

Communication

Macrocyclic Spermidine Alkaloids from *Androya decaryi* L. Perrier

Anne-Cécile Le Lamer ^{1,*}, Nehal Ibrahim ¹, Frédéric Manjary ², Sonia Mallet-Ladeira ³, Cynthia Girardi ¹, Alexis Valentin ¹, Nicolas Fabre ¹ and Claude Moulis ¹

¹ Université de Toulouse; UMR 152 UPS-IRD (Pharmacochimie et Pharmacologie pour le Développement-PHARMA DEV), F-31062 Toulouse Cedex 9, France

² Laboratoire de Chimie de la Faculté des Sciences, Université de Toliara, BP 185 Maninday Toliara 602, Madagascar

³ Institut de Chimie de Toulouse, Université de Toulouse, UPS, FR2599, 118 route de Narbonne, F-31062 Toulouse Cedex 9, France

* Author to whom correspondence should be addressed; E-Mail: anne-cecile.le-lamer@univ-tlse3.fr.

Received: 18 February 2013; in revised form: 28 March 2013 / Accepted: 29 March 2013 /

Published: 4 April 2013

Abstract: Three new spermidine alkaloids and two known compounds were isolated from the leaves of *Androya decaryi*. Their structures were elucidated on the basis of their spectroscopic data (NMR and mass spectrometry), by X-Ray diffraction and by comparison with literature values. Evaluation of the *in vitro* antiplasmodial properties of the isolated compounds revealed they did not possess any significant activity.

Keywords: *Androya*; Scrophulariaceae; spermidine alkaloids

1. Introduction

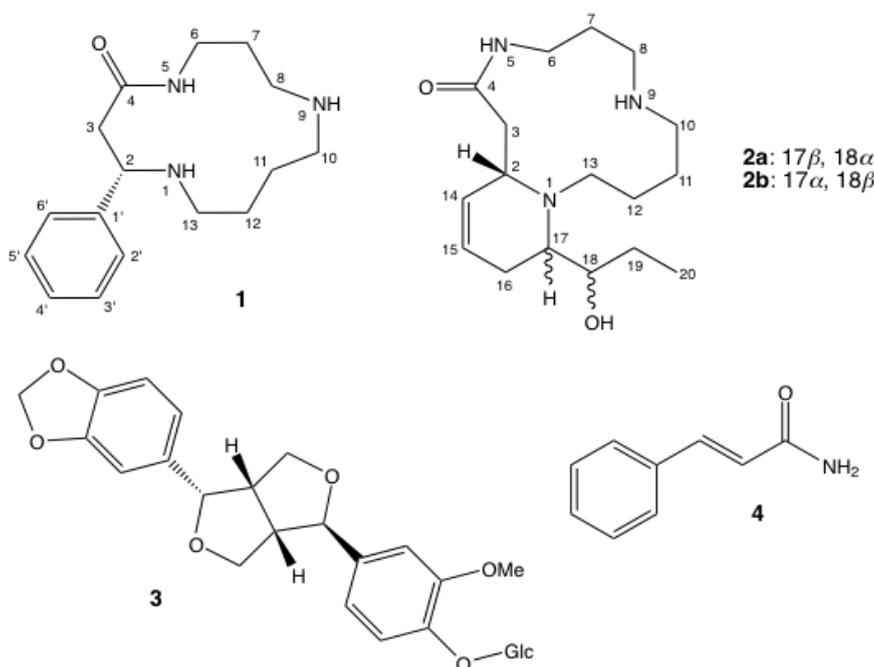
Androya decaryi L. Perrier (Scrophulariaceae) [1] is an evergreen shrub endemic to the southwest of Madagascar. Traditionally used in folk medicine, the decoction of the aerial parts is used to treat yellow fever and malaria-like symptoms (according to local ethnopharmacological investigations). Considering the need for new antiplasmodial compounds due to the emergence of resistant strains of *Plasmodium falciparum* [2], and the fact that natural products and phytomedicines are still promising sources of antimalarial agents [3–5], the aerial parts of *A. decaryi* were phytochemically investigated.

Herein, we report the isolation and structural identification of three new macrocyclic spermidine alkaloids **1-2a/b**, along with a lignan **3** and (*E*)-cinnamamide (**4**) from the leaves *A. decaryi*, as well as the evaluation of their *in vitro* antiplasmodial activity. To the best of our knowledge, this is the first report on phytochemistry of the genus *Androya* and on spermidine alkaloids from a plant belonging to the Scrophulariaceae.

2. Results and Discussion

Since alkaloids have been detected by TLC in the leaves of *A. decaryi*, a leaf alkaloid extract was studied. Compounds **1** and **2a** (Figure 1) were isolated from the crude alkaloid fraction. Minor alkaloids were also detected; a second extraction was therefore performed on a larger amount of *A. decaryi*, leading to the isolation of the alkaloid **2b** and compounds **3** and **4** (Figure 1).

Figure 1. Structures of compounds **1-4**.



Compound **1**, a yellow oil, had the molecular formula $C_{16}H_{25}N_3O$ as established by HR-ESIMS (m/z 276.2078 $[M+H]^+$). The 16 carbon signals (Table 1) were identified with *J*-modulated ^{13}C -NMR spectrum as eight sp^3 methylenes, one sp^3 methine, five sp^2 methines and two sp^2 quaternary carbons, including one amide carbonyl. The 1H -NMR and HSQC spectra displayed signals for three pairs of aliphatic methylene protons between δ_H 1.40 and 1.85, five pairs of deshielded methylene protons between δ_H 2.33 and 3.74, one deshielded methine (δ_H 4.01), and a monosubstituted benzene ring. These data were consistent with a 13-membered cyclic spermidine alkaloid bearing an aromatic ring. Analysis of COSY and HMBC spectra, measurement of a negative optical rotation ($[\alpha]_D^{25} = -13$ (c 3.0, MeOH), and comparison with literature values indicated that compound **1** was (–)-(2*S*)-2-phenyl-1,5,9-triazacyclotridecan-4-one [6]. Although this alkaloid has been used as a building block for the synthesis of other natural spermidine alkaloids [6–8], it is reported here for the first time as a natural product.

Table 1. ^1H and ^{13}C -NMR data for compound **1** in CDCl_3 .

	δ_{H} (J in Hz) ^a	δ_{C}
2	4.01 dd (11.5, 2.9)	60.0
3	2.48 dd (15.0, 2.9); 2.56 dd (15.0, 11.5)	45.0
4	-	171.7
5	8.57 bs	-
6	3.19 dt (13.7, 5.5); 3.70 dt (13.7, 5.2)	39.6
7	1.80–1.85 m	27.8
8	2.86 dt (12.1, 5.0) 2.96 dt (12.1, 5.6)	49.7
10	2.74–2.75 m	49.1
11	1.55–1.61 m; 1.74–1.80 m	27.9
12	1.40–1.49 m; 1.63–1.69 m	27.6
13	2.33 ddd (12.5, 8.8, 1.9); 2.52–2.56 m	45.8
1'	-	142.7
2'/6'	7.23–7.27 m	126.4
3'/5'	7.30–7.36 m	128.7
4'	7.23–7.27 m	127.3

Compound **2a** was obtained as colorless crystals (CH_2Cl_2). In the HR-ESIMS spectrum, it exhibited a quasi-molecular ion at m/z 310.2501 $[\text{M}+\text{H}]^+$, establishing a molecular formula of $\text{C}_{17}\text{H}_{31}\text{N}_3\text{O}_2$. The molecular formula and the general framework of the ^1H and ^{13}C -NMR spectra (Table 2) indicated that compound **2a** was also a macrocyclic lactam spermidine alkaloid. The J -modulated ^{13}C -NMR and HSQC spectra displayed signals for an amide carbonyl carbon at δ_{C} 173.2, two methylenes adjacent to an amide function at δ_{C} 37.4 (CONH- CH_2) and δ_{C} 41.9 (CH_2 -CONH), three nitrogenated CH_2 (δ_{C} 45.8, 46.3 and 54.4), three methylenes (δ_{C} 22.2, 22.5 and 26.9) and one methine carbon at δ_{C} 56.5. These signals were consistent with a macrocyclic spermidine moiety and the partial structure was confirmed by analysis of the ^1H -NMR spectroscopic data (Table 2), and COSY and HMBC correlations (Figure 2). Moreover, two additional methylene carbons (δ_{C} 20.5, 28.9), one methyl at δ_{C} 10.7, and four deshielded methine, including one oxygenated (δ_{C} 73.0) and two olefinic methines at δ_{C} 123.7 and 126.6, were observed. The ^1H -NMR data and COSY correlations supported the presence of an ethyl moiety adjacent to an oxymethine and a double bond (δ_{H} 5.59, d, $J = 10$ Hz and 5.79–5.84, m). The connectivity between H-2/H-14, H-14/H-15, H-15/H₂-16, H₂-16/H-17, H-17/H-18, H-18/H₂-19 and H₂-19/H-20 was evident from COSY correlations suggesting that an unsaturated and hydroxylated chain was linked to the macrocyclic lactam through C-2. Further HMBC correlations revealed that C-17 was also connected through N-1 to the macrocyclic lactam (C-17/H-2 and C-17/2H-13). This was in accordance with the downfield chemical shift of C-17 (δ_{C} 63.5). The structure of **2a** was finally confirmed by single-crystal X-ray diffraction (Figure 3). Moreover, owing to the presence of chloride atoms in the crystal, the absolute configurations of C-2, C-17 and C-18 were assigned as 2*S*, 17*S* and 18*S* using the refinement of the Flack parameter [9]. On the basis of the above findings and the measurement of a positive optical rotation $[\alpha]_{\text{D}}^{25} = +12$ (c 0.84, MeOH), the structure of **2a** was unambiguously elucidated and the trivial name (+)-decaryine A was proposed. (+)-Decaryine A is different from palustrine previously isolated from *Equisetum palustre* [10,11] with respect to the orientation of the spermidine moiety in the macrocyclic lactam.

Table 2. ^1H and ^{13}C -NMR data for compounds **2a** and **2b** in CDCl_3 .

	2a		2b	
	δ_{H} (<i>J</i> in Hz) ^a	δ_{C}	δ_{H}	δ_{C} ^a
2	3.56–3.62 m	56.5	3.71–3.73, m	57.05
3	2.22 dd (13.7, 2.3); 2.52–2.62 m	41.9	2.40 dd (14.4, 3.8); 2.56–2.61 m	40.4
4	-	173.2	-	173.9
5	7.45–7.53 m	-	8.10 br s	-
6	2.97–3.03 m; 4.12–4.21 m	37.4	3.05–3.11 m; 3.76–3.81 m	38.8
7	2.02–2.06 m; 2.42–2.52 m	26.9	1.98–2.03 m; 2.20–2.26 m	25.8
8	2.87–2.95 m; 3.27–3.36 m	46.3	3.05–3.11 m; 3.16–3.22 m	49.5
10	2.87–2.95 m; 3.21–3.27 m	45.8	2.88 ddd (13.2, 11.5, 2.4); 2.97–3.01 m	50.1
11	1.70–1.77 m; 1.94–2.02 m	22.5	1.67–1.77 m; 1.98–2.09 m	26.4
12	1.73–1.82 m	22.1	1.57–1.78 m	24.7
13	2.28–2.37 m; 2.87–2.93 m	54.4	2.46–2.53 m; 2.56–2.61 m	46.8
14	5.59 d (10.4)	126.6	5.55–5.58 m	126.95
15	5.79–5.84 m	123.7	5.83 ddd (10.0, 4.3, 3.7)	127.0
16	1.85–1.92 m; 2.28–2.37 m	20.5	1.83–1.85 m	22.2
17	2.73–2.79 m	63.5	3.16–3.22 m	57.0
18	3.53 dt (11.0, 1.8)	73.0	3.54, ddd (9.5, 6.4, 5.2)	72.2
19	1.61–1.77 m	28.9	1.50–1.56 m	26.7
20	1.13 t (7.4)	10.7	1.06 t (7.3)	10.4

^a Assignment of ^{13}C -NMR data based on *J*-modulated, HSQC and HMBC spectra.

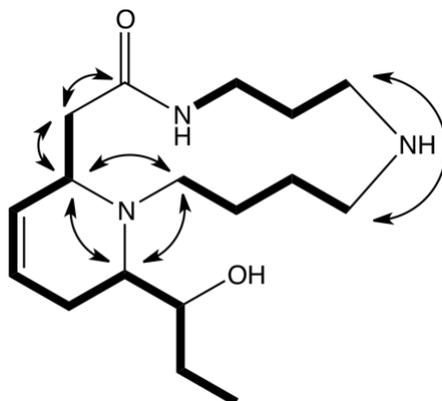
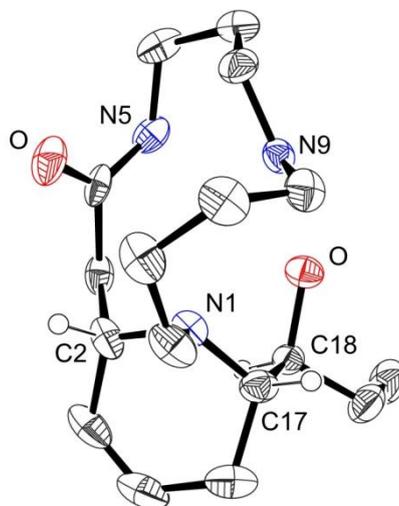
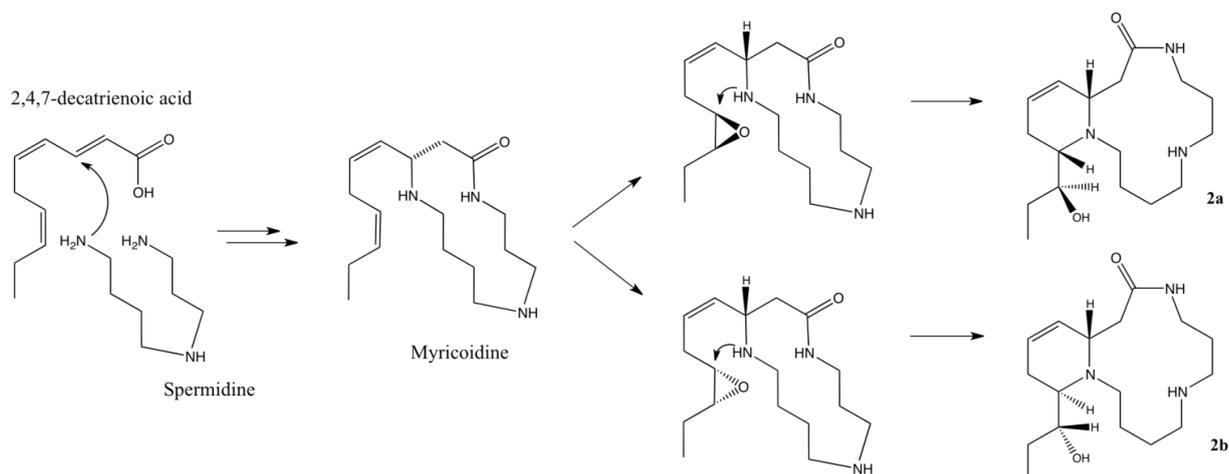
Figure 2. Selected COSY (**bold**) and HMBC correlations of compound **2a**.

Figure 3. Perspective drawing of the X-Ray structure of compound **2a** hydrochloride. Thermal ellipsoids are set at the 50% probability level. The solvent molecule, the chloride counter anion and all hydrogen atoms, except those on the chiral carbons, have been omitted for clarity.



Compound **2b** ($[\alpha]_D^{25} = -15$ (c 0.13, MeOH)) was isolated as a yellow oil. The molecular formula was determined as $C_{17}H_{31}N_3O_2$ by HR-ESIMS (m/z 310.2497 $[M+H]^+$). The ^{13}C -NMR data (Table 2) in combination with analysis of HSQC and HMBC spectra revealed the presence of 17 carbons similar to those of compound **2**. These data along with 1H -NMR data (Table 2) and extensive analysis of COSY and HMBC correlations suggested that compound **2b** is a diastereoisomer of **2a**. The relative and absolute configurations of C-2, C-17 and C-18 could not have been assessed by spectroscopic analyses. Nevertheless, considering the fact that all spermidine alkaloids, including those described here, exhibit the same three-dimensional orientation at C-2 [7,12–19], we can speculate that the latter carbon is (*S*)-configured. Schultz *et al.* [12] suggested that the first step of the biogenesis of spermidine alkaloids involves an enzyme-catalyzed Michael addition that would lead to the same stereochemistry at C-2 (Scheme 1). They also proposed a potential biogenetic pathway for palustrine [13] that can be adapted to the alkaloids of *A. decaryi* (Scheme 1). Indeed, a Michael addition of the primary amine of the butyl unit of spermidine to the α,β double bond of 2,4,7-decatrienoic acid, followed by amidification of the second primary amine may give a first intermediate, an alkaloid named myricoidine. Epoxidation of the double bond in the 17,18-position would then allow a nucleophilic attack by the nitrogen atom N-1 of the secondary amine at C-17 that induce the inversion of C-17 stereochemistry, with a concomitant 6-membered ring closure and hydroxyl group formation at C-18. In that case, we can postulate that epoxidation may occur on both faces of the double bond. Epoxide ring opening could then yield both (*2S*, *17S*, *18S*) and (*2S*, *17R*, *18R*) diastereoisomers as depicted in Scheme 1. Detection of dihydromyricoidine [19] in crude alkaloidal extract by MS analysis (see Supplementary Material) and notable difference in chemical shift of C-17 (δ_C 63.5 and δ_C 57.0 for compound **2a** and **2b**, respectively) are in agreement with this hypothesis. Therefore, the configuration of compound **2b**, named (–)-decaryine B, was proposed as *2S*, *17R*, *18R*.

Scheme 1. Potential biogenetic pathway for compounds **2a** and **2b**.

The known compounds lantibeside D (**3**) [20] and (*E*)-cinnamamide (**4**) [21] exhibited spectral data consistent with that reported. Finally, compounds **1,2a** and **3,4** were evaluated for their *in vitro* antiplasmodial activity. Compounds **1, 2a** and **4** were not active ($IC_{50} > 100 \mu M$), whereas the lignan **3** exhibited a weak antiplasmodial activity ($IC_{50} = 20.7 \mu M$).

3. Experimental

3.1. General

IR spectra were taken on a Perkin Elmer FT-IR Paragon 1000. 1H and ^{13}C -NMR spectra were recorded on a Bruker Avance 500 instrument in $CDCl_3$ or MeOD. HR-ESIMS spectra were recorded on Waters GTC Premier and Waters LCT spectrometers. Single-crystal-X-ray diffraction analysis was performed on a Bruker AXS SMART APEX II diffractometer at low temperature (193(2)K) and using Mo K_{α} radiation ($\lambda = 0.71073 \text{ \AA}$). Optical rotations were measured with a Perkin Elmer 241 polarimeter. Column chromatographies were performed over Merck silica gel 60A (40–63 μm) and Sephadex[®] LH-20. Analytical and semi-preparative TLC were performed on precoated Kieselgel 60 F₂₅₄ plates (Merck, 0.5 mm and 20 × 20, 1 or 2 mm, respectively) using Dragendroff reagent for visualization. CCDC 923477 contains the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/conts/retrieving.html (or from the CCDC, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033; e-mail: deposit@ccdc.cam.ac.uk).

3.2. Plant Material

Androya decaryi was collected twice by Frédéric Manjary (in 2008) and Claude Moulis (in 2009) in the Saint-Augustin region, in southwest Madagascar. Voucher specimens (ADFM2008 and ADCM2009) were deposited at the herbarium of the Institut Supérieur de Technologie of Toliara, Madagascar.

3.3. Extraction and Isolation

The dried and powdered leaves of a first sample of *A. decaryi* (108 g) were macerated with 100 mL of NH₄OH (28% aqueous) and then successively percolated with dichloromethane (2 L) and MeOH (1 L). The CH₂Cl₂ extract was evaporated under reduced pressure at 40 °C to give 3.58 g of a crude extract which was dissolved in CH₂Cl₂ (200 mL), and extracted with HCl (0.5 M, 3 × 100 mL). The combined aqueous layers were then made alkaline by adding NaOH to pH 12 and extracted three times with CH₂Cl₂ (200 mL) to obtain the crude alkaloidal fraction (582 mg). This fraction was subjected to Si gel CC eluting with a CH₂Cl₂-MeOH gradient to yield seven fractions (F1–7). F7 (110 mg) was subjected to a second Si gel CC in the same conditions to afford compound **1** (40 mg). F5 (150 mg) was rechromatographed over Si gel CC (CH₂Cl₂-MeOH gradient) and Sephadex LH 20 (CH₂Cl₂ then MeOH) to yield F521 (75 mg). Compound **2a** was repeatedly precipitated in MeOH/Et₂O (1/1) from F₅₂₁, and further crystallization of compound **2a** in CH₂Cl₂ afforded colorless crystal (34 mg).

The second sample of *A. decaryi* (1.5 kg) was extracted and partitioned according to the same procedure. A portion (2.8 g) of the crude alkaloidal fraction were further partitioned in a mixture of water (500 mL) and EtOAc (500 mL). The aqueous neutral layer was extracted thrice with EtOAc (500 mL) and the combined organic layers were concentrated under reduced pressure to yield AD2E1 (1 g). After alkalization to pH 12 (28% NH₄OH), the aqueous layer was extracted with EtOAc (3 × 500 mL) to give after concentration 1.5 g of AD2E2. A white solid precipitated in CH₂Cl₂ from AD2E1 to afford after filtering (*E*)-cinnamamide **4** (125 mg). The resulting filtrate was further concentrated (880 mg) and fractionated on Si gel CC (CH₂Cl₂-MeOH gradient) to yield compound **3** (30 mg). AD2E2 was subjected to combinations of Sephadex LH 20 (CH₂Cl₂), Si gel (CH₂Cl₂-MeOH gradient) chromatography and final semi-preparative TLC (CH₂Cl₂/MeOH/NHEt₂ 9/0.5/0.5) yielding compound **2b** (1.3 mg).

(–)-(2*S*)-2-Phenyl-1,5,9-triazacyclotridecan-4-one (**1**). Yellow oil; $[\alpha]_D^{25} = -13$ (c 3.0, MeOH); IR (film) ν_{\max} 3420, 3250, 3075, 2920, 1639; ¹H and ¹³C-NMR see Table 1; ESI-MS positive m/z [M+H]⁺ 276 (100); HR-ESIMS m/z [M+H]⁺ 276.2078 (calc for C₁₆H₂₆N₃O, 276.2076).

Decaryine A (**2a**). Colorless crystals; $[\alpha]_D^{25} = +12$ (c 0.84, MeOH); IR (film) ν_{\max} 3383, 3060, 2926, 1648, 1093; ¹H and ¹³C-NMR see Table 2; ESI-MS positive m/z [M+H]⁺ 310 (100); HR-ESIMS m/z [M+H]⁺ 310.2501 (calc for C₁₇H₃₂N₃O₂, 310.2495).

Decaryine B (**2b**). Yellow oil; $[\alpha]_D^{25} = -15$ (c 0.13, MeOH); ¹H and ¹³C-NMR see Table 2; ESI-MS positive m/z [M+H]⁺ 310 (100); HR-ESIMS m/z [M+H]⁺ 310.2497 (calc for C₁₇H₃₂N₃O₂, 310.2495).

Lantibeside D (**3**). White amorphous powder; $[\alpha]_D^{25} = +9$ (c 2.0, MeOH); ¹H and ¹³C-NMR see [20]; ESI-MS positive m/z [M+Na]⁺ 541 (100), [2M+Na]⁺ 1059 (63).

3.4. Antiplasmodial Bioassay

Parasites (FCM-29 strain) were cultured according to the method described by Trager and Jensen [22] with modifications described by Benoit *et al.* [23]. The cultures were synchronized every 48 h by 5% D-sorbitol lysis [24] (Merck, Darmstadt, Germany). The FCM-29 strain was considered as a

chloroquine-resistant strain (chloroquine IC₅₀: 145 nM). *In vitro* antimalarial activity testing was performed by [³H]-hypoxanthine (Amersham, Orsay, France) incorporation as described by Desjardins *et al.* [25] with modifications [26].

4. Conclusions

The phytochemical investigation of *Androya decaryi* led to the isolation of three macrocyclic spermidine alkaloids named (–)-(2*S*)-2-phenyl-1,5,9-triazacyclotridecan-4-one (**1**), (+)-decaryine A (**2a**) and (–)-decaryine B (**2b**). To the best of our knowledge, compound **1** was previously synthesized, but it is reported here for the first time as a natural product. This is also the first report of spermidine alkaloids in Scrophulariaceae. The absolute configurations of alkaloids **1** and **2a** were unambiguously elucidated, supporting the hypothesis described by Hesse and co-workers of a common initial biogenetic step. A similar biogenetic pathway was then postulated for compound **2a** and **2b**, allowing us to propose an absolute configuration for (–)-decaryine B (**2b**). The structural complexity of macrocyclic spermidine alkaloids has stimulated numerous syntheses, especially to unambiguously determine their absolute configuration [12,13,18,19,27,28]. Thus, enantioselective synthesis of (–)-decaryine B would confirm these hypotheses. Evaluation of their antiplasmodial properties showed that compounds **1**, **2a**, **3** and **4** exhibited no significant activity against *Plasmodium falciparum*.

Supplementary Materials

Supplementary materials can be accessed at: <http://www.mdpi.com/1420-3049/18/4/3962/s1>.

Acknowledgments

The authors gratefully acknowledge Amandine Fabre and Paul-Louis Ricard for technical assistance.

References

1. Oxelman, B.; Kornhall, P.; Olmstead, R.G.; Bremer, B. Further disintegration of Scrophulariaceae. *Taxon* **2005**, *54*, 411–425.
2. Petersen, I.; Eastman, R.; Lanzer, M. Drug-resistant malaria: Molecular mechanisms and implications for public health. *FEBS Lett.* **2011**, *585*, 1551–1562.
3. Bero, J.; Frédrich, M.; Quetin-Leclercq, J. Antimalarial compounds isolated from plants used in traditional medicine. *J. Pharm. Pharmacol.* **2009**, *61*, 1401–1433.
4. Bero, J.; Quetin-Leclercq, J. Natural products published in 2009 from plants traditionally used to treat malaria. *Planta Med.* **2011**, *77*, 631–640.
5. Nogueira, C.R.; Lopes, L.M.X. Antiplasmodial natural products. *Molecules* **2011**, *16*, 2146–2190.
6. Kuehne, P.; Linden, A.; Hesse, M. Asymmetric synthesis of the alkaloids mayfoline and *N*(1)-acetyl-*N*(1)-deoxymayfoline. *Helv. Chim. Acta* **1996**, *79*, 1085–1094.
7. Kuehne, P.; Guggisberg, A.; Hesse, M. Synthesis and chiroptical properties of 13-membered spermidine alkaloids (–)-(*S*)-Celacinine. (0)-(*S*)-celabenzine, (–)-(*S*)-celafurine, and (+)-(*S*)-viburnine. *Helv. Chim. Acta* **1997**, *80*, 1802–1808.

8. Wassermann, H.H.; Matsuyama, H.; Robinson, R. β -lactams as building blocks in the synthesis of macrocyclic spermine and spermidine alkaloids. *Tetrahedron* **2002**, *58*, 7177–7190.
9. Flack, H.D. On enantiomorph-polarity estimation. *Acta Cryst.* **1983**, *A39*, 876–881.
10. Kerrer, P.; Eugster, C.H. An alkaloid from *Equisetum palustre*. *Helv. Chim. Acta* **1948**, *31*, 1062–1066.
11. Natsume, M.; Ogawa, M. Total synthesis of (\pm)-palustrine and structure revision. *Chem. Pharm. Bull.* **1984**, *32*, 3789–3791.
12. Schultz, K.; Kuehne, P.; Häusermann, U.A.; Hesse, M. Absolute configuration of macrocyclic spermidine alkaloids. *Chirality* **1997**, *9*, 3789–3791.
13. Guggisberg, A.; Hesse, M. Natural polyamine derivatives—New aspects of their isolation, structure elucidation and synthesis. In *The Alkaloids, Chemistry and Biology*; Academic Press: San Diego, CA, USA, 1998; Volume 50, pp. 219–256.
14. Murata, T.; Miyase, T.; Yoshizaki, F. Meehanines A–K, Spermidine alkaloidal glycosides from *Meehania urticifolia*. *J. Nat. Prod.* **2009**, *72*, 1049–1056.
15. Murata, T.; Miyase, T.; Yoshizaki, F. Meehanines L–W, Spermidine alkaloidal glycosides from *Meehania urticifolia*. *J. Nat. Prod.* **2009**, *72*, 1937–1943.
16. Murata, T.; Miyase, T.; Yoshizaki, F. Cyclic spermidine alkaloids and flavone glycosides from *Meehania fargesii*. *Chem. Pharm. Bull.* **2010**, *58*, 696–702.
17. Wang, S.Q.; Ren, D.M.; Xiang, F.; Wang, X.N.; Zhu, C.J.; Yuan, H.Q.; Sun, L.M.; Ly, B.B.; Sun, X.J.; Lou, H.X. Dracotanoides A–D, Spermidine glycosides from *Dracocephalum tanguticum*: Structure and amide rotational barrier. *J. Nat. Prod.* **2009**, *72*, 1006–1010.
18. Khanjin, N.A.; Hesse, M. Synthesis of the spermidine alkaloids (–)-(2*R*,3*R*)- and (–)-(2*R*,3*S*)-3-hydrocelacinnine: Macrocyclisation with oxirane-ring opening and inversion via cyclic sulfamidates. *Helv. Chim. Acta* **2003**, *86*, 2028–2057.
19. Häusermann, U.A.; Linden, A.; Song, J.; Hesse, M. Total synthesis of (–)-(2*R*)-dihydromyricoidine. *Helv. Chim. Acta* **1996**, *79*, 1995–2003.
20. Li, T.; Hao, X.J.; Gu, Q.Q.; Zhu, W.M. Minor furofurano lignans from the Tibetan herb. *Lancea tibetica*. *Planta Med.* **2008**, *74*, 1391–1396.
21. Cai, M.Z.; Zhao, H.; Hu, W.Y.; Hale, P.D. Silica-supported Arsine Palladium(0) complex: A highly active and stereoselective catalyst for arylation of butyl acrylate and acrylamide. *Chin. J. Chem.* **2005**, *23*, 443–447.
22. Trager, W.; Jensen, J.B. Human malaria parasites in continuous culture. *Science* **1976**, *193*, 673–675.
23. Benoit, F.; Valentin, A.; Pelissier, Y.; Marion, C.; Dakuyo, Z.; Mallie, M.; Bastide, J.M. Antimalarial activity *in vitro* of *Cochlospermum tinctorium* tubercle extracts. *Trans. Roy. Soc. Trop. Med. Hyg.* **1995**, *89*, 217–218.
24. Lambros, C.; Vanderberg, J.P. Synchronization of *Plasmodium falciparum* erythrocytic stages in culture. *J. Parasitol.* **1979**, *65*, 418–420.
25. Desjardins, R.E.; Canfield, C.J.; Haynes, J.D.; Chulay, J.D. Quantitative assessment of antimalarial activity *in vitro* by a semiautomated microdilution technique. *Antimicrob. Agent Chemother.* **1979**, *16*, 710–718.

26. Valentin, A.; Benoit-Vical, F.; Moulis, C.; Stanislas, E.; Mallie, M.; Fouraste, I.; Bastide, J.-M. *In vitro* antimalarial activity of penduline, A bisbenzylisoquinoline from *Isopyrum thalictroides*. *Antimicrob. Agent Chemother.* **1997**, *41*, 2305–2307.
27. Häusermann, U.A.; Hesse, M. Total syntheses of (+)-(2*S*)-dihydromyricidine and (+)-(2*S*)-myricidine. *Tetrahedron Lett.* **1998**, *39*, 257–260.
28. Bienz, S.; Bisegger, P.; Guggisberg, A.; Hesse, M. Polyamine alkaloids. *Nat. Prod. Rep.* **2005**, *22*, 647–658.

Sample Availability: Samples of the compounds **1** and **2a** are available from the authors.

© 2013 by the authors; licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution license (<http://creativecommons.org/licenses/by/3.0/>).