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Javier Moraga, Berengere Dalmais, Inmaculada Izquierdo-Bueno, Josefina Aleu, James R. Hanson, et al.. Genetic and molecular basis of botrydial biosynthesis. Connecting cytochrome P450-encoding genes to biosynthetic intermediates.. ACS Chemical Biology, 2016, 11 (10), pp.2838-2846. 10.1021/ac-schembio.6b00581 . hal-01531674

HAL Id: hal-01531674

<https://hal.science/hal-01531674>

Submitted on 31 Jan 2024

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Genetic and Molecular Basis of Botrydial Biosynthesis: Connecting Cytochrome P450-Encoding Genes to Biosynthetic Intermediates

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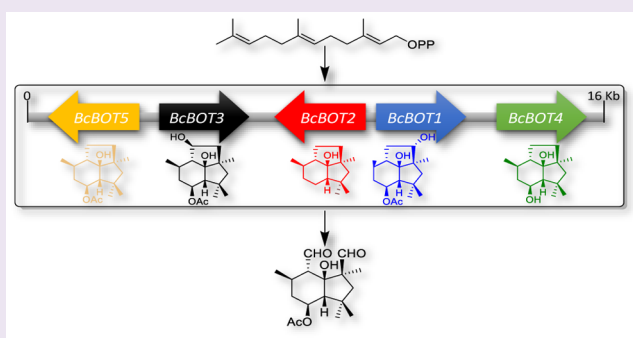
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S Supporting Information

ABSTRACT: Over two hundred species of plants can be infected by the phytopathogenic fungus *Botrytis cinerea* under a range of different environmental conditions. In response to these, the fungus produces unique terpenoid and polyketide metabolites. Parts of the plants may be killed by the phytotoxin botrydial, enabling the fungus to feed on the dead cells. In this paper, we describe the genetic and molecular basis of botrydial biosynthesis and the function of the five genes of the genome of *B. cinerea* that together constitute the botrydial biosynthetic gene cluster. Genes *BcBOT3* and *BcBOT4*, encoding two cytochrome P450 monooxygenases, were inactivated by homologous recombination and were shown to catalyze regio- and stereospecific hydroxylations at the carbons C-10 and C-4, respectively, of the presilphiperfolan-8 β -ol skeleton. The null mutants, *bcbot3 Δ and *bcbot4 Δ , accumulated key intermediates in the botrydial biosynthesis enabling the complete genetic and molecular basis of the botrydial biosynthetic pathway to be established. Furthermore, the *bcbot4 Δ mutant overproduced a significant number of polyketides, which included, in addition to known botcinins, botrylactones and cinbotolide A, two new botrylactones and two new cinbotolides, cinbotolides B and C.***



Fungal secondary metabolites have a wide impact on human health, medicine, and agriculture. While many secondary metabolites including antibiotics, immunosuppressants, and phytohormones are beneficial, other such as phytotoxins and mycotoxins are harmful to plants, humans, and animals.¹ In order to manage or modify the biosynthesis of secondary metabolites, we need to identify the location of the genes encoding the relevant biosynthetic enzymes. The genes that encode the biosynthesis of structurally related secondary metabolites are often clustered. The horizontal gene transfer (HGT)² between distant fungi may be facilitated by such clustering, conferring an evolutionary advantage.^{3–6}

The Ascomycete *Botrytis cinerea* is the cause of “gray mold” disease of a wide range of dicoyledons under a variety of environmental conditions in which its secondary metabolites (SM) confer a selective advantage. *B. cinerea* has the characteristics of a necrotroph in which the infection may be mediated by the production of phytotoxins and reactive oxygen species, as well as by inducing an oxidative burst in the plant.^{7,8} A number of unique terpenoids and polyketides have been obtained from the mycelium. These include two groups of nonspecific phytotoxins. The first are the sesquiterpenoid botryanes,⁹ botrydial (1) and dihydrobotrydial (2). The second are the polyketide lactones, botrylactone (3)¹⁰ together with

botcinic (4) and botcinic acid (5),¹¹ and their cyclic relatives, the botcinins (6, 7).¹² Cinbotolide (8) is a new polyketide that has been reported and contains a carbon chain that possesses structural features and a stereochemistry reminiscent of the botcinins and botrylactone (Figure 1).¹³

Botrydial (1) is the major phytotoxin^{14,15} of *B. cinerea*, and it has been shown to reproduce the symptoms of the infection.^{16,17} Spectroscopic, chemical, and X-ray methods^{18–20} were used to establish the irregular sesquiterpenoid structures of these botryane metabolites. Their biosynthesis via farnesyl diphosphate (FPP, Figure 2) has been elucidated.^{21–24}

The genome of *B. cinerea* has been sequenced leading to the identification of the genes that are responsible for the synthesis of its secondary metabolites. Forty-four genes that encode key enzymes (KE) that are responsible for the committed biosynthetic steps were identified from sequences in two strains, BO5.10 (B cero five.ten) and T4.^{25,26} Typical of the secondary metabolism of filamentous fungi, the majority of the genes that encode these key enzymes are part of biosynthetic

Received: July 6, 2016

Accepted: August 16, 2016

Published: August 16, 2016

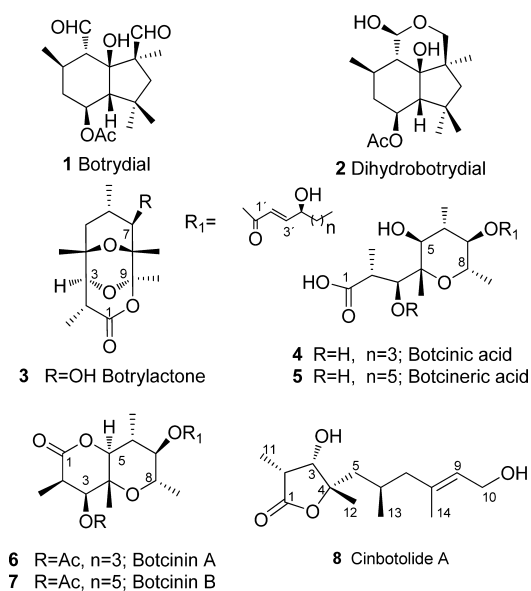


Figure 1. Some metabolites isolated from *Botrytis cinerea*.

gene clusters. Among them, the botrydial biosynthetic gene cluster has been identified (Figure 3).^{27,28} The cluster, which consists of five genes (*BcBOT1* to *BcBOT5*) (for *Botrytis cinerea*



Figure 3. Scheme of the gene cluster involved in the biosynthesis of botrydial. Dashed line indicates AT-rich DNA (more than 80% of A and T content).

BOTrydial) involves one gene coding for a sesquiterpene cyclase (*BcBOT2*), three monooxygenases (*BcBOT1*, *BcBOT3*, and *BcBOT4*), and an acetyl transferase (*BcBOT5*).

Gene deactivation studies, which had been carried out previously, had shown that the *BcBOT1* gene, encoding a P450 monooxygenase is responsible for one of the final steps in botrydial biosynthesis,²⁷ while *BcBOT2* codes for a sesquiterpene cyclase (presilphiperfolan-8 β -ol (PSP) synthase),²⁸ which is an early step in botrydial biosynthesis (Figure 2).

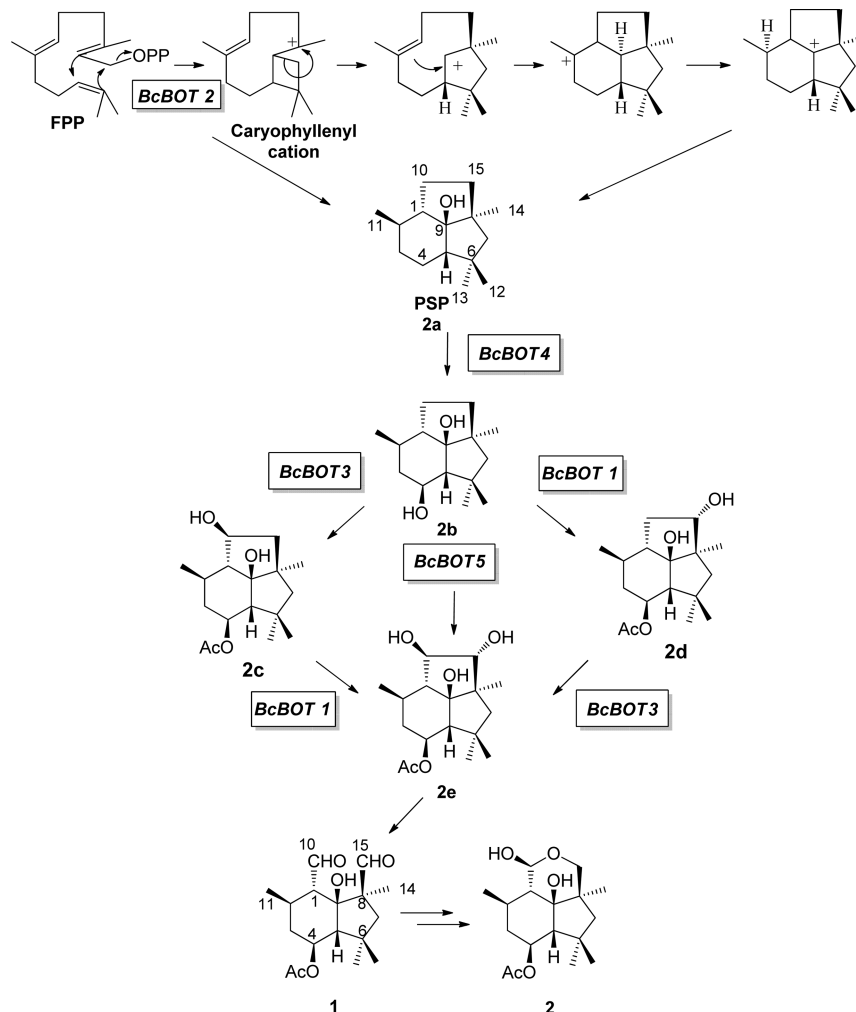


Figure 2. Biosynthetic pathway to botrydial (1).

The P450 monooxygenases, under mild reaction conditions, catalyze regio- and stereospecific oxidations of nonactivated hydrocarbons, to yield interesting chemical transformations in biosynthetic chemistry.²⁹

This paper describes the molecular sequence of botrydial biosynthesis and the functions of the three cytochrome P450 monooxygenases involved in botrydial biosynthesis in terms of the hydroxylations at C-4, C-10, and C-15 on the PSP carbon skeleton. The structures of four new polyketide metabolites, which were isolated from the *bcbot4Δ* mutant, are also reported.

RESULTS AND DISCUSSION

Previous experiments using isotopically labeled acetate and mevalonate established the origin of the carbon skeleton of botrydial (**1**) and the occurrence of a backbone rearrangement and 1,3-hydride shift during the cyclization of FPP.^{9,24,30} The original mechanistic proposal (Figure 2) has been corroborated by the incubation of labeled FPP with a recombinant PSP synthase (BcBOT2) to afford the labeled intermediate tricyclic alcohol, presilphiperfolan-8β-ol (**2a**, PSP, probotryane).³¹ This intermediate **2a** is converted to botrydial (**1**) by the action of the three cytochrome P450s encoded by the co-regulated genes *BcBOT1*, *BcBOT3*, and *BcBOT4* together with the acetyltransferase (BcBOT5), which function in an as yet unknown sequence.

Deletion of the Genes Encoding BcBOT3 and BcBOT4 Monooxygenases. Previous studies of the amino acid sequences of the BcBOT3 and BcBOT4 proteins have revealed the presence of conserved P450 monooxygenase domains, which suggested that they could be involved in botrydial biosynthesis.^{27,28} A gene knockout (KO) approach was used to confirm this hypothesis. Two KO DNA cassettes containing a nourseothricin resistance gene in place of the *BcBOT* genes, were constructed by PCR fusion and employed to transform protoplasts of the B05.10 wild-type strains (for details, see Methods). The transformation resulted in two independent *bcbot3Δ* and two independent *bcbot4Δ* mutants. PCR analyses verified integration of the replacement fragment at the targeted loci of the genome and the absence of wild-type nuclei. Both mutants grow and produced conidia similarly to the parent strain B05.10 on a standard rich medium (data not shown).

Chemical Characterization of the *bcbot3Δ* and *bcbot4Δ* Mutants. The KO *bcbot3Δ* and *bcbot4Δ* mutants, which had been derived from the B05.10 strain, were grown on a solid malt agar medium. The metabolites were recovered in ethyl acetate and purified by column chromatography on silica gel. The products were finally purified by HPLC to yield the metabolites shown in Figure 4. The results indicated that none of the mutants produced botrydial (**1**), confirming that these mutations were involved in botrydial biosynthesis.

The chromatographic study, characterization, and quantification of the metabolites biosynthesized by the *bcbot3Δ* mutant showed, in addition to small amounts of botcinin A (**6**), high levels, (35 mg/L), of the biosynthetic intermediate 4β-acetoxy-15α-hydroxyprobotryane (**2d**), Figure 2, which was identified by extensive NMR studies and comparison with authentic samples. In addition, the new metabolite **9** was isolated and characterized.

Compound **9** was obtained as colorless oil. It showed a molecular ion in its HRMS (ESI+) at $m/z = 253.1807$ [$(C_{15}H_{25}O_3)$, $M + H$]⁺ in agreement with the ¹³C NMR and DEPT spectra. The ¹H and ¹³C NMR spectra of compound **9**

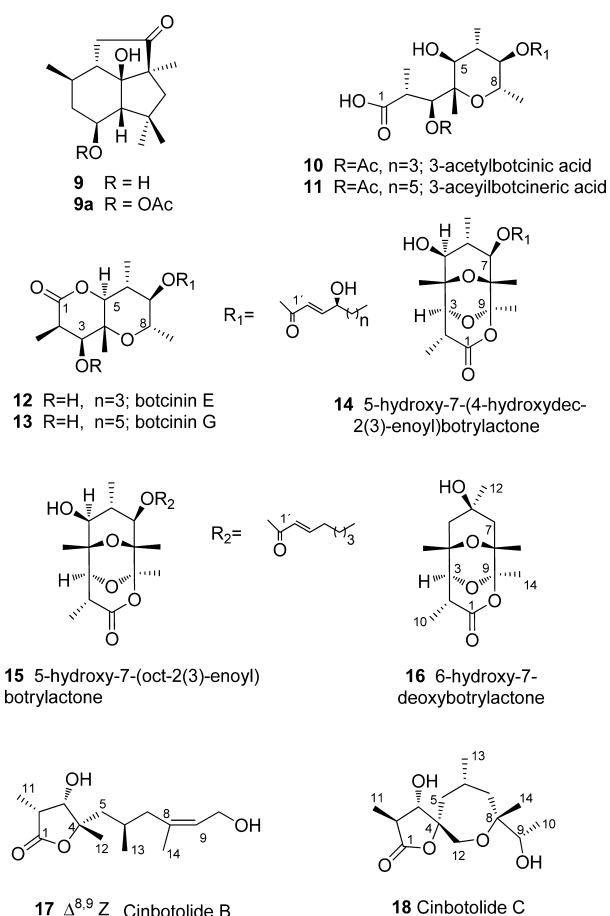


Figure 4. Some metabolites isolated from mutants *bcbot3Δ* and *bcbot4Δ*.

contained a similar pattern of signals to the probotryane intermediate (**2d**). However, the signal for the C-4 CH(OH) appeared at δ 4.04, while the signals for the acetate group and the C-15 CH(OH) were missing. There was a signal in the ¹³C NMR spectrum at δ 211.0 characteristic of a cyclopentanone, while the H-10 proton resonances were shifted. These spectroscopic data indicated that compound **9** was the oxidation product of the biosynthetic intermediate **2d** possessing a carbonyl group at C-15. Acetylation of compound **9** yielded a monoacetylated derivative **9a** that exhibited the characteristic ¹H NMR signals of a methyl group at δ 2.03 and a signal, ddd, at δ 5.11 for the proton geminal to the acetate. The proton and carbon-13 NMR signals of both compounds **9** and **9a** were assigned by detailed analysis of their COSY, NOESY, HSQC, and HMBC spectra (Supplementary Table S1). These confirmed their structures and relative stereochemistry.

The fermentation broth of mutant *bcbot4Δ* was chromatographed following the same procedure used for the KO mutant *bcbot3Δ*. Final purification by HPLC and GC-MS yielded the metabolites **2a** and **3–8** (Figures 1 and 2) and **10–18** show, in Figure 4. Interestingly, in addition to the key probotryane intermediate **2a**, from the cyclization of FPP by the sesquiterpene cyclase BcBOT2, an overproduction of polyketides was observed yielding the new polyketides **15–18**, Figure 4.

The probotryane **2a**, which was accumulated by the *bcbot4Δ* mutant, was identified in the hexane fraction by comparison with an authentic sample. Although it had not been found in



Figure 5. Comparative metabolites profiled by GC-MS analysis, for determination of probotryane **2a** in the strain broths. (A) Chromatograms of **2a** and of the extract obtained from the wild-type, B05.10 at 4, 8, and 10 days. (B) Chromatograms of **2a** and of the extract obtained from the mutant *bcbot4Δ* at 4, 8, and 10 days.

the wild-type (Figure 5A), it was readily identified in the *bcbot4Δ* mutant fermentation after 4, 8, and 10 days (Figure 5B).

Chemical Characterization of Polyketides Isolated from *bcbot4Δ* Mutant. The metabolites **15** and **16** showed a pattern of NMR signals that were characteristic of the botrylactones.¹¹ The ¹³C NMR contained 22 signals indicating that the side chain contained eight carbon atoms. The ¹H NMR of compound **15** resembled that of **14**. The most significant differences were the multiplicity of the signals corresponding to the double bond and the absence of the signal of a geminal proton to a hydroxyl group in the side chain. Thus, the vinylic proton signals appeared as doublet of triplets at δ 7.04, H-3', and a doublet corresponding to H-2', δ 5.84, indicating that compound **15** possessed a side chain that was similar to that of compound **14** but with eight carbon atoms and without a hydroxyl group at C-4'.

The HRMS, which was supported by the ¹³C NMR spectrum, showed that compound **15** possessed a molecular

formula of C₂₂H₃₄O₇. The COSY, HSQC and HMBC spectra, and NOE experiments (Supplementary Table S2) led to the structure 5-hydroxy-7-(oct-2-enoyl)botrylactone for compound **15**.

Compound **16** showed ¹H NMR signals that were comparable to botrylactone (**3**).¹¹ The main differences were the lack of the signal typical of the C-7 CH(OH) while the signal assigned to the C-6 methyl group was a singlet rather than a doublet. The HRMS and ¹³C NMR spectra indicated that the compound **16** had a molecular formula C₁₄H₂₂O₃. All the carbons and their associated proton signals were assigned from an examination of the COSY, HSQC, HMBC, and NOESY spectra (Table S3), in accord with the structure and relative stereochemistry of compound **16**. The relative stereochemistry of the chiral center at C-6 followed from the NOE experiments. The rigid structure of the compound meant that a β configuration of the C-6 methyl group would produce NOE interactions with the C-4 and C-8 methyl groups (Supplementary Figure S1), which would not be observed

with its epimer. However, irradiation of the C-4 methyl group gave NOE interactions with H-2 and H-5 β while irradiation of the C-8 methyl signals revealed interactions with H-7 β and H-14. Furthermore, irradiation of the C-6 methyl signal only gave NOE interactions with H-5 α and H-7 α indicating an α rather than a β configuration for this methyl group. These data led to the structure and stereochemistry of 6- β -hydroxy-7-deoxybotrylactone for **16** (Table S3, Figure S1).

The ^1H NMR signals of compound **17** resembled those of cinbotolide A (**8**).¹⁵ There was one primary and one secondary hydroxyl group and one double bond in its structure. The main differences were the multiplicities of the $-\text{CH}_2\text{OH}$ signal and the C-9 vinylic proton resonance. In addition to the signals assigned to the methyl groups at C-2, C-4, C-6, and C-8, there was a double-doublet at δ 5.39 (H-9) and signals at δ 4.15 (dd, $J = 12.0, 6.9$ Hz) and 4.05 (d, $J = 5.4$ Hz) attributed to H-10 and H-3, respectively.

The HRMS together with the ^{13}C NMR spectrum was consistent with a molecular formula $\text{C}_{14}\text{H}_{24}\text{O}_4$. Examination of the HSQC, HMBC, and COSY spectra enabled all the carbons and their associated ^1H NMR signals to be assigned (Table S4). Thus, compound **17**, which was named cinbotolide B, was the “Z” isomer of cinbotolide A (**8**).

The molecular formula of compound **18** was shown by HRMS to be $\text{C}_{14}\text{H}_{24}\text{O}_5$. The ^{13}C NMR spectrum contained signals corresponding to four methyl, three methylene, and four methine groups, together with three quaternary carbons. The latter included a signal at δ 177.3 ppm attributed to a lactone. The four methyl group signals in the ^1H NMR spectrum were three doublets (δ 0.98, 1.14, and 1.26) and a singlet (δ 1.21). The pattern of signals was typical of a cinbotolide lactone containing two hydroxyl groups. The proposed structure **18** was consistent with the HSQC, HMBC, and COSY spectra (Table S5). The stereochemistry was deduced from NOE experiments. When H-3 was irradiated, there were enhancements to the signals corresponding to H-2, H-11, and H-5 β , and when H-2 was irradiated there were enhancements to the signals corresponding to H-3, H-11, and in particular H-5 α . Consequently compound **18** was a C-2 epimer (H-2 α) of the cinbotolides A and B (**8** and **17**). This epimerization at C-2 has been observed in other polyketides biosynthesized by this fungus,¹¹ and it was supported by conformational study carried out with molecular mechanics calculation (Figures S1 and S2). When H-6 was irradiated, NOE effects were observed at H-14, H-12, H-5, and H-7. A NOE effect on H-2 and H-5 β was observed on irradiation of H-5 α , while effects on H-12, H-9, H-6, and H-10 were produced by irradiation of H-14. These effects were consistent with the structure and stereochemistry of **18**, named cinbotolide C. In this cinbotolide, epimerization has taken place at C-2, and the C-4 methyl group (C-12) has been oxidized and an ether bridge has been formed to C-8.

Genetic and Molecular Basis of Botrydial Biosynthesis. Although the biochemical function of the enzymes BcBOT1 and BcBOT2 is known, the functional role of the proteins BcBOT3, BcBOT4, and BcBOT5 are still unknown. In order to understand the molecular basis and to complete the biosynthetic pathway to botrydial (**1**), the KO mutants for the cytochrome P450 monooxygenases BcBOT3 and BcBOT4 encoding genes were constructed, and their secondary metabolism was studied from the chemical point of view to identify the potential biosynthetic intermediates that were produced and to characterize both of the P450 monooxygenases.

Previous studies showed that BcBOT2 protein catalyzes the cyclization of FPP to the sesquiterpene alcohol PSP, **2a**, and is required for the production of botryane and probotryane sesquiterpenes, which firmly established that BcBOT2 is responsible for the initial step of the botrydial biosynthesis (**1**).^{28,31} On the other hand, inactivation of the *BcBOT1* gene demonstrated that those mutants were unable to produce 10 β ,15 α -dihydroxyprobotryane (**2e**), botrydial (**1**), or some of its relatives. Instead the intermediate **2c** was accumulated, indicating that the monooxygenase BcBOT1 was responsible for the regiospecific hydroxylation at C-15, in intermediate **2b**, to give **2d** in botrydial biosynthesis, Figure 2.²⁷

In this context, the knock out *bcbot3 Δ* and *bcbot4 Δ* mutants were fermented, and their broths were examined to identify the secondary metabolites that were produced. Compounds **2d** and **9** (Figures 2 and 4) were isolated from the broth of the mutant *bcbot3 Δ* and characterized. The large amount of the metabolite **2d** that was isolated clearly indicated that the monooxygenase BcBOT3 is involved in the regio- and stereospecific β hydroxylation at C-10 on the skeleton of intermediate **2b**, yielding intermediate **2c**, and thus the hydroxylated derivative at C-15, **2d**, accumulated.

It was observed that the *bcbot3 Δ* mutant did not produce botrydial (**1**) and only produces small amounts of botcinin A (**6**). This result was unexpected, since we had previously established that the suppression or inhibition of the biosynthetic pathway leading to botrydial (**1**) induced the overproduction of the second family of toxins, botcinic and botcinic acids (**4**, **5**) and botcinins A and B (**6**, **7**).³² These results seem to suggest that the metabolic system of the fungus behaved as though the production of botrydial (**1**) was not inhibited, keeping the gene expression and the production of botcinins low. This result leads us to propose that this P450 monooxygenase as a good target for controlling the fungus and its pathogenicity by its inhibition.

The mutant *bcbot4 Δ* accumulated the probotryane **2a** in the less polar fractions. This was established by GC-MS using an authentic sample of **2a** as reference, Figure 5. These results showed that the monooxygenase BcBOT4 catalyzes the first regio- and stereospecific hydroxylation of the probotryane skeleton **2a** at C-4.

The previously reported isolation of intermediates **2b** and acetylated derivatives **2c** and **2d**²³ led us to propose that the acetyl transferase BcBOT5 brought about the acetylation of the hydroxyl group at C-4 of probotryane **2b**.

These results lead us to propose the complete biosynthetic pathway to botrydial (**1**), Figure 2. Thus, direct and indirect evidence for the cyclization of FPP to the parent tricyclic alcohol **2a**, by the sesquiterpene cyclase BcBOT2, has been reported.^{28,31} In addition, the accumulation of intermediate probotryanes **2a**, **2c**, and **2d** confirmed the proposed biosynthetic sequence in which the BcBOT4 monooxygenase produced the hydroxylation at C-4 to give intermediate **2b**. Acetylation of the hydroxyl at C-4 was carried out by the acetyl transferase BcBOT5, followed by the combined action of the P450 monooxygenases BcBOT3 and BcBOT1, to yield finally the glycol **2e**, via the regio- and stereospecific hydroxylations at C-10 and C-15 of the intermediates **2c** and **2d**, respectively.

The cleavage of the C-10–C-15 bond of probotryane skeleton is an intriguing and chemically important reaction, which could be mediated by some of the monooxygenases or by a combination of them. Many P450 monooxygenases carry out the oxidative cleavage of vicinal diols, such as the side chain

cleavage (scc) enzyme in testosterone biosynthesis that cleaves the 20,22-diol.³³ Although the mechanism of this cleavage has never been settled in detail, in this case, it is possible that in the formation of **2e**, either BcBOT3 or BcBOT1 would oxidize either the 10- or the 15-hydroxy group to the hydroperoxide derivative, which would then undergo heterolytic fragmentation to give the dialdehyde botrydial (**1**).

The products isolated from these mutants shed an interesting light on a possible mechanism for the cleavage of the C-10–C-15 bond in botrydial biosynthesis. Earlier studies had shown that this takes place with the formation of botrydial rather than dihydrobotrydial and the use of stereospecifically labeled mevalonates implicated a trans C-10–C-15 glycol.^{21,22,24} The formation of a C-15 ketone (**9**) when BcBOT3 is knocked-out but BcBOT1 remains suggests that the oxidation of the C-15 alcohol to a ketone can be mediated by this enzyme. Since BcBOT1 is a monooxygenase rather than a dehydrogenase, the oxidation may occur by the reaction indicated in Figure 6A, where ketone (**9**) is obtained, via hydroperoxide, from the alcohol intermediate **2d**.

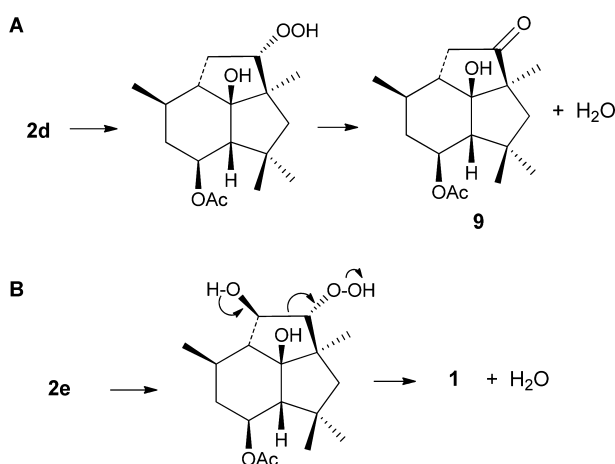


Figure 6. (A) Formation of compound **9** from intermediate **2d**. (B) Formation of botrydial (**1**) from intermediate **2e**.

A corollary to this suggestion is that the cleavage of the C-10–C-15 bond is also mediated by this enzyme. This would involve the fragmentation of a C-15 hydroperoxide derived from **2e** (Figure 6B). This would accord with the labeling pattern from the mevalonates and in particular the trans relationship between the functional groups.²⁴

Here we have reported the genetic and molecular basis of botrydial biosynthesis and the assignment of the biochemical function to the five genes that are colocalized in the genome of

B. cinerea and constitute the botrydial biosynthesis gene cluster. Genes BcBOT3 and BcBOT4, encoding two cytochrome P450 monooxygenases, catalyzed regio- and stereospecific hydroxylations at the carbons C-10 and C-4, respectively, of the presilphiperfolan-8 β -ol skeleton. Interestingly, the *bcbot4* Δ mutant overproduced a significant number of polyketides, which included the four new metabolites **15–18**.

METHODS

Fungal Strains. The strain of *Botrytis cinerea* Pers. Fr. employed in this work, B05.10, was isolated from a *Vitis* field.³⁴

Media and Culture Conditions. *B. cinerea* strains and mutants were cultivated at 23 °C with 12 h daylight per day. Cultures were grown on malt agar medium (MA; 20 g/L malt extract, 5 g/L yeast extract, 15 g/L agar) and minimal medium (MMII; 20 g/L glucose, 2 g/L NaNO₃, 1 g/L KH₂PO₄, 500 mg/L MgSO₄·7H₂O, 500 mg/L KCl, 10 mg/L FeSO₄·7H₂O, 15 g/L agar). The transformation medium was a minimal medium supplemented with saccharose (MMV; 20 g/L glucose, 200 g/L saccharose, 2 g/L NaNO₃, 1 g/L KH₂PO₄, 500 mg/L MgSO₄·7H₂O, 500 mg/L KCl, 10 mg/L FeSO₄·7H₂O, 15 g/L agar). The *bcbot3* and *bcbot4* mutants were selected with nourseothricin fungicide (Werner Bioagent, Germany; 70 μ g/mL).

Standard Molecular Methods: Gene Deletion of the Cytochrome P450-Encoding Genes BcBOT3 and BcBOT4. The *B. cinerea* genes BcBOT3 and BcBOT4 (GenBank AY277723.2) were deleted by homologous recombination. The 5' genomic noncoding regions of BcBOT3 and BcBOT4 (882 bp and 1442 bp, respectively) were amplified by PCR. In the same way, we obtained the 3' flanking sequences of BcBOT3 and BcBOT4 (1027 bp and 866 bp, respectively) and the nourseothricin resistance gene (*nat*). For each gene, a knockout (KO) cassette made of the 5' flanking sequence of *bcbot*, the *nat* resistance gene, and the 3' flanking region of BcBOT was obtained by double-joint PCR fusion³⁵ as previously described for *B. cinerea*.³⁶ All the PCR primers are listed in Table 1.

Protoplasts of the B05.10 strain were transformed as previously reported.³⁷ Transformants were selected and purified on medium agar containing 70 μ g/mL nourseothricin. Gene inactivation was verified by PCR using primers designed outside the KO cassette and primers designed in the resistance gene *nat* (primers verify integration, Table 1). In addition, the absence of the BcBOT3 or BcBOT4 WT alleles in the transformant was also checked by PCR. *B. cinerea* mutants used in this study are described at <http://botbioger.versailles.inra.fr/botmut/>.

Chemical Methods and Analysis of the *bcbot3* Δ and *bcbot4* Δ Metabolite Profile. **General Procedure.** *B. cinerea* WT strain and mutants were grown on malt agar medium³² at 25 °C to produce mycelium plugs (0.8 cm) that were further used to inoculate the medium for metabolite production. All strains were fermented in solid malt agar medium following the reported protocol.³² Extraction of the solid agar malt medium was carried out with an ultrasonic bath, using ethyl acetate (3 \times 0.5 vol). The solvents were dried over Na₂SO₄ and concentrated to dryness.

Purification of the metabolites was carried out by column chromatography (CC) and semipreparative and analytical HPLC.^{28,32} Optical rotation was recorded with a digital polarimeter.

Table 1. PCR Primers Used for BcBOT3 and BcBOT4 Gene Deletion

primer name pair	sequence fwd	sequence rev	usage
BD48/BD51	tccattaagtctgtttggataaagt	GGGAATGCGGCTCTAGTCGGGTTGCTCTTCTCCAT	BcBOT3 homologous 5' region
BD49/BD50	gggcttgaagaattgtattg	cttgagcggcctgcaaatggatgcgaactctgc	BcBOT3 homologous 3' region
BD44/BD46	gcttctgtccttccgctac	GGGAATGCGGCTCTAGTTCACCTTCTCCGATTTCATG	BcBOT4 homologous 5' region
BD43/BD45	TATGACCGGAGAGCTCGAAA	cttgagcggcctgcaacttaagccctatgacaga	BcBOT4 homologous 3' region
K7Nat1 Fwd/Rev	TAGAGCCGATTCCCGATTCGG	TGCAGGCCGCTCAGGGGCAGGG	nourseothricin K7
BD52/BD12	gctacagcggtgtatctc	CTAAGGGCTCGGGAGTTG	BcBOT3 purity
BD52/Nat1Fverif	gctacagcggtgtatctc	GACACCGCCTGTACGAC	BcBOT3 verify integration
BD2/BD13	cgcggaagaagtcttaagga	gctgaattcccaggatctg	BcBOT4 purity
BD47/Nat1Fverif	TCATTCACCTCGACATACTTCC	GACACCGCCTGTACGAC	BcBOT4 verify integration

IR spectra were determined on a FT-IR spectrophotometer and reported as wavenumber (cm^{-1}).¹³

¹H and ¹³C NMR spectra were obtained on Agilent 400 and 500 MHz spectrometers with SiMe₄ as the internal reference at 25 °C.¹³ Mass spectrometry (MS) and high resolution mass spectrometry (HRMS) were performed on Thermoquest Voyager spectrometer and QTOF mass spectrometer, respectively.¹³

bcbot3Δ Metabolite Analysis. Ten days after inoculation, the solid agar malt medium was extracted³² yielding a dense oil from *bcbot3Δ* culture (350 mg) that was separated by column chromatography on silica gel, eluted with mixtures containing increasing percentages of ethyl acetate/hexane (10–100%) and methanol as solvent.

Final purification of selected fractions was carried out by HPLC (hexane (Hex)–ethyl acetate (EtAc) 40:60; flow 1 mL min⁻¹). Three compounds were isolated and characterized: 4β-acetoxy-15α-hydroxyprobotryane (**2d**) (39 mg), 4β-hydroxyprobotryan-15-ona (**9**) (1.1 mg), and botcinin A (**6**) (1.5 mg).

4β-Hydroxyprobotryan-15-one (9). Colorless oil; $[\alpha]_{\text{D}}^{25} = -9.62$ ($c = 0.106$ mg, CHCl₃). IR ν_{max} (film) 3425.38, 2961.33, 1729.85, 1454.75, 913.80, 771.28 cm⁻¹. ¹H NMR (600 MHz, CDCl₃) δ 4.04 (1H, ddd, $J = 11.2, 9.6, 4.0$ Hz, H-4), 2.69 (1H, dd, $J = 18.1, 8.6$ Hz, H-10), 2.63 (1H, dd, $J = 18.1, 3.3$ Hz, H-10'), 2.13 (1H, d (br), H-7), 1.91 (1H, ddd, $J = 12.8, 4.0, 2.9$ Hz, H-3), 1.82 (1H, m, H-2), 1.64 (1H, ddd, $J = 11.6, 8.6, 3.3$ Hz, H-1), 1.46 (1H, d, $J = 9.6$ Hz, H-5), 1.37 (3H, s, H-14), 1.34 (3H, s, H-13), 1.32 (3H, s, H-12), 0.97 (3H, d, $J = 6.4$ Hz, H-11). ¹³C NMR (151 MHz, CDCl₃) δ 211.0 (s, C-15), 94.4 (s, C-9), 70.0 (d, C-4), 63.2 (s, C-8), 60.6 (d, C-5), 48.7 (d, C-1), 47.2 (s, C-6), 44.2 (t, C-7), 44.1 (t, C-3), 42.6 (t, C-10), 36.2 (q, C-12), 33.4 (d, C-2), 27.8 (q, C-13), 23.9 (q, C-14), 21.0 (q, C-11). HRMS (ES⁺) m/z calcd for C₁₅H₂₅O₃ [M + H]⁺ 253.1804, found 253.1807.

Acetylation of 4β-Hydroxyprobotryan-15-one (9). Compound **9** (5 mg) was dissolved in the minimum quantity of dry pyridine (0.5 mL), and acetic anhydride (30 mg) was added to the solution at 25 °C. The mixture was kept at RT for 24 h, and the solvent was evaporated. The crude was chromatographed over silica gel (hexane–ethyl acetate 8:2) and HPLC (Hex–EtAc) 80:20; flow 1 mL min⁻¹; $t_{\text{R}} = 19.5$ min) to give derivative **9a** (4.5 mg).

4β-Acetoxyprobotryan-15-one (9a). Colorless oil; $[\alpha]_{\text{D}}^{25} = -9.40$ ($c = 0.1$ mg, CHCl₃); IR ν_{max} (film) 2350.4, 1727.5, 1248, 776.61, 669.12 470.26 cm⁻¹. ¹H NMR (600 MHz, benzene) δ 5.11 (1H, td, $J = 10.9, 4.0$ Hz), 2.70 (1H, dd, $J = 18.1, 8.6$ Hz, H-10), 2.64 (1H, dd, $J = 18.1, 3.4$ Hz, H-10'), 2.17–2.08 (2H, m), 2.03 (3H, s, CH₃COO–), 2.02 (1H, dt, $J = 7.0, 5.4$ Hz, H-3), 1.95 (1H, m, H-2), 1.37 (3H, d, $J = 0.9$ Hz), 1.29 (3H, s), 1.24 (2H, t, $J = 3.6$ Hz), 1.16 (3H, s), 1.23–1.01 (2H, m), 0.95 (3H, d, $J = 6.4$ Hz, H-11). ¹³C NMR (151 MHz, CHCl₃) δ 210.5 (s, C-15), 170.4 (s, CH₃COO–), 94.0 (s, C-9), 72.2 (d, C-4), 63.1 (s, C-8), 56.2 (d, C-5), 48.6 (d, C-1), 47.15 (s, C-6), 44.1 (t, C-7), 42.7 (t, C-10), 39.6 (t, C-3), 35.8 (q, C-12), 33.2 (d, C-2), 27.5 (q, C-13), 24.1 (q, C-14), 21.4 (CH₃COO–), 20.9 (d, C-11). HRMS (ES⁻) m/z calcd for C₁₇H₂₅O₄ [M – H]⁻ 293.1753, found 293.1779.

bcbot4Δ Metabolite Analysis. The analysis was carried out as described for *bcbot3Δ*. Evaporation of the solvent under reduced pressure afforded a dense oil from the *bcbot4Δ* culture (360 mg) that was chromatographed by CC on silica gel. Final purification by HPLC and GC-MS yielded the metabolite probotryane **2a** and a significant number (15) of metabolites with polyketide skeletons. In addition to the known polyketides the following were isolated and characterized: botrylactone (**3**, 21.7 mg), botcinic and botcinic acids (**4** and **5**, 6.9 and 4.5 mg), botcinins A, B, E, and G (**6**, **7**, **12**, and **13**, 63.6, 37.6, 10.1, and 7.9 mg), cinbotolide (**8**, 12 mg), acetylbotcinic and acetylbotcinic acids (**10** and **11**, 8.3 and 7.0 mg), and 5-hydroxy-7-(4-hydroxydec-2(3)-enoyl)botrylactone (**14**, 12.2 mg); the new botrylactones, 5-hydroxy-7-(oct-2(3)-enoyl) botrylactone (**15**, 11.5 mg; semipreparative (Semprep) HPLC, Hex–EtAc–acetone (Act) 70:20:10; flow 2.5 mL min⁻¹; $t_{\text{R}} = 27$ min) and 6-hydroxy-7-deoxybotrylactone (**16**, 15 mg, semprep HPLC, Hex–EtAc–Act 70:20:10; flow 2.5 mL min⁻¹; $t_{\text{R}} = 38.5$ min), and the new cinbotolides B (**17**, 2.4 mg, semprep HPLC, Hex–EtAc–Act 50:40:10; flow 2.5 mL

min⁻¹; $t_{\text{R}} = 40$ min) and C (**18**, 11.8 mg, semprep HPLC, Hex–EtAc 75:25; flow 3 mL min⁻¹; $t_{\text{R}} = 45$ min).

Presilphiperfolan-8β-ol (Probotryane) (2a). The less polar fraction obtained from the *bcbot4Δ* extract with hexane was studied by GC-MS. Fermentations, for 4, 8, and 10 days, of wild-type B05.10 and of the mutant *bcbot4Δ* were carried out, and the less polar fractions obtained by extraction with hexane was studied by GC-MS. An authentic sample of the presilphiperfolan-8β-ol (**2a**) was used as reference to detect the compound in the extracts. Compound **2a** was not detected in the hexane fractions from the wild-type at 4, 8, and 10 days. However, the presence of compound **2a** was clearly revealed in all fractions from the fermentations of the mutant *bcbot4Δ* on different days (see Figure S).

2R,3S,4S,6S,8R,9R-5-Hydroxy-7-(oct-2(3)-enoyl) botrylactone (15). Yellow oil; $[\alpha]_{\text{D}}^{25} 4.7$ ($c = 0.1$ mg, CHCl₃); IR ν_{max} (film) 3348, 2929.48, 1721.36, 1456.89, 1260.51, 1105.94, 1015.96, 719.49 cm⁻¹. ¹H NMR (400 MHz, CDCl₃) δ 7.04 (1H, dt, $J = 15.6, 6.9$ Hz, H-3'), 5.84 (1H, dt, $J = 15.6, 1.6$ Hz, H-2'), 4.93 (1H, d, $J = 10.9$ Hz, H-7), 3.68 (1H, d(br), $J = 11.1$ Hz, H-5), 3.52 (1H, d, $J = 0.5$ Hz, H-3), 2.76 (1H, qd, $J = 7.4, 0.5$ Hz, H-2), 2.21 (2H, m, H-4'), 1.97 (1H, m, H-6), 1.55 (3H, s, C₉–CH₃), 1.47 (3H, d, $J = 7.4$ Hz, C₂–CH₃), 1.38–1.24 (m, H-5', H-6', H-7'), 1.15 (3H, s, C₄–CH₃), 1.10 (3H, s, C₈–CH₃), 1.04 (3H, d, $J = 6.5$ Hz, C₆–CH₃), 0.90 (3H, t, $J = 6.9$ Hz, H-8'). ¹³C NMR (100 MHz, CDCl₃) δ 171.0 (s, C-1), 165.5 (s, C-1'), 151.4 (d, C-3'), 120.2 (d, C-2'), 104.2 (s, C-9), 80.8 (d, C-3), 79.3 (d, C-8), 76.2 (s, C-4), 75.2 (d, C-5), 73.8 (d, C-7), 36.5 (d, C-6), 34.5 (d, C-2), 32.3 (t, C-4'), 31.3 (t, C-5'), 27.6 (t, C-6'), 22.4 (t, C-7'), 21.7 (q, C₉–CH₃), 18.5 (q, C₂–CH₃), 18.2 (q, C₄–CH₃), 18.1 (q, C₈–CH₃), 13.9 (q, C-8'), 13.8 (q, C₆–CH₃). HRMS (ES⁺) m/z calcd for C₂₂H₃₅O₇ [M + H]⁺ 411.2383, found 411.2370.

6-Hydroxy-7-deoxybotrylactone (16). White amorphous powder; $[\alpha]_{\text{D}}^{25} 1.16$ ($c = 1.146$ mg, CHCl₃); IR ν_{max} (film) 3447.87, 2938.54, 1740.30, 1258.90, 1065.66, 976.23, 691.56 cm⁻¹; ¹H NMR (500 MHz, CDCl₃) δ 3.33 (1H, d, $J = 0.7$ Hz, H-3), 3.27 (1H, s(br), –OH), 2.74 (1H, qd, $J = 7.3, 0.7$ Hz, H-2), 1.91–1.69 (4H, m, H-5, H-7), 1.44 (3H, d, $J = 7.3$ Hz, C₂–CH₃), 1.44 (3H, s, C₉–CH₃), 1.30 (3H, s, C₈–CH₃), 1.27 (3H, s, C₄–CH₃), 1.24 (3H, s, C₆–CH₃). ¹³C NMR (126 MHz, CDCl₃) δ 171.1 (s, C-1), 104.4 (s, C-9), 81.6 (d, C-3), 76.5 (s, C-8), 72.8 (s, C-4), 70.3 (s, C-6), 45.0 (t, C-5), 43.1 (t, C-7), 34.4 (d, C-2), 28.2 (q, C₆–CH₃), 27.5 (q, C₄–CH₃), 26.4 (q, C₈–CH₃), 22.06 (C₉–CH₃), 18.6 (q, C₂–CH₃). HRMS (ES⁻) m/z calcd for C₁₄H₂₁O₅ [M – H]⁻ 269.1389, found 269.1385.

Cinbotolide B (17). Yellow oil; $[\alpha]_{\text{D}}^{25} -16.03$ ($c = 0.131$ mg, CHCl₃); IR ν_{max} (film) 3417.83, 2925.61, 1747.89, 1455.50, 1383.58, 1217.32, 996.51, 934.66, 748.70 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 5.39 (1H, q, $J = 6.9$ Hz, H-9), 4.15 (2H, dd, $J = 12.0, 6.9$ Hz, H-10), 4.05 (1H, d, $J = 5.5$ Hz, H-3), 2.94 (1H, dq, $J = 7.3, 5.5$ Hz, H-2), 2.26 (1H, dd(br), $J = 13.1, 6.4$ Hz, H-5), 2.02–1.75 (3H, m, H-5', H-6, H-7), 1.68 (3H, d, $J = 6.9$ Hz, C₈–CH₃), 1.71–1.6 (1H, m, H-7'), 1.36 (3H, s, C₄–CH₃), 1.24 (3H, d, $J = 7.3$ Hz, C₂–CH₃), 0.95 (3H, d, $J = 6.2$ Hz, C₆–CH₃). ¹³C NMR (100 MHz, CDCl₃) δ 177.6 (s, C-1), 137.3 (s, C-8), 125.2 (d, C-9), 87.8 (s, C-4), 76.5 (d, C-3), 60.0 (t, C-10), 44.5 (t, C-7), 40.7 (t, C-5), 40.0 (d, C-2), 27.2 (d, C-6), 23.2 (q, C₄–CH₃), 21.4 (q, C₆–CH₃), 13.2 (q, C₈–CH₃), 8.3 (q, C₂–CH₃). HRMS (ES⁺) m/z calcd for C₁₄H₂₅O₄ [M + H]⁺ 257.1747, found 257.1758.

Cinbotolide C (18). Colorless oil; $[\alpha]_{\text{D}}^{25} 52.5$ ($c = 0.12$ mg, CHCl₃); IR ν_{max} (film) 3426.45, 2926.73, 1752.33, 1643.35, 1456.46, 1046.65, 771.09 cm⁻¹. ¹H NMR (400 MHz, CDCl₃) δ 4.18 (1H, d, $J = 5.5$ Hz, H-3), 4.01 (2H, d, $J = 1.3$ Hz, H-12), 3.65 (1H, q, $J = 6.5$ Hz, H-9), 2.86 (1H, dq, $J = 7.3, 5.5$ Hz, H-2), 2.26 (1H, m, H-6), 1.78 (1H, ddd, $J = 14.5, 2.8, 1.6$ Hz, H-5), 1.66 (1H, dd, $J = 14.2, 11.4$ Hz, H-7'), 1.57 (1H, dd, $J = 14.5, 10.9$ Hz, H-5'), 1.47 (1H, dt, $J = 14.2, 1.7$ Hz, H-7), 1.26 (3H, d, $J = 7.3$ Hz, C₂–CH₃), 1.21 (3H, s, C₈–CH₃), 1.14 (3H, d, $J = 6.5$ Hz, C₉–CH₃), 0.98 (3H, d, $J = 6.7$ Hz, C₆–CH₃). ¹³C NMR (100 MHz, CDCl₃) δ 177.3 (s, C-1), 87.7 (s, C-8), 79.8 (s, C-4), 76.6 (d, C-3), 74.4 (d, C-9), 63.1 (t, C-12), 45.7 (t, C-5), 42.7 (t, C-7), 39.7 (d, C-2), 24.6 (q, C₆–CH₃), 24.6 (d, C-6), 18.9 (q, C₈–CH₃), 17.3 (q, C₁₀–CH₃), 8.3 (q, C₂–CH₃). HRMS (ES⁺) m/z calcd for C₁₄H₂₅O₅ [M + H]⁺ 273.1702, found 273.1701.

■ ASSOCIATED CONTENT

■ Supporting Information

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Details of spectroscopic and characterization data, and reproductions of ¹H- and ¹³C NMR spectra (PDF)

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Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

This research was supported by a grant from Ministerio de Economía y Competitividad (AGL2015-65684-C2-1-R) (MINECO-FEDER). Use of NMR facilities at the Servicio Centralizado de Ciencia y Tecnología (SCCYT) of the University of Cádiz is acknowledged. We thank D. Cane for the generous gift of a sample of compound **2a**.

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