Stereoselective Domino Rearrangement peri-Annulation of Cinchona Alkaloid Derivatives with 8-Bromo-1-naphthyl Grignard

Przemysław J. Boratyński*

ABSTRACT: The unexpected domino coupling and rearrangement of the Cinchona alkaloid skeleton has been found to occur in the reaction of 9-chloro-9-deoxy-alkaloids with Grignards from peri-dihalogenonaphthalene. The cyclization and migration of the central quinuclidinylmethyl group (C9) from position C-4 to position C-3' of quinoline formed a novel chiral ring system of 5-aza-7H-benzo[no]tetraphene, yielding products of unlike configuration. The proposed reaction pathway involves radical intermediates.

INTRODUCTION

The Cinchona alkaloids are of relevance to medicinal chemistry and the development of asymmetric catalytic methods. Many valuable organocatalysts and metal ligands were made by the manipulation of the central 9-hydroxyl group of quinine. In 2008, a synthetically viable substitution of 9-chlorodeoxy-Cinchona alkaloids with sp2 Grignard reagents providing 9-arylated Cinchona derivatives emerged. The reactions proceeded stereoconvergently, producing only an 8,9-like diastereoisomer from both 9R and 9S epimers of chloro derivatives. This stereochemical outcome was justified by the coordination of magnesium by the quinuclidine nitrogen atom. In previous reactions of Grignard reagents from meta- and para-dihalobenzene, halophenyl derivatives were rather efficiently prepared (23–80% yield), while di-Grignard reagents produced the corresponding dimeric products. Reactions of unmodified alkaloids with Grignard reagents have been shown to result in nucleophilic additions at positions 4' and 2' (Scheme 1A).

Cinchona alkaloids functionalized at position 9 with the 1,2-disubstituted naphthyl group have been utilized in several asymmetric transformations. The naphthalene ring facilitates π-interactions and gives versatility to functional group placement. Although the most effective spatial control would be enforced through 1,8-substitution (peri arrangement), no such product has been described, hence the attempt to couple a Cinchona alkaloid with the 8-functionalized naphthalene ring amenable for further derivatization. The reactivity of 1,8-dihalomnapthalene-derived Grignards with 9-chloro-9-deoxy-quinine resulted in the formation of an additional carbon–carbon bond and an unprecedented rearrangement within the quinoline part. Previously, some modifications at position 9 caused rearrangements of the alkaloid structure, mostly by the breakage of the adjacent N-1–C-8 bond, which resulted in quinuclidine ring opening or ring expansion. There is a reaction of a likely radical mechanism, in which the quinuclidine methyl group separates from the quinoline ring on treatment with the LiAlH4/O2 system (Scheme 1B). However, no migration of this group to other positions of the quinoline ring has been reported. Since the migration of any group from position 4 to position 3 of pyridine requires special circumstances, it is rare.

RESULTS AND DISCUSSION

In order to selectively modify Cinchona alkaloids with the previously developed method for C-9–C bond formation, 9S-chlorodeoxyquinine was treated with the Grignard reagents obtained separately from 1,8-diodonaphthalene and 1,8-dibromonaphthalene. The electrospray mass spectrometry of the crude reaction mixtures only revealed traces of naphthalene 8-halogenated derivatives. Instead, the most abundant signal originated from unexpected product 1, which was isolated in up to 30% yield (Scheme 1C). In this product, the quinuclidine methylene unit (the C-9 atom) migrated from position 4' to position 3' of the quinoline ring and the newly introduced naphthalene ring became fused between the central C-9 and quinoline C-4' carbon atoms. The product contains an unprecedented fusion of five rings with one nitrogen and one sp3 carbon atom of defined stereochemistry.

By way of model experiments on the reaction mechanism, the reaction of 1,8-dibromonaphthalene with magnesium in tetrahydrofuran (THF) was initially found to provide a mixture of mono- and bis-Grignard reagents (3:1 to 10:1) after 1 h of reaction time and only moderately correlated with the ratio of...
Scheme 1. (A) Reactivity of Cinchona Alkaloids and Their Derivatives with Grignard Reagents; (B) Rearrangements and Bond Dissociations in Cinchona Alkaloid Chemistry Involving the C-9 Atom; (C) New Domino Coupling—Rearrangement; Essential NOESY Interactions and Traditional Atom Numbering Are Shown

| Table 1. Domino Coupling—Rearrangement of 9-Chloro-9-deoxy Cinchona Alkaloids |
|---------------------------------|-----|-----------------|-----------------|
| parent alkaloid                | R²  | R³              | product config  | product, %²   |
| quinine                        | OMe | C₇H₈            | (8S,9R)         | 16–30        |
| quinidine                      | OMe | C₇H₈            | (8R,9S)         | 20–15        |
| cinchonine                     | H   | C₇H₈            | (8R,9S)         | 3–5          |
| dhydrodrcinchonidine           | H   | Et              | (8S,9R)         | 2–4          |

²Isolated yield. bUnder optimized conditions with 2 equiv of 8-bromo-1-naphthyl magnesium bromide.

reactants. When the reaction was carried out for 18 h, bis-Grignard¹⁰ and mono-Grignard were both separately prepared in an estimated >92% selectivity by controlling the magnesium to dibromonaphthalene ratio (2.1:1 and 1.07:1) as evidenced by quenching experiments (for details, see the SI). In the subsequent reaction with 9-chloro-9-deoxyquinine, an increase in bis-Grignard quantity led to a significant deterioration of yields. An opposite effect was seen with pure 8-bromo-1-naphthylmagnesium bromide.

First, the substitution of quinine 9-halide with 8-bromonaphthalene occurred thus precluding thermodynamic equilibration of the product. The reaction of any 9 epimer of 9-chlorodeoxyquinine resulted in the formation of the same isomer of product 1. This is partly consistent with our previous finding that the Wurtz-type coupling of Grignard reagents only produced a single like stereoisomer of the product regardless of the configuration at position 9 of the starting material.² In later experiments it was shown that thermodynamic base-promoted equilibration produced a mixture of stereoisomers in comparable quantities.³

Here, for the reaction quenched in D₂O no observable incorporation of deuterium into the molecule occurred thus precluding thermodynamic equilibration of the product.

A tentative reaction mechanism can be outlined (Scheme 2). First, the substitution of quinine 9-halide with 8-bromonaphthalene-magnesium bromide according to the previously described pathway² would produce intermediate Int.A. Proximity of another Grignard molecule could initiate single electron transfer (SET) analogous to the one postulated for metal—halogen exchange in main group organometallic chemistry, particularly at elevated temperatures.¹³ This could produce aryl radical Int.B.¹⁴ This localized nucelophic radical can attack the C-4 atom of the quinoline ring, forming a spirocyclic radical Int.C.¹⁵ The ensuing fragmentation of the C-4/C-9 bond produces Int.D in an overall radical substitution reaction from Int.B.¹⁶ In this intermediate, the radical is of a highly delocalized benzyl
type and as such is expected to be more stable than Int.B*. Furthermore, sufficient lifetime of this species may result in the loss of stereochemistry at the sp\(^2\) carbon at the former position 9. Productive intramolecular addition\(^{16}\) in the intermediate Int.D* can result in the formation of a bond between C-9 and quinoline C-3', giving diarylmethyl-type radical Int.E*, which has a complete carbon skeleton of the end product 1. A similar mechanism was proposed for radical rearrangement annulation involving nitrogen-centered (aminyl) attacking and leaving radicals.\(^{17}\)

Overall, the presence of a radical pathway is partly supported by the observation of a faint EPR signal after 3 h of reaction time and by the trapping experiments with DMPO and TEMPO. The adduct with DMPO showed an intense EPR trace, which could not be easily interpreted. The diamagnetic coupling product with TEMPO was identifiable in the ESI-MS. The observed value (m/z 590) is consistent with the formula of isomeric intermediate radicals Int.B**–F** (for details, see the SI).

The end radical Int.E/Int.F* will eventually become diamagnetic 1, either by hydrogen abstraction and oxidative rearromatization during workup or by electron abstraction in another SET process. The latter explanation may be consistent with the unchanged ESI-MS spectral pattern following the workup of the reaction mixture under reductive conditions (NaBH\(_4\)). For a brief discussion of alternative reaction pathways, see the SI.

Some stabilization of the proposed intermediates Int.C–F* may be offered by forming a coordination bond between the quinoline nitrogen atom and magnesium ions.\(^{12,15}\) DFT calculations on simplified models (MgBr\(^{2-}\) removed or replaced with a proton) were conducted at the DFT/B3LYP/CC-pVDZ and M06-2X/CC-pVDZ levels of theory. These indicate that the radical isomerization pathway from Int.B* to Int.F* is energetically favorable. For the observed (9\(^R\))-1, additional stabilization can be offered by the interaction between quinoline C-2' and quinuclidine nitrogen atoms (Int.Fr*). In the radical cation model, the geometry of Int.Er* converges into Int.Fr*.

This intermolecular nucleophilic addition is spatially not accessible for the unobserved (9\(^S\))-1 and is likely the cause of the observed stereoselectivity in the reaction (Figure 1).

Attempts to extend the scope of the reaction for either other peri-substituted arenes or non-Cinchona alkaloid derivatives were synthetically ineffective. For the reaction of similar 5,6-dibromoacenaphthene, the most abundant signal in the mass spectrometry corresponded to alkaloid 9-dimer. The presence of a likely cyclized product was evident (m/z = 459), but the quantity was low and the isolation of a sample of sufficient purity
The obtained products can be defined as nitrogen-containing polycyclic aromatic systems which may be valued for their electronic and associated fluorescent properties. The large nearly planar polycyclic aromatic system with a nitrogen atom in 1–4 is the cause of fluorescent properties on the TLC plate and in the solution. In the absence of external acid, blue light is emitted, while in 15 mM TFA the solution of 1 becomes deeply orange and green fluorescence emerges with a similar quantum yield (Φ_F = 0.32–0.45, Figure 2A). The corresponding emission maxima for quinine-derived 1 are 436 nm for neutral and 522 nm for acidic samples. The presence of acid also increases the Stokes shift by a factor of 2 (56 nm vs 94 nm). In contrast, the results for the cinchonine derivative 3 (Φ_F = 0.47–0.58) show that the methoxy group is not responsible for fluorescence (for details, see the SI). The structure of the modified natural products as well as acidity-dependent fluorescence prompted the evaluation of its utility for biological staining in a simple assay. The microscopic live plant cell imaging with the quinine derivative 1 revealed preferential fluorescence staining of some globular cell cytoplasm structures surrounding the nucleus and cell walls (Figure 2B).

CONCLUSIONS

In summary, the unprecedented rearrangement involving carbon bond migration from position 4 to position 3 of quinoline without transition metals and under nonic conditions forms a novel chiral 5-aza-7H-benzo[no]tetraphene ring system with fluorescent properties. While the isolated products are limited to Cinchona alkaloid derivatives, the transformation may be relevant to other lepidine and peri-naphthalene derivatives.

EXPERIMENTAL SECTION

General Comments. NMR spectra were collected on a 600 MHz Bruker Avance II instrument. Spectra were internally referenced to tetramethylsilane (TMS, δ_C = 0 and δ_H = 0). Structural assignments were made with additional information from gCOSY, gHSQC, gHMBC, and NOESY experiments. Electrospray (ESI) MS and HRMS spectra were recorded on a Waters LCT Premier XE apparatus with a TOF analyzer. ECD spectra were measured on a Jasco J-1500 circular dichroism spectrophotometer. UV−vis spectra were taken on a Jasco V-670 spectrophotometer. Fluorescence spectra were taken on a Horiba Fluoromax-4 spectrofluorimeter and are uncorrected. Flash chromatography was performed on standard silica gel 230–400 mesh (Merck). Automated flash-chromatography system CombiFlash NextGen 300 (ISCO, Teledyne) was used in some isolations. TLC plates with F256 indicator (Sigma-Aldrich) were illuminated by a dual UV lamp at 254 and 365 nm. Cinchona alkaloids were purchased from Buchler (Braunschweig, Germany). 9-Chloro-9-deoxyquinine and other 9-deoxy-9-halogeno-alkaloids were obtained by the treatment of the corresponding Cinchona alkaloid with thionyl chloride (56–84% yield) as described in the literature. THF was purified and dried by sequential distillation from LiAlH₄ and distillation from sodium/hydroquinone, and toluene was dried by storing over sodium/potassium metal. All other reagents were purchased from commercial suppliers (Merck/Sigma-Aldrich and Fluorochem) and used as received.

Magnesium (190 mg, 7.88 mmol, 1.5 equiv) was activated with iodine (ca. 10 mg) and suspended in dry THF (26 mL) under argon. 1,8-Dibromonaphthalene (2.08 g, 7.26 mmol, 1.4 equiv) was added, and the mixture was stirred under reflux in an oil bath. 1,2-Dibromoethane was added in small portions (total 50 μL, 0.58 mmol, 0.1 equiv), and after ca. 1–1.5 h almost all magnesium dissolved. Then, a solution of 9-chloro-9-deoxyquinine (1.78 g, 5.19 mmol, 1 equiv) in toluene (25 mL) was added to the reaction mixture, and stirring was continued in an oil bath at 85–90 °C for 18 h. The mixture gradually took a deep brown color. The heating was discontinued, and at room temperature the reaction was quenched with a saturated NH₄Cl solution (15 mL), extracted with DCM (1 × 60 mL, 3 × 10 mL), and dried over MgSO₄ in a flask open to air. After 18 h the mixture was filtered, evaporated, and subjected to column chromatography on silica gel with DCM/MeOH (3% to 5% gradient), and fractions containing a bright fluorescent spot on TLC were collected. Obtained 0.449 g of 1 as a light brown, amorphous solid (20%).

Figure 2. (A) Emission (solid lines) and excitation (dashed lines) for a 10⁻³ M solution of 1 in DCM without acid (blue lines) and in 15 mM TFA (red lines). (B) Fluorescence micrograph (280× magnification, FITC setup) of live plant skin from Allium cepa stained with 1.
Repeated reactions on 0.6–3.5 g (2–10 mmol) scale gave 15–20% yields. The reaction performed with 1,8-diodonaphthalene instead of 1,8-dibromonaphthalene gave a 9% yield. An increase of relative quantity of magnesium vs 1,8-dibromonaphthalene or use of >2 equiv of magnesium generally caused a decrease in isolated pure product yield. Reactions run for significantly shorter time (3 h) or at room temperature showed much lower conversion. Similar yields were obtained for the reactions run in benzene/THF instead of toluene/THF. A reaction performed with 2.0 equiv of 8-bromo-1-naphthylmagnesium bromide gave a 30% yield.

**Preparation of 8-Bromo-1-naphthylmagnesium Bromide.** In a sealed Schlenk tube under an argon atmosphere magnesium (55 mg, 2.28 mmol, 1.07 equiv) was placed and activated with bromine (ca. 20 μL) at 150–200 °C for 3 min, bromine was evacuated, and a solution of 1,8-dibromonaphthalene (664 mg, 2.32 mmol) in dry THF was added (10 mL). The container was sealed and heated at 80 °C in an oil bath for 18 h, producing a dark brown but transparent solution, which was used directly for the preparation of 1 (26–30% yield).

1H NMR (600 MHz, CDCl₃, TMS): δ 8.75 (s, 1H), 8.44 (d, J = 7.2 Hz, 1H), 8.05 (d, J = 9.1 Hz, 1H), 8.01 (d, J = 2.6 Hz, 1H), 7.92 (d, J = 8.2 Hz, 1H), 7.76–7.79 (m, 1H), 7.68 (t, J = 7.7 Hz, 1H), 7.47–7.49 (m, 2H), 7.32 (d, J = 9.1, 2.6 Hz, 1H), 5.33 (ddd, J = 17.1, 10.3, 8.1 Hz), 4.72 (dt, J = 17.1, 1.1 Hz, 1H), 4.64 (dt, J = 10.3, 1.5 Hz, 1H), 4.19 (d, J = 10.1 Hz, 1H), 3.92 (s, 3H), 3.37–3.43 (m, 1H), 2.94 (ddd, J = 14.1, 9.9 Hz, 1H), 2.82–2.88 (m, 1H), 2.32 (q, J = 9.5 Hz, 1H), 2.19 (dt, J = 14.1, 2.9 Hz, 1H), 2.03–2.07 (m, 1H), 1.58–1.63 (m, 2H), 1.45–1.52 (m, 1H), 1.07–1.12 (m, 1H), 0.73–0.79 (m, 1H) ppm. 13C (H) NMR (151 MHz, CDCl₃, TMS): δ 158.0, 150.3, 144.4, 142.4, 134.5, 133.38, 133.29, 132.6, 131.7, 130.4, 128.8, 128.6, 126.14, 126.02, 126.00, 125.95, 125.5, 122.5, 119.4, 113.8, 104.9, 62.8, 56.6, 55.5, 47.2, 413, 40.5, 28.7, 28.2, 27.7 ppm. 13C (H) NMR (151 MHz, CDCl₃, TMS): δ 158.1, 150.7, 145.3, 142.1, 133.8, 133.79, 133.37, 132.8, 132.5, 130.6, 129.1, 128.5, 126.1, 125.98, 125.92, 125.87, 125.3, 125.1, 119.3, 113.4, 104.9, 63.0, 56.5, 54.6, 47.0, 410, 40.4, 28.5, 28.3, 27.8 ppm. HRMS (ESI-TOF) m/z: calcd for [C₁₉H₂₈N₂O⁺ + H]⁺ 433.2274, found 433.2275.

(7S)-5-Aza-2-methoxy-7-(15,2R,5R)-5-vinylquinuclidin-2-yl)-7H-benzo[n]tetraphene (2); Cinchonidine Derivative 2.

The product was obtained as described for 1 starting from magnesium (0.198 g, 8.22 mmol, 1.5 equiv), 1,8-dibromonaphthalene (2.21 g, 7.72 mmol, 1.4 equiv), and 9R-chloro-9-deoxyquinicline (1.74 g, 5.49 mmol, 1 mmol) instead of chlorodeoxyquinicline. Following chromatography on silica gel with DCM/MeOH 3% to 5% gradient gave a brown amorphous solid (121 mg, 5%). 1H NMR (600 MHz, CDCl₃, TMS): δ 8.89 (s, 1H), 8.61 (d, J = 8.7 Hz, 1H), 8.39 (d, J = 7.3 Hz, 1H), 8.15 (dd, J = 8.7, 1.1 Hz, 1H), 7.93 (d, J = 8.2 Hz, 1H), 7.79 (d, J = 8.1 Hz, 1H), 7.66 (t, J = 7.9 Hz, 1H), 7.63–7.67 (m, 1H), 7.48–7.55 (m, 3H), 5.87 (ddd, J = 17.3, 10.4, 7.3 Hz, 1H), 5.08 (dt, J = 10.4, 1.4 Hz, 1H), 5.04 (d, J = 17.3 Hz, 1H), 4.33 (ddd, J = 9.6 Hz, 1H), 3.06 (ddd, J = 13.6, 10.1 Hz, 1H), 2.91–2.96 (m, 1H), 2.60–2.67 (m, 1H), 2.42–2.48 (m, 1H), 2.38 (q, J = 9.0 Hz, 1H), 2.20 (q, J = 8.2 Hz, 1H), 1.55–1.58 (m, 1H), 1.36–1.41 (m, 1H), 1.28–1.34 (m, 1H), 1.12–1.19 (m, 1H), 0.47–0.52 (m, 1H) ppm. 13C (H) NMR (151 MHz, CDCl₃, TMS): δ 152.6, 148.3, 140.3, 135.9, 133.4, 133.0, 131.9, 130.41, 130.26, 129.1, 128.3, 128.0, 126.8, 126.4, 126.21, 126.06, 125.7, 125.5, 125.3, 125.0, 115.4, 62.8, 49.4, 47.9, 46.4, 39.9, 28.3, 26.12, 26.03 ppm. HRMS (ESI-TOF) m/z: calcd for [C₂₉H₂₆N₂O⁺ + H]⁺ 436.2175, found 436.2175.

(7R)-5-Aza-2-(15,2S,5S)-5-ethylquinuclidin-2-yl)-7H-benzo[n]tetraphene (4); Dihydrocinchonidine Derivative 4.

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/jacs.2c01249.

![Diagram](https://pubs.acs.org/joc)
Plots of NMR and EPR spectra, assignment of NMR signals for compounds 1 and 3, plots of ESI-MS, UV−vis, CD, and fluorescence spectra, micrographs, and computational details (PDF)
FAIR data, including the primary NMR FID files, for compounds 1−4 (ZIP)

AUTHOR INFORMATION
Corresponding Author
Przemysław J. Boratyński − Department of Organic and Medicinal Chemistry, Wrocław University of Technology, Wrocław 50-370, Poland; orcid.org/0000-0002-7139-8568; Email: przemyslaw.boratynski@pwr.edu.pl

Complete contact information is available at: https://pubs.acs.org/10.1021/acs.joc.2c01249

Notes
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