

Multiple Monohydroxylation Products from *rac*-Camphor by Marine Fungus *Botryosphaeria* sp. Isolated from Marine Alga *Bostrychia radicans*

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This manuscript describes the biooxidation of *rac*-camphor using whole cells of marine-derived fungus *Botryosphaeria* sp. CBMAI 1197. The main biotransformation products of this monoterpenes were achieved via a hydroxylation reaction and occurred with 5 days of *rac*-camphor incubation. Products were identified by means of gas chromatography mass spectrometry (GC-MS) and nuclear magnetic resonance (NMR) data. The major hydroxylated products were 6-*endo*-hydroxycamphor, 6-*exo*-hydroxycamphor, 5-*exo*-hydroxycamphor, 5-*endo*-hydroxycamphor, 3-*exo*-hydroxycamphor and 8-hydroxycamphor. The 6-*exo*-hydroxycamphor was obtained through a retro-aldol reaction when 6-*endo*-hydroxycamphor was maintained in presence of CDCl₃; this isomerization was confirmed by ¹H NMR and GC-MS data.

Keywords: terpene, hydroxylation, biocatalysis, marine endophytic fungi

Introduction

The mono and sesquiterpenes belong to the largest class of secondary metabolites. They are known not only as raw materials for flavor and fragrance, but also as biologically active substances.¹ The (–)-camphor is a bicyclic monoterpenes and is one of the oldest known organic compounds. It is a major constituent of essence of dalmatian sage (*Salvia officinalis*).^{2,3} It has a camphoraceous odor and is used commercially as a moth repellent and in skin care formulations for cosmetic and pharmaceutical purposes.⁴

Monoterpene in plants have mainly ecological roles. They repel herbivores, minimize fungus, and attract

pollinators.¹ This class of compounds has been widely used for bactericidal, virucidal, fungicidal, antiparasitical, insecticidal, medicinal and cosmetic applications especially in pharmaceutical, sanitary, cosmetic, agricultural and food industries.^{5,6}

Considering the wide variety of applications and activities of monoterpenes, there are attempts to modify its basic structure to improve its properties. This can be accomplished by chemical synthesis and biotransformation reactions. However, organic synthesis is not typically satisfactory for performing structural modifications of terpenoids because of its low reactivity. This produces substances in poor yields.

The oxidation of inactivated C–H bonds under mild conditions has no equivalent in classical synthetic methodologies; P450 enzymes also have significant

potential in biotransformation applications. For example, cytochrome P450 from *Pseudomonas putida* catalyzed the selective oxidation of camphor to 5-*exo*-hydroxycamphor.⁷

There are few articles in the literature describing monohydroxylation of camphor. Nakahashi and Miyazawa⁸ used *Salmonella typhimurium* OY1000/2A6 of human cytochrome P450, employing the cofactor NADPH-P450 reductase. This hydroxylated the (–)-5-*exo*-hydroxycamphor and (–)-8-hydroxycamphor. Miyazawa and Miyamoto⁹ used the larvae of the common cutworm (*Spodoptera litura*) to biotransform *rac*-camphor to 5-*endo*-hydroxycamphor, 5-*exo*-hydroxycamphor (**5**) and 8-hydroxycamphor. However, to the best of our knowledge, there is not report on camphor biotransformation by marine fungi.

We recently investigated the bio-oxidation of natural products using marine-derived fungi.¹⁰ In this study, we used whole cells of Brazilian marine fungus *Botryosphaeria* sp. CBMAI 1197 in the biohydroxylation reaction of monoterpene *rac*-camphor (**1**). The fungus *Botryosphaeria* sp. CBMAI 1197 was isolated from red marine alga *Botrychia radicans*.¹¹ Therefore, different hydroxylated compounds were produced by fungal bioconversion of monoterpene.

Experimental

General methods

The *rac*-camphor (**1**) (> 96%) used in the biotransformation experiments was purchased from Sigma-Aldrich. Technal T-421 and Superohm orbital shakers were employed in the biocatalytic transformation experiments. Sterile materials were used for the biotransformation experiments, and *Botryosphaeria* sp. CBMAI 1197 was handled in a Veco biological safety cabinet. Thin-layer chromatography (TLC) was performed on pre-coated plates (aluminum foil, silica gel 60 F₂₅₄ Sorbent, 0.25 mm, from Merck).

Enzymatic products were purified by column chromatography (CC) on silica gel (SiO₂; 230-400 mesh, from Across) eluted with *n*-hexane/EtOAc mixtures in increasing polarity degrees. The products of *rac*-camphor (**1**) analyses were carried out using Shimadzu 2010 GC/FID equipped with an auto-injector AOC20i and DB-5MS (30 m × 0.25, 0.25 µm) column. The injector and detector temperature were maintained at 250 °C, the split ratio of the injector was 1:20, the carrier gas was He at 88.2 kPa, and the flux was 1.5 mL min⁻¹. The program used for gas chromatography (GC) analysis was as follows: initial temperature of 50 °C, final temperature of 232 °C, ramp rate of 7 °C, and final time of 30 min.

The mass spectrometry (MS) used electron capture detection, electronic impact of 70 eV, scanning speed of 1,000 Da sec⁻¹, scan interval of 0.50 fragments s⁻¹ and fragments detected from 40-500 Da. The ¹H and ¹³C nuclear magnetic resonance (NMR) spectra were obtained on a Bruker/AC-200 spectrometer (¹H at 200 MHz and ¹³C at 50 MHz). The spectra were obtained in deuterium chloroform (CDCl₃), and the chemical shifts are given in ppm with tetramethylsilane (TMS) as the internal standard.

Isolation, identification and preparation of stock cultures of marine-derived fungus *Botryosphaeria* sp. CBMAI 1197

The *Botryosphaeria* sp. CBMAI 1197 fungus strain was isolated from red marine alga *Botrychia radicans* collected in Ubatuba in the South Atlantic Ocean off the northern coast of the state of São Paulo, Brazil, in September 2007 by Prof Hosana M. Debonsi (FCFRP/ USP).¹¹ This alga was identified by conventional taxonomic methods by N. S. Yokoya (Botanical Institute of São Paulo, IBOT/SP, Brazil). A voucher specimen was deposited at the Herbarium of Botanical Institute of São Paulo, Brazil, under accession number SP 365678.

Stock cultures of the marine-derived fungus was stored in Petri dishes containing a 2% malt solid culture medium based on artificial sea water (ASW) [CaCl₂·2H₂O (1.36 g L⁻¹), MgCl₂·6H₂O (9.68 g L⁻¹), KCl (0.61 g L⁻¹), NaCl (30.0 g L⁻¹), Na₂HPO₄ (0.014 mg L⁻¹), Na₂SO₄ (3.47 g L⁻¹), NaHCO₃ (0.17 g L⁻¹), KBr (0.1 g L⁻¹), SrCl₂·6H₂O (0.040 g L⁻¹), H₃BO₃ (0.030 g L⁻¹)], agar (20 g L⁻¹), and malt extract (20 g L⁻¹). The pH of the medium was adjusted to 8 with 3 mol L⁻¹ KOH. The Petri dishes were maintained at 4 °C for later use.

Biotransformation of *rac*-camphor (**1**) by marine-derived fungus *Botryosphaeria* sp. CBMAI 1197

The fungus was cultured in 250 mL Erlenmeyer flasks containing 100 mL of liquid culture medium with 2% malt extract (3 days, at 32 °C) on an orbital shaker (150 rpm) in ASW. After this time, the *rac*-camphor (**1**) (50.0 mg) previously dissolved in 400 µL of dimethyl sulfoxide was added and mixed into the liquid culture medium. The biocatalytic reactions were analyzed periodically (24, 48, 72, and 120 h) by collecting 2 mL of samples and adding 2 mL of ethyl acetate with vortex mixing and then centrifuged at 6000 rpm for 6.0 min in a HERMLE Z-200 A. The organic phase was analyzed by GC-FID and GC-MS. The experiments were conducted in triplicate. After 120 h, the biotransformation products were isolated.

Purification of hydroxylated products obtained by biotransformation of *rac*-camphor (**1**)

The isolation of the hydroxylated products was performed using CC; with stationary phase flash silica gel 60 (230-400 mesh) from Across. The separation progress was monitored by TLC on pre-coated silica gel 60 F254 from Sorbent Technologies. The samples were observed under UV light (254 nm) and developed with anisaldehyde solution (100 mL of acetic acid, 1 mL of concentrated sulfuric acid and 1 mL of *p*-anisaldehyde) or alcoholic solution of vanillin (125 mL ethanol, 25 mL of concentrated sulfuric acid and 6 g vanillin). Hexane and ethyl acetate was used as the mobile phase for elution of components. This started with 100% hexane (175 mL) and the polarity gradually increased: 9.5:0.5 (75 mL), 9.0:1.0 (75 mL), 8.5:1.5 (75 mL), 8.0:2.0 (150 mL), 7.5:2.5 (200 mL), 7.0:3.0 (200 mL), and 6.5:3.5 (100 mL).

Results and Discussion

Biotransformation of *rac*-camphor (**1**) by the whole cells of marine-derived fungus *Botryosphaeria* sp. CBMAI 1197

Botryosphaeria sp. CBMAI 1197 transformed the *rac*-camphor (**1**) to different hydroxylated products **2-7**. The hydroxylation mediated by *Botryosphaeria* genus has been described in the literature in different natural products including biooxidation reactions of 3-keto-bisnorcholeno, nootkatone, ambrox, sclareol, sclareolide and valencene.¹²⁻¹⁴

We initially carried out a kinetic study of the biotransformation reaction of *rac*-camphor (**1**) up to 120 h. After this period, the chromatographic profile obtained by GC-flame ionization detection (FID) analysis was unchanged indicating that the biotransformation was stabilized in these conditions. The biotransformation of *rac*-camphor (**1**) was mediated by whole cells from *Botryosphaeria* sp. CBMAI

1197 fungus. This demonstrated that products **2-7** formed at different reaction times (24, 72 and 120 h) and were maintained by the same chromatographic profile (had no change) during the biotransformation process (Figure 1). A control stability experiment was performed in which the liquid culture medium (absent of fungus) was added to the *rac*-camphor (**1**); the compound was stable for at least 120 h (Figure 1).

Several compounds may undergo spontaneous reactions. These are sometimes mistakenly associated with enzymatic conversions. Thus, *rac*-camphor (**1**) was added to the culture medium in the absence of fungus to study stability. The compound remained intact. Thus, the peaks in Figure 1 are products **2-7** due to biotransformation of *rac*-camphor (**1**) by enzymes from *Botryosphaeria* sp. CBMAI 1197.

In the biotransformation of *rac*-camphor (**1**) by *Botryosphaeria* sp. CBMAI 1197 were obtained six main metabolites **2-7**, which were previously identified by mass spectrometry data. In addition, four substances (**3, 5-7**) were isolated in mixture by chromatographic column and identified by NMR and MS analyses. All of the data were compared with those described in the literature (Scheme 1).

MS analysis of metabolites **2-7** showed a specific molecular ion peak (M^{+*}) at m/z 168, indicating the presence of a hydroxyl group added to the substrate (152 g mol⁻¹) by *Botryosphaeria* sp. CBMAI 1197 (Figure 2).

Figure 3 shows the chromatogram (GC-MS) fractions (A-D) containing compounds **3-7** produced by biotransformation. The compounds were partially purified through normal phase chromatography column. These fractions showed the oxidized products **3-6**. Compounds **3** and **5** were isolated with 81 and 75% purity, respectively, and were identified by NMR analyses (Figures 3A-B). Metabolites **6** and **7** were obtained as a mixture. These molecules were structurally identified by NMR data (Figures 3C-D). The fraction obtained by chromatographic separation and analyzed by GC-MS showed compound **3** in a mixture of substances **5-6** (Figure 3C). Therefore,

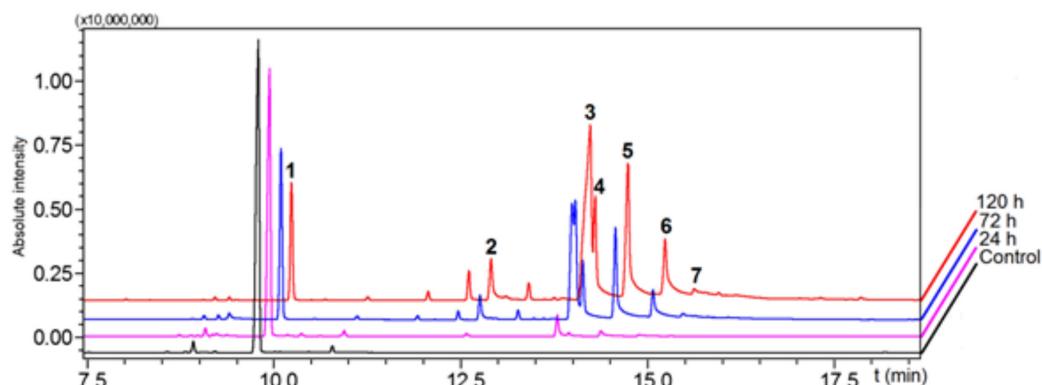
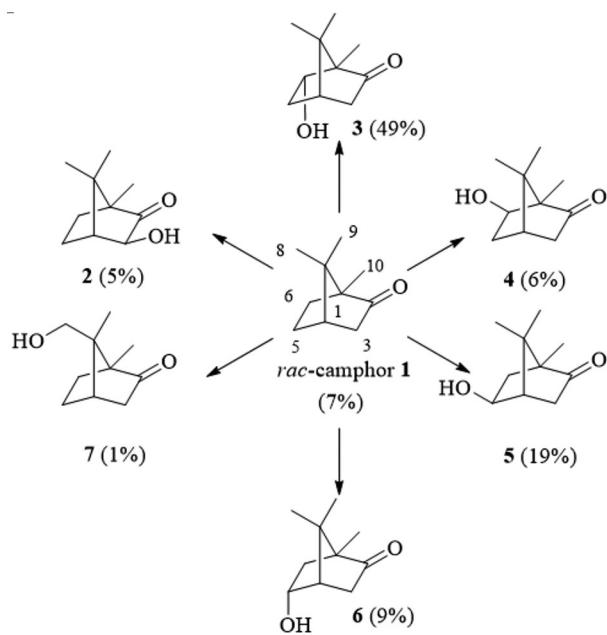


Figure 1. GC-FID analyses of biotransformation reaction of *rac*-camphor (**1**) by marine endophytic fungus *Botryosphaeria* sp. CBMAI 1197 in different times. Control: liquid culture medium plus *rac*-camphor (**1**) in fungus absence.



Scheme 1. Biohydroxylation of *rac*-camphor (**1**) by whole cells of *Botryosphaeria* sp. CBMAI 1197 yielding the compounds **2-7**.

metabolite **6** was in mixture of compound **3** and **5**. Compound **3** (majority in fraction C) was obtained in mixtures with metabolites **5-7**. Compound **7** was identified in this mixture and could be seen in fraction D (Figure 3D). In addition, compound **2** was not isolated, and compound **4** was obtained from isomerization of compound **3**.

The identification and characterization of hydroxylated compounds **2-7** are described here. The exact position of the hydroxyl group was confirmed by NMR and MS analyses of the substances (even in mixture). These were purified by flash chromatography column and compared with literature data.⁹ These substances obtained via biotransformation of *rac*-camphor (**1**) by *Botryosphaeria* sp. CBMAI 1197 were obtained as white solids.

The ¹H NMR spectra of the carbinolic hydrogen region showed that the fractions contained mixture of metabolites **3-7**. This explained the position of the hydroxyl groups in *rac*-camphor (**1**) (Supplementary Information).

Fraction A (Figure 3A) had a peak at δ 9.86. Surprisingly, this signal was not assigned to any hydrogen

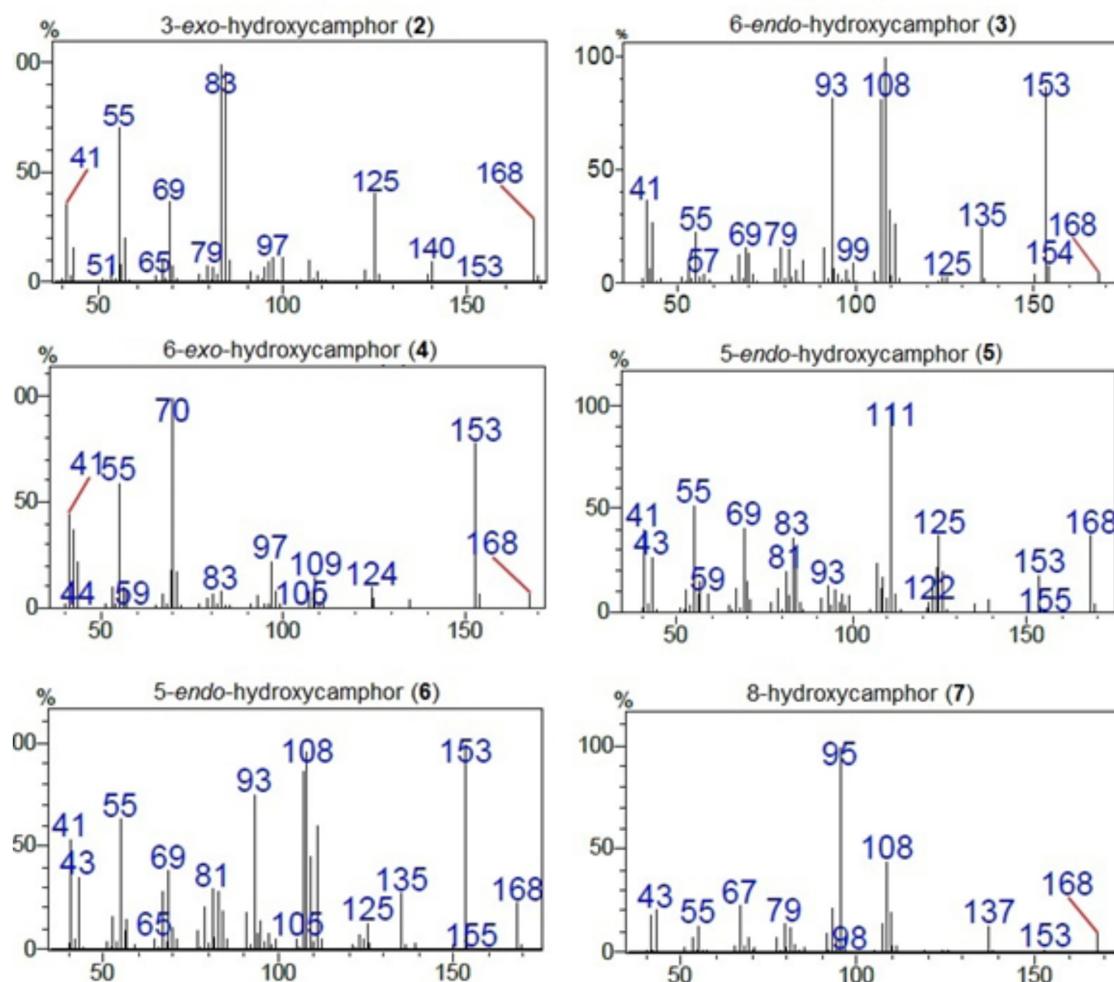
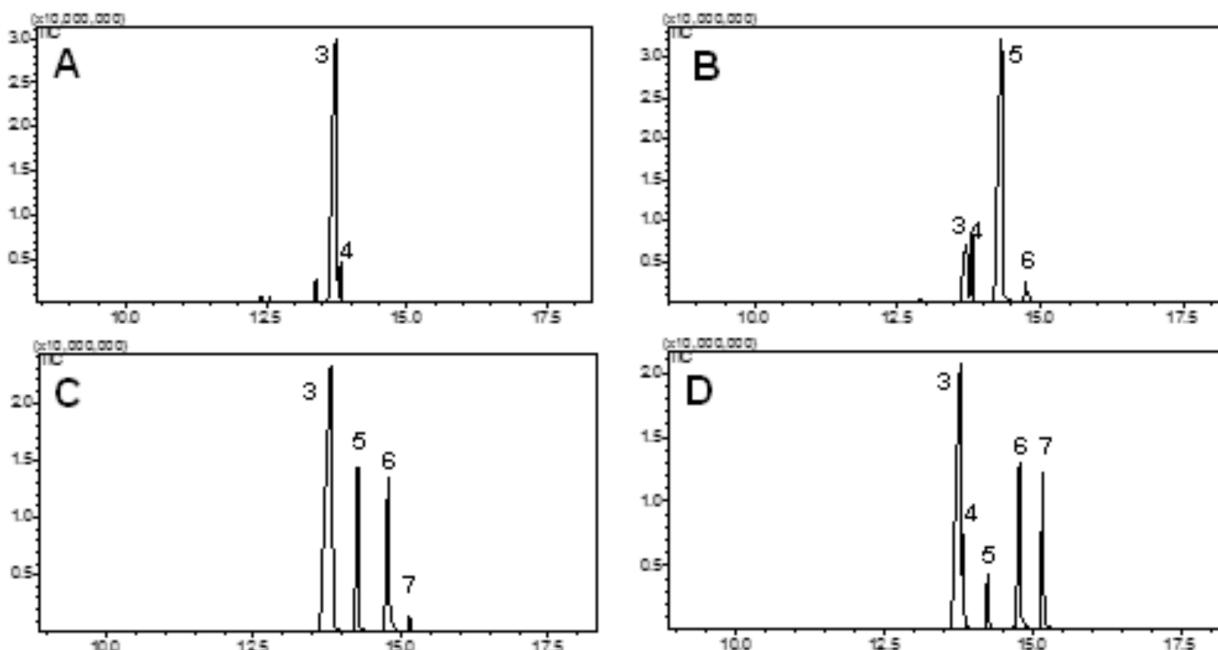
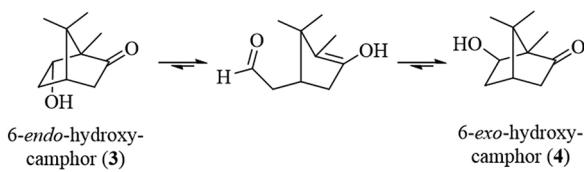


Figure 2. Mass spectra for the hydroxylated derivative metabolites **2-7** from *rac*-camphor (**1**) obtained by *Botryosphaeria* sp. CBMAI 1197.



Figures 3. (A-D) GC-MS (70 eV) chromatograms of the fractions 3A-D containing the compounds 3-7 yield by biotransformation of *rac*-camphor (**1**) using *Botryosphaeria* sp. CBMAI 1197.

of the hydroxyl-camphor metabolite. The signal at δ 9.86 was obtained from an aldehyde intermediate when the sample of 6-*endo*-hydroxycamphor (**3**) was solubilized in deuterated chloroform. According to Grogan *et al.*,¹⁵ the 6-*endo*-hydroxycamphor (**3**) undergoes an isomerization through a retro-aldol reaction confirmed with ^1H NMR shield at 9.86 ppm. Scheme 2 shows the spontaneous isomerization mechanism of 6-*endo*-hydroxycamphor (**3**) to obtain the 6-*exo*-hydroxycamphor (**4**) (NMR spectrum, Supplementary Information).



Scheme 2. Spontaneous isomerization of 6-*endo*-hydroxycamphor (**3**) to 6-*exo*-hydroxycamphor (**4**) in presence of CDCl_3 .

GC-MS analysis after dilution of the sample containing 6-*endo*-hydroxycamphor (**3**) in presence of CDCl_3 after 16 hours confirmed the existence of isomerization (Figure 4 and Scheme 2). The 6-*exo*-hydroxycamphor (**4**) signal increased and the peak of 6-*endo*-hydroxycamphor (**3**) decreased (Figures 3A and 4). The increased stability of the 6-*exo*-diastereomer (**4**) explains the balance shift.

In the ^1H NMR spectrum of fraction A (Figure 3A and Supplementary Information), we noted a double doublet at δ_{H} 4.18 ($J = 8.0$ and 2.0 Hz) that was attributed to 6-*exo*-

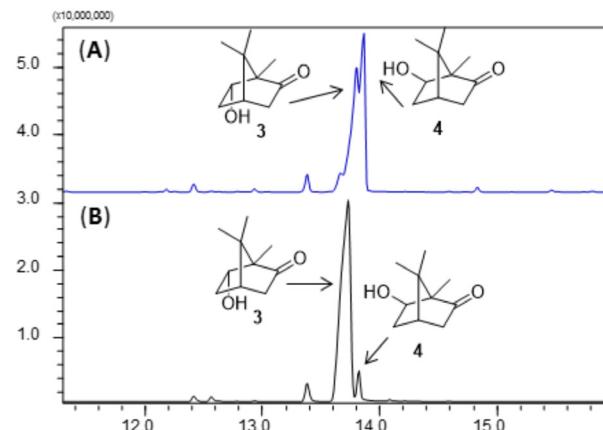


Figure 4. Chromatograms obtained by GC-MS analysis. (A) Spontaneous isomerization of 6-*endo*-hydroxycamphor (**3**) to 6-*exo*-hydroxycamphor (**4**) in presence of CDCl_3 ; (B) 6-*endo*-hydroxycamphor (**3**) in absence of CDCl_3 .

hydrogen of 6-*endo*-hydroxycamphor (**3**). Therefore, the structural characterization of hydroxylated metabolite **3** was carried out by ^1H NMR analysis. This showed the presence of methyl groups at 0.83 ppm (3H, s) and 0.98 ppm (6H, s). These signals were attributed to the hydrogens at positions 9, 8 and 10 of 6-*endo*-hydroxy-camphor (**3**) (Table 1). Also, the 6-*endo*-hydroxy-camphor (**3**) had coupling of 18 Hz between H-3_{endo} (1.99 ppm) and H-3_{exo} (2.35-2.48) for vicinal hydrogens in this position. At 2.15 ppm, we noted a triplet of the methinic hydrogen bound in position 4 of the 6-*endo*-hydroxy-camphor (**3**). The hydroxyl group assigned to the position H-6_{endo} was responsible for the signal at

4.18 ppm. This showed a constant coupling at 8.0 and 2.0 Hz. The complete structural determination of 6-*endo*-hydroxy-camphor (**3**) was confirmed by comparison of ¹H NMR data described in the literature (Table 1).⁹ Other characteristic signals can be seen in Table 1.

Table 1. ¹H NMR data of 6-*endo*-hydroxy-camphor (**3**) (200 MHz, CDCl₃) obtained by hydroxylation of *rac*-camphor (**1**) using *Botryosphaeria* sp. CBMAI 1197

Position	Experimental data for 6- <i>endo</i> -hydroxy-camphor (3)		Literature data for 6- <i>endo</i> -hydroxy-camphor (3) ⁹	
	δ_{H} (J in Hz) ^a / ppm	δ_{H} (J in Hz) ^b / ppm	δ_{H} (J in Hz) ^a / ppm	δ_{H} (J in Hz) ^b / ppm
3- <i>endo</i>	1.99 (d, 18.0)		1.98 (d, 18.5)	
3- <i>exo</i>	2.35-2.48 (m)		2.43 (ddd, 18.5, 5.0, 3.0)	
4	2.15 (t, 4.0)		2.14 (dd, 5.0, 4.5)	
5- <i>endo</i>	1.34 (dd, 14.0, 2.0)		1.33 (dd, 13.5, 2.5)	
5- <i>exo</i>	2.50-2.61 (m)		2.54 (dd, 13.5, 10.0, 4.5, 3)	
6- <i>exo</i>	4.18 (dd, 8.0, 2.0)		4.16 (dd, 10, 2.5)	
8 and 10	0.98 (s, 6H)		0.96 (s, 6H)	
9	0.83 (s, 3H)		0.82 (s, 3H)	

^a¹H NMR data (200 MHz, CDCl₃); ^b¹H NMR data (400 MHz, CDCl₃).

The fraction B chromatogram (Figure 3B) showed compound **5** as the majority presence. The carbinolic hydrogen region of the ¹H NMR spectrum showed a double doublet at 4.02 ppm that was attributed to the 5-*endo*-hydrogen yielding 5-*exo*-hydroxycamphor (**5**). This doublet had coupling constants of 6.0 and 4.0 Hz indicating an *endo*-*endo* correlation between the hydrogens in positions H-5 and H-6, and *endo*-*exo* between the 5-*endo*-hydrogen and 6-*exo*-hydrogen, respectively. The other signals were compared with the literature and corroborated the characterization of 5-*exo*-hydroxycamphor (**5**) (Table 2).⁹

The ¹H NMR spectrum of fraction C (Figure 3C and Supplementary Information) showed the presence of three groups with characteristic signal for carbinolic hydrogens. The majority signal at 4.18 ppm concerned the 6-*exo*-hydrogen of 6-*endo*-hydroxycamphor (**3**). The signal at 4.02 ppm was the 5-*endo*-hydrogen of 5-*exo*-hydroxycamphor (**5**), and the other signal type multiplet was at 4.58-4.66 ppm and was assigned to 5-*exo*-hydrogen of 5-*endo*-hydroxycamphor (**6**) (Figure 5). This characterization agrees with the data obtained by Miyazawa and Miyamoto⁹ who proposed substance **6** presents a shield in 4.64 ppm as a dddd (*J* = 9.5, 4.5, 4.0 and 2.0 Hz).

Fraction D (Figure 2D and Supplementary Information) showed hydroxylation at 6-*endo* for compound **3** (4.18 ppm, dd, *J* = 8.0, and 2.0 Hz), 5-*exo* for compound **5** (δ 4.02) and 5-*endo* for compound **6** (δ 4.58-4.66, m). In addition,

Table 2. ¹H NMR data of 5-*exo*-hydroxy-camphor (**5**) (200 MHz, CDCl₃) obtained by hydroxylation of *rac*-camphor (**1**) using *Botryosphaeria* sp. CBMAI 1197

Position	Experimental data for 5- <i>exo</i> -hydroxy-camphor (5)		Literature data for 5- <i>exo</i> -hydroxy-camphor (5) ⁹	
	δ_{H} (J in Hz) ^a / ppm	δ_{H} (J in Hz) ^b / ppm	δ_{H} (J in Hz) ^a / ppm	δ_{H} (J in Hz) ^b / ppm
3- <i>endo</i>	1.70 (d, 18.0)		1.70 (d, 18.5)	
3- <i>exo</i>	2.34 (d, 18.0, 6.0)		2.33 (dd, 18.5, 5.0)	
4	2.16 (d, 6.0)		2.16 (dd, 5.0, 1.0)	
5- <i>endo</i>	4.02 (dd, 6.0, 4.0)		4.02 (dd, 7.5, 3.5)	
6- <i>endo</i>	1.85 (m)		1.85 (dd, 14.0, 7.5)	
6- <i>exo</i>	1.81 (m)		1.79 (ddd, 14.0, 3.5, 1.0)	
8	0.85 (s, 3H)		0.85 (s, 6H)	
9	1.25 (s, 3H)		1.25 (s, 3H)	
10	0.93 (s, 3H)		0.93 (s, 3H)	

^a¹H NMR data (200 MHz, CDCl₃); ^b¹H NMR data (400 MHz, CDCl₃).

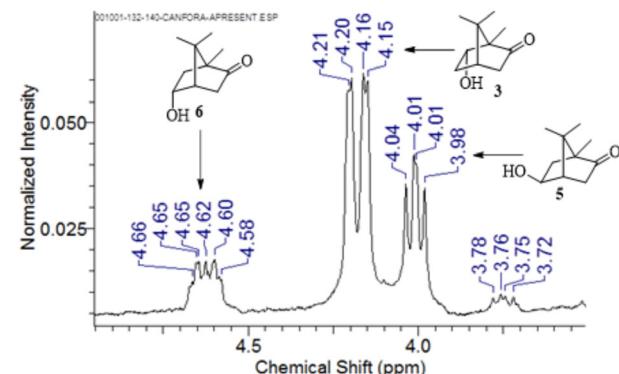


Figure 5. Specific signals obtained by ¹H NMR spectrum for carbinolic hydrogens of monohydroxylated derivatives (**3**, **5** and **6**) of *rac*-camphor (**1**) obtained from fraction C.

there were two doublets at 3.50 ppm (d, *J* = 12.0 Hz) and 3.74 ppm (d, *J* = 12.0 Hz). This is usually designated as the roof effect and was allocated to two diastereotopic carbinolic hydrogens of 8-hydroxycamphor (**7**) (Figure 6). Miyazawa and Miyamoto⁹ assigned readings on 3.52 ppm (d, *J* = 11.0 Hz) and 3.74 ppm (d, *J* = 11.0 Hz) to hydrogens of C-8 of 8-hydroxycamphor (**7**).

Finally, compound **2** was not isolated, and there are ten sites of hydroxylation in the *rac*-camphor (**1**) (Figure 7). Compound **2** might have been a possible hydroxylated stereoisomer according to the data of spectrum mass. This was suggested at position-2 (Figure 2 and Supplementary Information).

This study showed the camphor biotransformation by a marine environment fungus. The identification of monohydroxylated products from the oxidation of camphor is important for the identification of metabolites or the use of hydroxylated derivatives of camphor in pharmaceutical formulations.

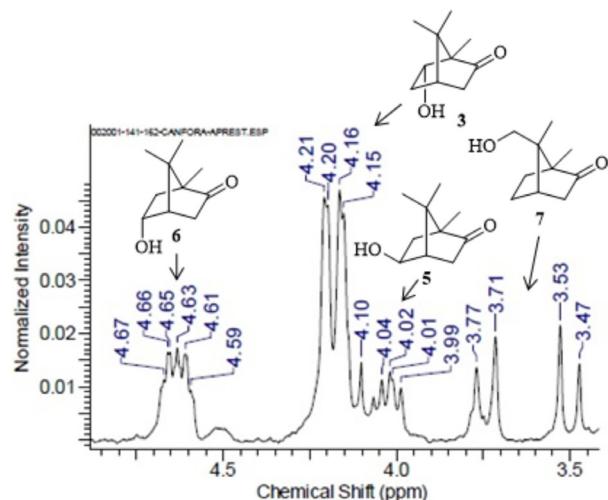


Figure 6. Specific signals obtained by ^1H NMR spectrum for carbinolic hydrogens of monohydroxylated derivatives (**3**, **5**, **6** and **7**) of *rac*-camphor (**1**) obtained from fraction D.

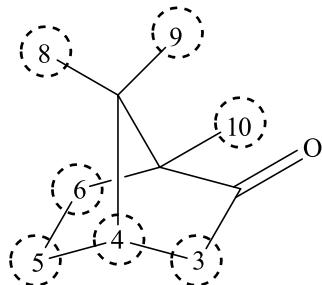


Figure 7. Possible sites of monohydroxylation of *rac*-camphor (**1**) to obtain stereoisomers. Compounds identified had the positions C-3, C-5, C-6 and C-8 hydroxylated.

Conclusions

The Brazilian marine-derived fungus *Botryosphaeria* sp. CBMAI 1197 is an efficient biocatalyst that promotes mono-biohydroxylation of *rac*-camphor (**1**). The main biotransformation product was 6-*endo*-hydroxycamphor (**3**) reaching 49% after 5 days of reaction followed by 5-*exo*-hydroxycamphor (**5**), 5-*endo*-hydroxycamphor (**6**), 6-*exo*-hydroxycamphor (**4**) and 8-hydroxycamphor (**7**). Chemical modifications in non-activated carbon atoms such as sp^3 camphor carbons usually cannot be done by conventional organic synthesis.

Supplementary Information

We would like to inform that this manuscript presents to the reader the supplementary information (NMR spectra and proposal of MS fragmentation for identified compounds), available free of charge at <http://jbcs.sbj.org.br>.

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