

Induced Daldinin A, B, C with a New Skeleton from Cultures of the Ascomycete *Daldinia concentrica*

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Abstract Daldinin A, B, C with a new skeleton, together with four known compounds, were induced and isolated from cultures of the ascomycete *Daldinia concentrica*. Their structures were elucidated by spectroscopic analysis, and that of daldinin A was confirmed by single-crystal X-ray diffraction.

Keywords daldinin A, daldinin B, daldinin C, *Daldinia concentrica*, ascomycete, induction

Introduction

Natural products are an attractive source of varied structures that exhibit potent biological activities, and desirable pharmacological profiles. Historically, natural products have been the source of many new drugs. Their importance is well documented in the recent literatures [1–3]. The search for bioactive compounds is one of the central subjects of industrial and academic natural products chemistry. With new emerging targets for drug discovery, continued demand for new diverse natural products serve as a vehicle for probing biological activity.

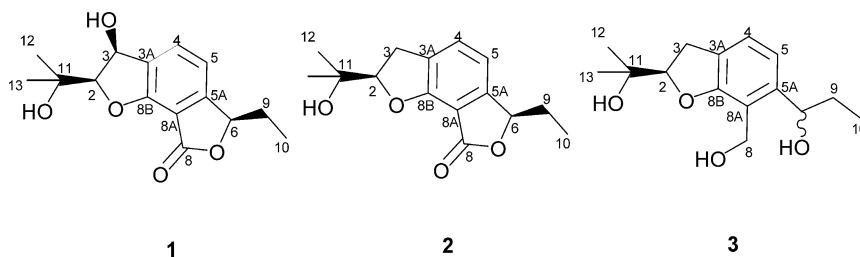
Variation of cultivation parameters to induce the production of formerly unknown compounds is the most simple approach to increase the number of secondary metabolites from one single organism for fungi or bacteria. This way of releasing nature's chemical diversity was

termed the 'OSMAC (One Strain-Many Compounds)' approach [4, 5]. It is based on the observation that individual strains are able to produce more metabolites than normally detected in a routine screening program and very small changes in the cultivation parameters (for example, culture vessel, media composition, addition of enzyme inhibitors) can completely shift the metabolic profile of various microorganisms. It is a very simple but even more random approach to the improvement of fermentations to obtain maximum production titers of desired compounds. This approach was successfully used as a valuable tool to exploit natural products diversity in the past, for example, actinomycetes (*Streptomyces* sp.) and fungi (*Aspergillus* sp., *Sphaeropsidales* sp.) produce additional compounds after variation of the culture conditions [6–8].

As shown in former investigations, an anti-HIV-1 agent concentricolide (4) and other six new compounds were isolated from the fruiting bodies of *Daldinia concentrica* [9–13]. Concentricolide showed an activity to inhibit HIV-1 induced cytopathic effects. The EC₅₀ value was 0.31 μg/ml. The therapeutic index (TI) was 247. Concentricolide exhibited the blockage (EC₅₀ 0.83 μg/ml) on syncytium formation between HIV-1 infected cells and normal cells. *Daldinia concentrica* (strain S 0318) produces concentricolide and a pair of novel hept-6-ene-2,4,5-triols when cultivated in reagent bottles (color: brown; size: 500 ml; volume of medium: 300 ml) with cultivation medium (PD): potato (peel off) 200 g, glucose 20 g, KH₂PO₄ 3.0 g, MgSO₄ 1.5 g, citric acid 0.1 g and thiamin hydrochloride 10 mg in 1.0 liter of de-ionized water. Applying the OSMAC approach, we intended to examine whether the production of new metabolites could be induced in this strain.

Following the OSMAC approach, our strain S 0318 has been cultivated with Erlenmeyer flask instead of reagent

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Structures of daldinin A (**1**), daldinin B (**2**), daldinin C (**3**)

bottles. The surprises were obtained. In the culture broth extract from Erlenmeyer flasks (color: white; size: 500 ml; media: 300 ml) we detected, by thin layer chromatography and HPLC, rich additional metabolites besides concentricolide. Careful investigation on the culture of *D. concentrica* strain (S 03180) led to the isolation of new compounds daldinins A~C (**1**~**3**) and known compounds: concentricolide (**4**) [9], isoochracein (**5**) [14], 4-hydroxyisoochracein (**6**) [15] and 2-(hydroxymethyl)-3-(1'-hydroxypropyl) phenol (**7**) [16] by the alteration of culture parameter (form of culture vessel). The fermentation, isolation and structure elucidation of these compounds are reported here.

Materials and Methods

General

NMR experiments were performed on a Bruker AM-400 or a DRX-500 spectrometer with TMS as internal standard. Mass spectra were recorded on a VG Auto Spec-3000 or an API QSTAR Pulsar 1 spectrometer. IR spectra were recorded on a Bruker Tensor 27 spectrometer with KBr pellets. Optical rotations were measured on a Horiba SEPA-300 polarimeter. Column chromatography was performed on silica gel (200~300 mesh; Qingdao Marine Chemical Ltd., Qingdao, People's Republic of China) and Sephadex LH-20 (Amersham Biosciences, Sweden). TLC analysis was carried out on silica gel GF254 precoated plates (0.20~0.25 mm; Qingdao) with detection by heating silica gel plates sprayed with 10% H₂SO₄ in EtOH.

Fungal Material

Strain (S 0318) was isolated from tissue culture of the fruiting bodies of *Daldinia concentrica* collected at Laojunshan, Yunnan Province, P. R. China, in July 2003 and identified by Prof. Mu Zang, Kunming Institute of Botany, Chinese Academy of Sciences. A voucher specimen (HKAS 40992) was deposited at the herbarium of Kunming Institute of Botany, Chinese Academy of Sciences.

Cultivation

Strain S 0318 was grown on agar slopes at 22°C until sporulation, and then stored at 4°C. The culture medium consisted of potato (peel off) 200 g, glucose 20 g, KH₂PO₄ 3.0 g, MgSO₄ 1.5 g, citric acid 0.1 g and thiamin hydrochloride 10 mg in 1.0 liter of de-ionized water. The fungus was grown in Erlenmeyer flasks (color: white; size: 500 ml; media: 300 ml). The pH was adjusted to 6.5 before autoclaving. Fermentation was carried out on a rotary shaker at 22°C and 150 rpm for 10 days.

Extraction and Isolation

The mycelium from 60 liters was filtered, and the filtrate was extracted five times with EtOAc (total 80 liters). The EtOAc extract was evaporated *in vacuo* and the deep brown gum (80 g) was subjected to column chromatography (silica gel) using a CHCl₃/MeOH gradient elution. Elution with CHCl₃/MeOH (98:2, v/v) produced **4** (20 mg), and **5** (17 mg). The fraction (20 g) from CHCl₃/MeOH (95:5) was further purified by twice chromatography over silica gel (petroleum ether/Me₂CO, from 20:1 to 10:1; CHCl₃/MeOH, from 30:1 to 10:1), and repeated Sephadex LH-20 (CHCl₃/MeOH, 1:1) to afford the pure compounds **2** (63 mg), and **6** (46 mg). The fraction (24 g) eluted with CHCl₃/MeOH (90:10) at the first chromatography was also further purified by repeated chromatography over silica gel (CHCl₃/MeOH, from 20:1 to 5:1) to yield compounds **1** (7 mg), **3** (60 mg), and **7** (87 mg).

Physico-chemical Properties

Daldinin A (**1**). Colorless prisms; m.p. 228~229 (C); [α]_D²⁶ -44.2° (*c* 0.39 in MeOH); IR (KBr): ν_{\max} 3258, 1760, 1633, 1607, 1475, 1452 cm⁻¹; UV (MeOH): λ_{\max} (log ϵ) 209 (4.48), 217 (4.43), 306 (3.83), 359 (2.56) nm; ¹H- and ¹³C-NMR data: see Table 1; EI-MS *m/z* (relative intensity) [M]⁺ 278 (1), 203 (94), 185 (41), 173 (100), 145 (83); Negative FAB-MS *m/z* (relative intensity) [M-H]⁻ 277(100); HR-ESI-MS *m/z* [M+Na]⁺ 301.1049 (calcd for C₁₅H₁₈O₅Na, 301.1051).

Table 1 NMR spectral data for daldinin A (**1**) in CD₃OD

	¹³ C	¹ H	¹ H, ¹ H-COSY	HMBC
2	94.0	4.46 (1H, d, 6.3)	H-3	C-3A, 8B, 12, 13
3	71.9	5.37 (1H, d, 6.3)	H-2	C-4, 8B, 11
3A	133.4			
4	133.5	7.70 (1H, d, 7.6)	H-5	C-3, 5A, 8B
5	115.2	7.06 (1H, d, 7.6)	H-4	C-3A, 6, 8A
5A	154.2			
6	84.3	5.51 (1H, dd, 4.1, 6.8)	H-9	C-5, 5A, 8, 8A, 10
8	170.4			
8A	110.2			
8B	158.7			
9	28.7	1.81 (1H, m) 2.14 (1H, m)	H-6, 10	C-5A, 6, 10
10	8.8	0.95 (3H, t, 7.4)	H-9	C-6, 9
11	73.2			
12	26.5	1.50 (3H, s)		C-2, 11, 13
13	27.0	1.48 (3H, s)		C-2, 11, 12

Table 2 NMR spectral data for daldinin B (**2**) in CD₃OD

	¹³ C	¹ H	¹ H, ¹ H-COSY	HMBC
2	92.6	4.48 (1H, dd, 8.8, 9.3)	H-3	C-3A, 8B, 12, 13
3	29.8	3.17 (1H, dd, 9.7, 16.1) 3.25 (1H, dd, 8.8, 16.1)	H-2	C-4, 8B, 11
3A	130.3			
4	131.3	7.44 (1H, d, 7.4)	H-5	C-3, 5A, 8B
5	113.6	6.89 (1H, d, 7.4)	H-4	C-3A, 6, 8A
5A	150.8			
6	82.4	5.34 (1H, dd, 4.4, 6.8)	H-9	C-5, 5A, 8, 8A, 10
8	168.2			
8A	108.8			
8B	158.4			
9	29.3	1.67 (1H, m), 2.01 (1H, m)	H-6, 10	C-5A, 6, 10
10	9.0	0.88 (3H, t, 7.3)	H-9	C-6, 9
11	71.3			
12	25.6	1.26 (3H, s)		C-2, 11, 13
13	25.5	1.19 (3H, s)		C-2, 11, 12

Crystallographic Data for 1

C₁₅H₁₈O₅, *M* 278.30, orthorhombic, space group *P*2₁2₁2₁, *a*=8.245 (1) Å, *b*=11.153 (1) Å, *c*=15.784 (1) Å, *V*=1451.4 (2) Å³, *Z*=4, crystal dimensions 0.20×0.40×0.60 mm were used for measurements on a MAC DIP-2030K diffractometer with a graphite monochromator ($\omega-2\theta$ scans, $2\theta_{\max}=50.0^\circ$), Mo K α radiation. The total number of independent reflections measured was 1517, of which 1407 were observed ($|F|^2 \geq 2\sigma|F|^2$). Final indices: *R*₁=0.058, *wR*₂=0.165. The crystal structure of **1** was

solved by direct methods using SHELXS-97 and expanded using difference Fourier techniques, refined by the program and method NOMCSDP and full-matrix least-squares calculations. The crystal structure of **1** was solved by direct methods using SHELXS-97 and expanded using difference Fourier techniques, refined by the program and method NOMCSDP and full-matrix least-squares calculations.

Daldinin B (**2**). Light yellow, oily solid; $[\alpha]_D^{19} -26.6^\circ$ (*c* 0.60 in Me₂CO); IR (KBr): ν_{\max} 3445, 1755, 1633, 1607, 1480, 1450 cm⁻¹; UV (MeOH): λ_{\max} (log ϵ) 212 (4.21), 218

Table 3 NMR spectral data for daldinin C (**3**) in CD₃OD

	¹³ C	¹ H	¹ H, ¹ H-COSY	HMBC
2	90.2	4.58 (1H, dd, 8.5, 9.4)	H-3	C-3A, 8B, 12, 13
3	31.4	3.10 (1H, dd, 9.6, 16.0) 3.25 (1H, dd, 8.5, 16.0)	H-2	C-4, 8B, 11
3A	127.0			
4	125.3	7.09 (1H, d, 7.6)	H-5	C-3, 5A, 8B
5	119.2	6.95 (1H, d, 7.6)	H-4	C-3A, 6, 8A
5A	144.4			
6	72.7	4.88 (1H, dd, 6.3, 7.2)	H-9	C-5, 5A, 8A, 10
8	58.7	4.71 (2H, s)		
8A	120.0			
8B	159.8			
9	32.4	1.77 (2H, m)	H-6, 10	C-5A, 6, 10
10	11.0	0.94 (3H, t, 7.4)	H-9	C-6, 9
11	72.6			
12	24.8	1.20 (3H, s)		C-2, 11, 13
13	25.7	1.30 (3H, s)		C-2, 11, 12

(4.44), 314 (3.79) nm; ¹H- and ¹³C-NMR data: see Table 2; EI-MS *m/z* (relative intensity) [M]⁺ 262 (24), [M-CH₃]⁺ 247 (8), [M-CH₃-H₂O]⁺ 229 (31), 215 (10), 204 (72), 186 (100), 175 (55), 171 (60), 158 (88), 145 (30), 129 (14); HR-ESI-MS *m/z* [M+Na]⁺ 285.1099 (calcd for C₁₅H₁₈O₄Na, 285.1102).

Daldinin C (**3**). Oily solid; [α]_D²⁵ -2.2° (*c* 0.46 in MeOH); IR (KBr): ν_{max} 3384, 1712, 1623, 1595, 1443 cm⁻¹; UV (MeOH): λ_{max} (log ε) 206 (4.53), 292 (3.59) nm; ¹H- and ¹³C-NMR data: see Table 3; EI-MS *m/z* (rel intensity) [M]⁺ 266 (7), 248 (1), 233 (2), 219 (97), 201 (19), 190 (22), 172 (75), 157 (100), 133 (46), 105 (11), 91 (13), 77 (14); Negative FAB-MS *m/z* (relative intensity) [M-H]⁻ 265 (100); HR-ESI-MS *m/z* [M-H]⁻ 265.1439 (calcd for C₁₅H₂₁O₄, 265.1439).

Results and Discussion

Daldinin A (**1**) was obtained as colorless prisms. Its molecular formula was found to be C₁₅H₁₈O₅ by mass spectrometry and confirmed by HR-ESI-MS (found 301.1049, calcd for C₁₅H₁₈O₅Na, 301.1051). The IR spectrum showed absorptions at 1760 (γ-lactone), 1633, 1607, 1475, and 1452 (aromatic ring) cm⁻¹. Fifteen signals in the ¹³C-NMR (DEPT) spectra were recognized including one ester carbonyl *C*-atom (δ 170.4), four aromatic quaternary *C*-atoms (δ 158.7, 154.2, 133.4 and 110.2) and two aromatic methines (δ 133.5 and 115.2). The remaining signals in the ¹³C-NMR spectra could be attributed to three

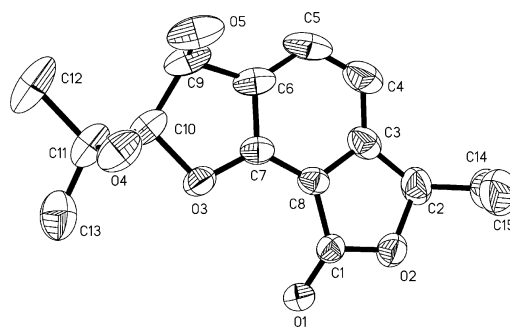


Fig. 1 Crystal structure of daldinin A (**1**) with crystallographic numbering pattern.

methyls (δ 8.8, 26.5 and 27.0), one methylene (δ 28.7), three methines bearing an *O*-atom (δ 94.0, 71.9 and 84.3), and one quaternary carbon bearing an *O*-atom (δ 73.2). The ¹H-¹H COSY data allowed establishment of three H-atoms systems, one at C(2) through C(3), other two at C(4) through C(5) and C(6) through C(10), by showing correlations between H-2 and H-3; H-4 and H-5; H-6 and H-9, H-9 and H-10. Combining with the information of HMBC, the basic skeleton of **1** was established. The structure of **1** was assigned by X-ray crystallographic analysis, which also offered information on the relative stereochemistry of **1** (Fig. 1). Thus, the structure of **1** was determined.

Daldinin B (**2**) was obtained as oily solid. Its molecular formula of C₁₅H₁₈O₄ was established on the basis of EI-MS, ¹³C-NMR, and DEPT NMR spectra and confirmed by HR-ESI-MS (found 285.1099, calcd for C₁₅H₁₈O₄Na,

285.1102). The IR spectrum showed absorptions at 1755 (γ -lactone), 1633, 1607, 1480, and 1450 (aromatic ring) cm^{-1} . Fifteen signals in the ^{13}C -NMR (DEPT) spectra were recognized including one ester carbonyl C-atom (δ 168.2), four aromatic quarternary C-atoms (δ 158.4, 150.8, 131.3 and 108.8) and two aromatic methines (δ 130.3 and 113.6). The remaining signals in the ^{13}C -NMR spectra could be attributed to three methyls (δ 9.0, 25.6 and 25.5), two methylenes (δ 29.8, and 29.3), two methines bearing an O-atom (δ 92.6, and 82.4), and one quarternary carbon bearing an O-atom (δ 71.3). The ^1H - ^1H COSY data allowed establishment of three H-atoms systems, one at C-2 through C-3, other two at C-4 through C-5 and C-6 through C-10, by showing correlations between H-2 and H-3; H-4 and H-5; H-6 and H-9, H-9 and H-10. Combining with the information of HMBC, the basic skeleton of **2** was established, which is similar to **1**. The structure of **1** was assigned by X-ray crystallographic analysis, which also offered information on the relative stereochemistry of **2**.

Daldinin C (**3**) was obtained as oily solid. The HR-EI-MS together with the NMR data were used to determine the molecular formula of **3** to be $\text{C}_{15}\text{H}_{22}\text{O}_4$, possessing four more hydrogen atoms than **2**. Also, the ^1H - and ^{13}C -NMR signals of **3** were virtually identical to those of **2**. The absence of a signal for a carbonyl (δ_{C} 168.2, s, C-8) of **2**, along with the observation of resonances for an additional oxygenated methyl (δ_{H} 4.71, 2H, s, H-8, δ_{C} 58.7, t, C-8), suggested the lack of a lactone ring in **3**, a feature found in **2**. Therefore, the structure of **3** was suggested.

The anti-HIV activity of daldinins A~C was tested with reported method [9], the results were negative.

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