059, 1034, 736 cm⁻¹.

HRMS (ESI-Quadrupole) *m/z* calc'd for [C₂₆H₃₁N₂O₂S³⁵Cl + H]⁺ 471.1873, found 471.1867.

(S)-2-((R)-tert-Butylsulfinyl)-9-(4-methoxybenzyl)-1-(prop-2-yn-1-yl)-2,3,4,9-tetrahydro-1H-pyrido[3,4-b]indole (11)

Sulfonamide (**10**) (268 mg, 0.56 mmol) in anhydrous DMF (2 mL) was added dropwise to a stirred suspension of sodium

hydride (36 mg of 57–63% w/w in mineral oil, 0.85 mmol) in anhydrous DMF (3 mL) at 0 °C. The mixture was stirred for 45 min and then quenched with saturated aq. NH₄Cl solution (10 mL). The mixture was diluted with EtOAc (20 mL) and extracted with water (3 × 10 mL). The combined organic layers were washed with brine (10 mL), dried (Na₂SO₄) and concentrated under reduced pressure. The residue was purified by flash chromatography (silica gel, 30% EtOAc/Hexanes) to give tricycle (**11**) as a colourless oil (252 mg, 97% yield).

¹H NMR (400 MHz, CDCl₃) δ 7.52 (m, 1H), 7.17–7.08 (m, 3H), 6.92–6.76 (m, 4H), 5.26 (s, 2H), 4.67–4.64 (m, 1H), 3.74 (s, 3H), 3.71–3.69 (m, 1H), 3.50–3.42 (m, 1H), 3.09–2.99 (m, 1H), 2.88–2.78 (m, 3H), 2.65–2.59 (m, 2H), 1.12 (t, *J* = 2.8 Hz, 1H), 1.15 (s, 9H).

¹³C NMR (100 MHz, CDCl₃) δ 158.9, 137.1, 134.8, 129.2, 127.2, 126.9, 122.0, 119.5, 118.3, 114.2, 109.9, 108.6, 80.7, 72.2, 58.5, 55.2, 46.7, 24.6, 22.7, 21.0.

$[\alpha]_D^{22} = +90.40$ (*c* = 1.00, CHCl₃).

FTIR (AT-IR) ν_{\max} 3298, 2928, 2300, 2117, 1612, 1512, 1462, 1245, 1075, 1032, 734 cm⁻¹.

HRMS (ESI-Quadrupole) *m/z* calc'd for [C₂₆H₃₀N₂O₂S + H]⁺ 435.2106, found 435.2105.

Methyl 4-((S)-2-((R)-tert-butylsulfinyl)-9-(4-methoxybenzyl)-2,3,4,9-tetrahydro-1H-pyrido[3,4-b]indol-1-yl)but-2-ynoate (12)

A solution of alkyne (**11**) (0.16 g, 0.36 mmol) in anhydrous MeOH (1.5 mL) was added to a stirred suspension of Pd(OAc)₂ (25 mg, 0.11 mmol) and triphenylphosphine (58 mg, 0.22 mmol) in anhydrous DMF (3 mL) at room temperature. The mixture was then placed under a mixture of CO and O₂ (~1 : 1, 1 atm) and stirred at room temperature for 16 h. Water (10 mL) was added and the mixture was filtered through a plug of Celite. The filtrate was diluted with EtOAc (25 mL) and water (15 mL). The layers were separated and the organic layer was washed with water (2 × 10 mL). The organic layer was washed with brine (15 mL), dried (Na₂SO₄), filtered, and concentrated under reduced pressure. The residue was purified by flash chromatography (silica gel, 40% EtOAc/Hexanes) to give ester (**12**) as a brown gum (157 mg, 86% yield).

¹H NMR (400 MHz, CDCl₃) δ 7.52 (d, *J* = 6.8 Hz, 1H), 7.18–7.07 (m, 3H), 6.93–6.77 (m, 4H), 5.29 (d, *J* = 17.2 Hz, 1H), 5.23 (d, *J* = 17.2 Hz, 1H), 4.74 (dd, *J* = 8.0, 4.4 Hz, 1H), 3.74 (s, 6H), 3.67 (dd, *J* = 14.8, 6.4 Hz, 1H), 3.51 (td, *J* = 11.6, 4.4 Hz, 1H), 3.11–3.03 (m, 1H), 2.95–2.72 (m, 3H), 1.17 (s, 9H).

¹³C NMR (100 MHz, CDCl₃) δ 159.0, 153.7, 137.1, 134.2, 129.1, 127.2, 126.8, 122.2, 119.5, 118.4, 114.3, 109.9, 108.8, 85.6, 77.2, 75.7, 58.6, 55.2, 52.6, 46.7, 24.9, 22.5, 20.6.

$[\alpha]_D^{22} = +10.74$ (*c* = 1.59, CHCl₃).

FTIR (AT-IR) ν_{\max} 2952, 2237, 1711, 1512, 1246, 1074, 736 cm⁻¹.

HRMS (ESI-Quadrupole) *m/z* calc'd for [C₂₈H₃₂N₂O₄S + H]⁺ 493.2161, found 493.2166.

Methyl (S)-4-(9-(4-methoxybenzyl)-2,3,4,9-tetrahydro-1H-pyrido[3,4-b]indol-1-yl)but-2-ynoate hydrochloride salt (13)

HCl in dioxane (4 M, 0.2 mL, 0.8 mmol) was added dropwise to a solution of ester (**12**) (49.7 mg, 0.1 mmol) in anhydrous



Et₂O (3 mL) at 0 °C. The mixture was stirred for 45 min, concentrated to dryness, and triturated with a mixture of hexane and Et₂O (1 : 1, 20 mL). The off-white hydrochloride salt was isolated by filtration, washing with Et₂O (2 × 10 mL) to give hydrochloride salt (**13**) as an off-white powder (42 mg) which was used directly in the next step without further purification.

¹H NMR (400 MHz, CDCl₃) δ 10.32 (brs, 2H), 7.54 (d, *J* = 7.2 Hz, 1H), 7.26–7.10 (m, 3H), 6.87–6.77 (m, 4H), 5.24 (d, *J* = 17.2 Hz, 1H), 5.14 (d, *J* = 17.2 Hz, 1H), 5.03 (m, 1H), 3.81–3.75 (m, 1H), 3.73 (s, 3H), 3.66 (s, 3H), 3.57–3.54 (m, 1H), 3.30–3.22 (m, 2H), 3.04–3.00 (m, 1H), 2.83–2.77 (m, 1H).

¹³C NMR (100 MHz, CDCl₃) δ 159.1, 153.2, 137.7, 128.3, 127.3, 126.9, 125.9, 123.3, 120.1, 118.9, 114.4, 110.1, 108.7, 82.6, 77.2, 55.2, 52.7, 48.2, 46.7, 38.6, 23.4, 18.3.

FTIR (AT-IR) ν_{max} 2951, 2447, 2240, 1713, 1512, 1248, 1176, 741 cm⁻¹.

HRMS (ESI-Quadrupole) *m/z* calc'd for [C₂₄H₂₄N₂O₃ + H]⁺ 389.1865, found 389.1868.

Methyl (S)-4-(9-(4-methoxybenzyl)-2-(prop-2-yn-1-yl)-2,3,4,9-tetrahydro-1H-pyrido[3,4-*b*]indol-1-yl)but-2-ynoate (**14**)

Propargyl bromide (80% w/w in toluene, 9.6 μL, 0.1 mmol) was added dropwise to a solution of hydrochloride salt (**13**) (42 mg, 0.099 mmol) and potassium carbonate (35 mg, 0.25 mmol) in anhydrous CH₃CN (3 mL) at 0 °C. The mixture was stirred at 0 °C for 2 h, allowed to warm to room temperature and stirred for 18 h. The mixture was diluted with EtOAc (20 mL) and water (10 mL). The layers were separated and the organic layer was washed with water (2 × 10 mL) and brine (15 mL), dried (Na₂SO₄), filtered, and concentrated under reduced pressure. The residue was purified by flash chromatography (silica gel, 30% EtOAc/Hexanes) to give the diyne (**14**) as a pale-yellow gum (34.5 mg, 65% yield over 2 steps).

¹H NMR (400 MHz, CDCl₃) δ 7.53 (d, *J* = 7.2 Hz, 1H), 7.21–7.08 (m, 3H), 6.89–6.76 (m, 4H), 5.26 (s, 2H), 4.25 (dd, *J* = 8.4, 4.4 Hz, 1H), 3.75 (s, 3H), 3.74 (s, 3H), 3.51 (dd, *J* = 16.4, 2.4 Hz, 1H), 3.40 (dd, *J* = 16.8, 2.0 Hz, 1H), 3.27–3.24 (m, 2H), 2.96–2.88 (m, 1H), 2.79–2.65 (m, 3H), 2.19 (t, *J* = 2.4 Hz, 1H).

¹³C NMR (100 MHz, CDCl₃) δ 158.9, 154.0, 137.3, 133.3, 127.3, 127.1, 126.7, 122.0, 119.4, 118.4, 114.3, 114.2, 109.8, 108.4, 87.0, 80.1, 77.2, 74.3, 72.7, 55.2, 54.2, 52.5, 46.5, 43.5, 42.7, 25.1, 17.6.

FTIR (AT-IR) ν_{max} 3282, 2924, 2302, 2239, 1713, 1513, 1275, 1260, 750 cm⁻¹.

HRMS (ESI-Quadrupole) *m/z* calc'd for [C₂₇H₂₆N₂O₃ + H]⁺ 427.2022, found 427.2025.

Methyl (S)-13-(4-methoxybenzyl)-5,7,8,13,13b,14-hexahydroindolo[2',3':3,4]pyrido[1,2-*b*]isoquinoline-1-carboxylate (**15**)

A solution of Cp*RuCl(COD) (1.9 mg, 5.04 × 10⁻³ mmol, 5 mol%) in anhydrous 1,2-dichloroethane (2 mL) was purged with acetylene gas for 10 min. A solution of diyne (**14**) (43 mg, 0.1 mmol) in anhydrous 1,2-dichloroethane (2 mL) was simi-

larly purged with acetylene gas for 10 min and then slowly added to the solution of Cp*RuCl(COD) at 70 °C over 4 h using a syringe pump. The mixture was cooled to room temperature and the mixture was filtered through a plug of Celite. The filtrate was concentrated under reduced pressure to give pentacycle (**15**) (48 mg) which was used directly in the next step without further purification.

¹H NMR (400 MHz, CDCl₃) δ 7.75–7.72 (m, 1H), 7.56–7.54 (m, 1H), 7.24–7.11 (m, 5H), 6.97–6.94 (m, 2H), 6.81–3.79 (m, 2H), 5.39 (d, *J* = 17.2 Hz, 1H), 5.27 (d, *J* = 17.2 Hz, 1H), 4.11–3.99 (m, 2H), 3.77–3.61 (m, 2H), 3.75 (s, 3H), 3.65 (s, 3H), 3.26–3.01 (m, 3H), 2.92–2.88 (m, 1H), 2.79–3.73 (m, 1H).

¹³C NMR (100 MHz, CDCl₃) δ 167.8, 158.8, 137.9, 136.4, 135.5, 135.2, 130.4, 129.8, 129.6, 128.9, 127.2, 126.9, 125.6, 121.5, 119.3, 118.2, 114.1, 109.8, 109.2, 58.0, 55.3, 55.2, 51.7, 50.2, 47.3, 33.3, 22.1, 21.

FTIR (AT-IR) ν_{max} 2952, 2929, 1717, 1513, 1249, 749 cm⁻¹.

HRMS (ESI-Quadrupole) *m/z* calc'd for [C₂₉H₂₈N₂O₃ + Na]⁺ 475.1998, found 475.2000.

(-)-Dihydrogambirtannine (**1**)

Trifluoroacetic acid (2.5 mL) was added to a solution of the crude pentacycle (**15**) (48 mg) in anisole (0.5 mL) at 0 °C under N₂ atmosphere. The mixture was warmed to room temperature and stirred for 3 days. The mixture was concentrated under reduced pressure and the residue was diluted with EtOAc (10 mL) and quenched with saturated aq. NaHCO₃ to pH ~10. The aqueous layer was extracted with EtOAc (2 × 10 mL). The combined organic layers were washed with brine (10 mL), dried (Na₂SO₄) and concentrated under reduced pressure. The residue was purified by flash chromatography (basic alumina (200–300 mesh), 7 : 2 : 1 Hexanes/CH₂Cl₂/EtOAc) to give dihydrogambirtannine (**1**) as a pale yellow gum (21 mg, 63% yield over 2 steps).

¹H NMR (400 MHz, CDCl₃) δ 8.01 (brs, 1H), 7.85 (d, *J* = 7.6 Hz, 1H), 7.51 (d, *J* = 7.6 Hz, 1H), 7.35 (d, *J* = 8.0 Hz, 1H), 7.29–7.21 (m, 4H), 4.14 (d, *J* = 15.2 Hz, 1H), 3.90 (s, 3H), 3.89–3.65 (m, 3H), 3.31 (dd, *J* = 11.2, 4.8 Hz, 1H), 3.19 (dd, *J* = 17.2, 11.6 Hz, 1H), 3.08–3.03 (m, 1H), 2.86–2.81 (m, 1H), 2.73 (td, *J* = 11.6, 4.4 Hz, 1H).

¹³C NMR (100 MHz, CDCl₃) δ 167.6, 136.3, 136.0, 135.6, 134.3, 130.9, 129.2, 128.9, 127.1, 125.7, 121.6, 119.4, 118.1, 110.9, 108.5, 58.1, 56.3, 52.4, 51.9, 33.5, 21.4.

[α]_D²⁴ = -145 (*c* = 0.33, CHCl₃).

FTIR (AT-IR) ν_{max} 3391, 2948, 1716, 1457, 1289, 1268, 1141, 744 cm⁻¹.

HRMS (ESI-Quadrupole) *m/z* calc'd for [C₂₁H₂₀N₂O₂ + H]⁺ 333.1603, found 333.1600.

Data availability

The data supporting this article have been included as part of the ESI.†



Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We thank the National Research Foundation, Singapore, NRF Competitive Research Programme (CRP), grant numbers NRF-CRP27-2021-0002 and NRF-CRP18-2017-01, for financial support. We thank NTU for support. V. P. thanks the URECA programme of NTU.

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