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Rhizovagine A, an unusual dibenzo- α -pyrone alkaloid from the endophytic fungus *Rhizopycnis vagum* Nitaf22†

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Rhizovagine A (**1**), a novel dibenzo- α -pyrone alkaloid with an unprecedented 5/5/6/6/6 fused pentacyclic skeleton, was isolated from the endophytic fungus *Rhizopycnis vagum* Nitaf22. The structure was elucidated by comprehensive spectroscopic analysis, in combination with quantum chemical ^{13}C NMR and electronic circular dichroism (ECD) calculations for configurational assignment. A plausible biosynthetic pathway for **1** was proposed. Compound **1** displayed acetylcholinesterase inhibitory activity.

Introduction

Dibenzo- α -pyrones, also known as 6*H*-dibenzo[*b,d*]pyran-6-ones, are a class of polyketides mainly produced by fungi.¹ Structurally, the diversity of this type of metabolites mainly arises from ring A by having different substitutions and different degrees of unsaturation, while ring B is exclusively aromatic though it might have different substituents at C-7 and C-9 (*i.e.* OMe, OH) (Fig. 1). In some unusual cases, ring A is contracted to a five-membered ring, as found in cephalosol,² or spiro-fused to a γ -lactone ring at C-4, as in rhizopycnolides A and B,³ and penicilliumolide A,⁴ or fused to a tetrahydrofuran, as in xinshenglin,⁵ or attached to a C₃ unit to form a diepoxy-cage structure, as in alternarilactone A.⁶ Ring C is six-membered in all reported structures except graphislactone D that contains a seven-membered ring.⁷ These metabolites display diverse biological activities, such as antimicrobial,^{2,3} cytotoxic,^{3,4} antioxidant,⁸ and larvicidal⁹ activities. The highly structural novelty as well as the interesting bioactivities made them attractive to the synthetic chemists. For example, the total synthesis of cephalosol, and graphislactones A–D has been completed.^{10–12}

Rhizopycnis vagum Nitaf22 is a fungal endophyte that was isolated from *Nicotiana tabacum*, which was found to produce dibenzo- α -pyrones in our previous research,³ as well as anisic acid derivatives.¹³ In our course to searching for new bioactive fungal metabolites, this fungal strain was re-fermented in the rice medium in a large scale, which led to the isolation of a novel alkaloid, rhizovagine A (**1**) (Fig. 1). Herein, we report the

isolation, structure elucidation, and the bioactivity of **1**. The biosynthetic pathway of **1** was also discussed.

Results and discussion

The fungal EtOAc extract was successively subjected to repeated column chromatography over silica gel, and ODS as well as semi-preparative HPLC to afford compound **1** (Fig. 1).

Rhizovagine A (**1**) was isolated as a minor, yellow amorphous solid. It exhibited a prominent pseudomolecular ion peak at *m/z* 386.1221 in the high-resolution electrospray ionization mass spectrometry (HRESIMS), indicating its molecular formula of C₂₀H₁₉NO₇, with 12 degrees of unsaturation. The UV spectrum displayed maximum absorptions at 203, 249, 340, and 400 nm, which were similar to the reported dibenzo- α -pyrones.³ The ^{13}C NMR spectrum (Table 1) displayed a total of 20 resonances, which were assigned to twelve sp² carbons (including two sp² CH at δ_{C} 101.7, 104.7), one oxygenated quaternary sp³ carbon (δ_{C} 86.2), one methine (δ_{C} 97.3), three methylene (δ_{C} 30.7, 25.4, 48.8), two methoxyl (δ_{C} 60.9, 55.9), and one methyl group (δ_{C}

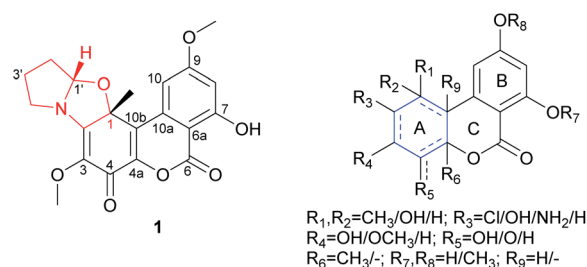


Fig. 1 Structure of rhizovagine A (**1**), and the basic skeleton of dibenzo- α -pyrones (with possible substitutions and unsaturation pattern indicated).

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† Electronic supplementary information (ESI) available: ECD and ^{13}C NMR calculation data, 1D and 2D NMR, and MS spectra of **1**. See DOI: 10.1039/d0ra05022a



Table 1 ^1H and ^{13}C NMR data of **1** (CDCl_3)

Position	δ_{C} , type	δ_{H} , mult. (J in Hz)
1	86.2, C	
2	160.3, C	
3	128.7, C	
4	171.9, C	
4a	143.7, C	
6	164.4, C	
6a	100.9, C	
7	164.7, C	
8	101.7, CH	6.62, d (2.4)
9	166.6, C	
10	104.7, CH	7.20, d (2.4)
10a	134.2, C	
10b	121.7, C	
1-CH ₃	26.2, CH ₃	1.86, s
3-OCH ₃	60.9, CH ₃	3.83, s
7-OH		11.41, s
9-OCH ₃	55.9, CH ₃	3.91, s
1'	97.3, CH	5.65, t (5.6)
2'	30.7, CH ₂	2.23, m
		1.81, m
3'	25.4, CH ₂	2.11, m
4'	48.8, CH ₂	3.61, m

26.2) by the aid of HSQC spectrum. Two *meta*-coupled aromatic protons (δ_{H} 7.20, 6.62, each d, $J = 2.4$ Hz) were seen in the downfield region of the ^1H NMR spectrum (Table 1), which were characteristic of the 7,9-dioxygenated substitutions in B ring, commonly found in dibenzo- α -pyrones.³ One oxymethine proton (δ_{H} 5.65, t), and six methylene protons (δ_{H} 1.74–3.70) were observed in the upfield region, together with two methoxyl groups (δ_{H} 3.91, 3.83) and one singlet methyl group (δ_{H} 1.86). In addition, one chelated hydroxyl group (δ_{H} 11.41) was present.

The gross structure of **1** was established by extensive analysis of the 2D NMR spectra (Fig. 2). The B ring was easily constructed by analysis of the HMBC spectra. The correlations from the chelated OH (δ_{H} 11.41) to C-7 (δ_{C} 164.7), C-6a (δ_{C} 100.9), and C-8 (δ_{C} 101.7), and from the methoxyl group (δ_{H} 3.91) to C-9 (δ_{C} 166.6), allowed the assignment of a 7-hydroxyl-9-methoxyl substituted B ring. The correlations from the methyl group (δ_{H} 1.86, s) to C-1 (δ_{C} 86.2), C-2 (δ_{C} 160.3), and C-10b (δ_{C} 121.7), together with the correlation from H-10 (δ_{H} 7.20, d) to C-10b, clearly positioned the methyl group to C-1. The second methoxyl group (δ_{H} 3.83) was attached to C-3 (δ_{C} 128.7) as

inferred from the cross-peak between them in the HMBC spectrum. Two sp^2 carbons (δ_{C} 171.9, 143.7) remained to be assigned, among which the former was a carbonyl group, while the latter had to be part of a double bond. By comparing the chemical shifts of these carbons with those of the co-occurring rhizopycnin B,³ suggested that this carbonyl was located at C-4, while the other positioned at C-4a (δ_{C} 143.7) to complete the A and C rings.

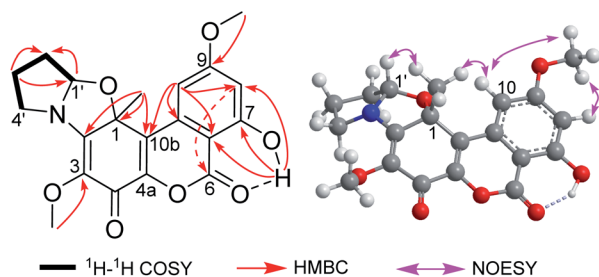
In addition, a 2-oxygenated pyrrolidine ring was recognized by analysis of the ^1H - ^1H COSY spectrum, in which correlations from H-1' (δ_{H} 5.65, t) to H₂-2' (δ_{H} 2.23, 1.81, m), H₂-2' to H₂-3' (δ_{H} 2.11, m), and in turn H₂-3' to H₂-4' (δ_{H} 3.61, m) were seen (Fig. 2), as well as by taken into consideration of the chemical shifts of these hydrocarbon groups, especially CH₂-4' (δ_{H} 3.61/ δ_{C} 48.8) and CH-1' (δ_{H} 5.65/ δ_{C} 97.3).¹⁴ The above functionalities only accounted for 11 degrees of unsaturation, thus hinting the presence of one additional ring to satisfy the required unsaturation index. Then, C-1 and C-1' had to be connected *via* an epoxy bridge, while C-2 was linked to the nitrogen atom to form a five-membered ring. This can explain the unusual downfield shift for C-1 (δ_{C} 86.2), comparing to that of rhizopycnin B (δ_{C} 69.5).³ Moreover, the NOESY correlation between H-1' and 1-CH₃ not only confirmed the above fusion pattern, but also defined the relative configuration of **1** (Fig. 2).

Meanwhile, in order to confirm the deduced structure, a GIAO ^{13}C NMR calculation was performed on mPW1PW91/6-311+G(2d,p)//B3LYP/6-31+G(d,p) level of theory using the reported procedure and scaling parameters.¹⁵ As shown in Table 2, the calculated ^{13}C NMR data of the proposed structure fitted

Table 2 GIAO ^{13}C NMR calculation of **1**

Position	Exp. δ_{C}	Cal. δ_{C}^a	$\Delta\delta$ (cal. – exp.)
1	86.2	88.0	1.8
2	160.3	161.0	0.7
3	128.7	128.4	-0.3
4	171.9	169.4	-2.5
4a	143.7	144.8	1.1
6	164.4	162.6	-1.8
6a	100.9	100.0	-0.9
7	164.7	163.0	-1.7
8	101.7	99.2	-2.5
9	166.6	164.4	-2.2
10	104.7	102.9	-1.8
10a	134.2	133.6	-0.6
10b	121.7	122.7	1.0
1-CH ₃	26.2	24.8	-1.4
3-OCH ₃	60.9	56.7	-4.2
9-OCH ₃	55.9	53.2	-2.7
1'	97.3	96.4	-0.9
2'	30.7	31.8	1.1
3'	25.4	26.8	1.4
4'	48.8	49.9	1.1
		MAE	1.59
		RMSD	1.82

^a Cal. δ_{C} was scaling corrected from calculated shielding tensor (σ) using the formula $\delta_{\text{C}} = (186.5242 - \sigma)/1.0533$.

Fig. 2 Key 2D NMR correlations of **1**.

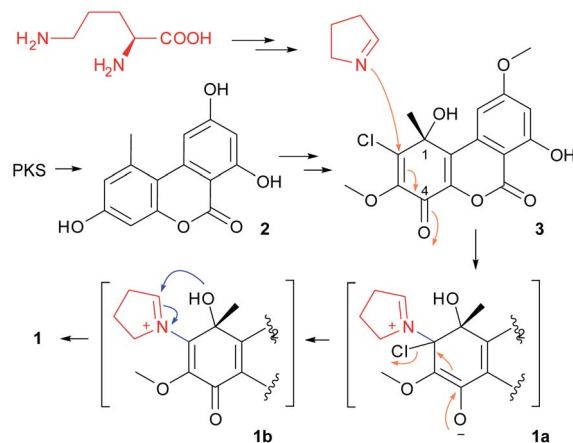
well with the experimental data with MAE 1.59 ppm and RMSD 1.82 ppm. Therefore, the structure of **1** was elucidated to be a hexahydropyrrolo[2,1-*b*]oxazole fused dibenzo- α -pyrone, hitherto not reported in nature.

To determine the absolute configuration of **1**, the TDDFT ECD computations were carried out on different level of theory (pbe0/TZVP, B3LYP/TZVP, BH&HLYP/TZVP) for the B3LYP/6-31G(d)-optimized conformers of **1**. The calculated spectrum of (1*R*,1'*S*)-**1** matched the experimental data (Fig. S1, ESI[†]), and the best agreement was found for the BH&HLYP/TZVP-calculated one (Fig. 3), thus the absolute configuration of **1** was unambiguously assigned.

Structurally, compound **1** was characterized as one pyrrolidine fused to a dibenzo- α -pyrone uniquely *via* an oxazolidine ring. The plausible biosynthetic pathway of **1** was proposed (Scheme 1). This should involve the formation of a key metabolite, TMC-264 (**3**), which was co-isolated from the titled fungus, and was synthesized *via* the polyketide pathway.¹⁶ This compound should be derived from alternariol (**2**) that was the first product of PKS,¹⁷ by several tailoring steps. The nucleophilic attack from Δ^1 -pyrroline that derived from L-ornithine, to the substrate **3**, would yield the iminium cation intermediate (**1a**), which then give rise to the derivative **1b** by ejecting the chloride ion (as a leaving group). The followed-up attack from 1-OH to C-1' in **1b** should give the oxazolidine (**1**). Iminium cations were reported to be widely involved in the biosynthesis of a range of alkaloids in nature.^{18,19}

Hexahydropyrrolo[2,1-*b*]oxazole or pyrrolidinooxazolidine alkaloids were not commonly found in nature. So far only 15 natural ones that incorporated in eight different skeletons have been reported, to our knowledge, which included two derivatives from the ladybird beetles, three from marine invertebrates, and ten from plants (Fig. S2, ESI[†]), however, none were reported from fungi. Some of these metabolites exhibited antibacterial, cytotoxic, antiplasmodial, antiviral, and phytotoxic activities,^{20–23} or playing a defense role against predators.²⁴

Compound **1** was thus screened for the antibacterial activity, however, it was inactive against the tested bacteria (MIC > 128 $\mu\text{g mL}^{-1}$). This compound was also tested for the antioxidant activity due to the presence of a phenolic group, whereas it did not exhibit strong activity in the DPPH radical scavenging assay



Scheme 1 Plausible biosynthetic pathway of **1**.

($\text{IC}_{50} > 200 \mu\text{g mL}^{-1}$). Meanwhile, in view of alkaloids being a major class of naturally-occurring acetylcholinesterase (AChE) inhibitors,²⁵ which could be of use as insecticides or a treatment for Alzheimer's disease, compound **1** was hence evaluated for its inhibitory activity against AChE. And a moderate inhibition ($\text{IC}_{50} 43.1 \mu\text{M}$) was found comparing to the positive control (tacrine hydrochloride, $\text{IC}_{50} 6.1 \mu\text{M}$).

Conclusions

In this study, rhizovagine A (**1**), the first fungal pyrrolidinooxazolidine alkaloid, bearing an unprecedented pentacyclic skeleton was isolated. The structure was elucidated by comprehensive spectroscopic analysis, together with quantum-chemical ¹³C NMR and ECD calculations. Compound **1** showed AChE-inhibitory activity, which could serve as a scaffold for developing new AChE inhibitors. Moreover, due to the structural novelty, **1** could be an intriguing target for realizing a total synthesis or biosynthesis.

Experimental section

General experimental procedures

The optical rotations were recorded on a Rudolph Autopol III automatic polarimeter (Rudolph Research Analytical, Hackensack, New Jersey). Ultraviolet (UV) spectra were recorded on a TU-1810 UV/Vis spectrophotometer (Beijing Persee General Instrument Co., Ltd., Beijing, China). Circular dichroism (CD) spectra were recorded on a JASCO J-815 CD spectrometer (JASCO Corp., Tokyo, Japan). High-resolution electrospray ionization mass spectrometry (HRESIMS) spectra were recorded on a LC 1260/Q-TOF-MS 6520 machine (Agilent Technologies, Santa Clara, CA). ¹H, ¹³C, and 2D NMR spectra were measured on Bruker Avance 400 NMR spectrometers (Bruker BioSpin, Zürich, Switzerland). ¹H and ¹³C NMR chemical shifts were expressed in δ (ppm) referring to the inner standard tetramethylsilane (TMS), and coupling constants in Hertz. HPLC-DAD analysis was performed on a Shimadzu LC-20A instrument equipping with a SPD-M20A photodiode array detector

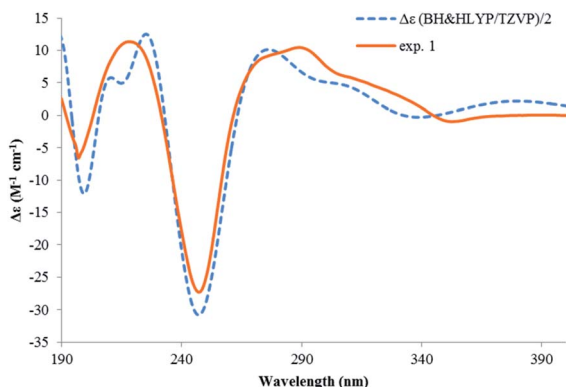


Fig. 3 Calculated and experimental ECD spectra of **1**.

(Shimadzu Corp., Tokyo, Japan) using an analytic C₁₈ column (250 mm × 4.6 mm i.d., 5 μm; Phenomenex Inc., Torrance, California). Semi-preparative HPLC separation was carried out on a Lumtech instrument (Lumiere Tech. Ltd., Beijing, China) equipped with a K-501 pump (flow rate: 3 mL min⁻¹) and a K-2501 UV detector (detection was set at 370 nm) using a Luna-C₁₈ column (250 mm × 10 mm i.d., 5 μm, Phenomenex Inc., Torrance, California) eluting with a mixture of MeOH and water.

Fungal material and fermentation

The endophytic fungus *Rhizopycnis vagum* Nitaf22 was isolated from *Nicotiana tabacum* previously in our lab.³

The fungus was cultured on potato dextrose agar for 5 days at 25 °C, and the resulting hyphae were used to inoculate 150 mL potato dextrose broth in a 250 mL flask. The culture was shaking for 7 days at 150 rpm and 25 °C, which was then used as inoculum to inoculate the autoclaved rice media in 1 L Erlenmeyer flasks each containing 100 g of rice and 110 mL of distilled water. The fermentation was carried out on a total of 10 kg of rice under static conditions at room temperature (RT) in the dark for 50 days.

Extraction and isolation

The culture was combined, dried and ground. The dry materials were extracted with EtOAc three times at RT, each for 5 days, which resulted in 163 g of extract after removal of the solvent.

The extract was subjected to vacuum liquid chromatography over silica gel (i.d. 8 cm × 30 cm) by eluting with a different mixture of petroleum ether and acetone, then CH₂Cl₂-MeOH. Fractions were pooled according to TLC, and five fractions were obtained (Frs. A-E). Fr. B was found to contain dibenzo- α -pyrones, which was then chromatographed by medium pressure liquid chromatography over silica gel using a gradient of CH₂Cl₂-MeOH (100 : 0-1 : 1, v/v) as the eluent to yield seven subfractions (Frs. B1-B7). Fr. B1 was further processed by chromatography over ODS using a gradient of MeOH/H₂O (35-100%) as the mobile phase, to afford nine further fractions (Frs. B1-1-B1-9). Among which, compound **1** (2.9 mg) was isolated from Fr. B1-5 by semi-preparative HPLC (63% MeOH/H₂O).

Rhizovagine A (1). Yellow amorphous powder; [α]_D²⁵ +163.2 (c 0.125, MeOH); UV (MeOH) λ_{\max} (log ϵ) 203 (4.57), 249 (4.66), 340 (3.86), 400 (3.84) nm; ECD (c = 4.33 × 10⁻⁴ M, MeOH) λ ($\Delta\epsilon$) 218 (+11.33), 247 (-27.33), 289 (+10.46), 352 (-0.97) nm; ¹H NMR (CDCl₃, 400 MHz), ¹³C NMR (CDCl₃, 100 MHz) see Table 1; HRESIMS *m/z* 386.1221 [M + H]⁺ (calcd for C₂₀H₂₀NO₇, 386.1234).

ECD calculation

Conformational searches using Merck Molecular Force Field (MMFF) were done by the MacroModel software. Geometry optimization and frequency calculations of the MMFF-conformers using the DFT method were performed by the Gaussian 09 package.²⁶ TDDFT ECD calculations of the low-energized conformers ($\geq 1\%$) without imaginary frequencies were carried out by Gaussian 09 using different functions (PBE0, B3LYP, BH&HLYP) and TZVP basis set with the

polarizable continuum model (PCM) for MeOH. The ECD spectrum of each conformer was simulated by the program SpecDis²⁷ using a Gaussian band shape with an exponential half-width σ of 0.25–0.3 eV, using the dipole-length computed rotational strengths. The equilibrium population of each conformer at 298.15 K was calculated from its relative Gibbs free energies using Boltzmann statistics. The Boltzmann-averaged ECD spectrum was then generated accordingly. The generated ECD spectra were compared with the experimental one to determine the absolute configuration. The calculated ECD spectra were UV-shifted (+7, +3, and +28 nm for PBE0, B3LYP, and BH&HLYP, respectively), for comparison with the measured spectrum.

¹³C NMR calculation

For the calculation of ¹³C NMR chemical shifts, B3LYP/6-31+G(d,p) method was used to optimize the selected conformers. NMR calculations were performed at the level of mPW1PW91/6-311+G(2d,p) with the gauge-independent atomic orbital (GIAO) method using the SMD model for CHCl₃. The ¹³C NMR chemical shifts were considered as the average value of the same atom in different conformers according to their Boltzmann distributions, using the relative Gibbs free energies as the weighting factors. The empirical scaling method was used to decrease the systematic error. The chemical shift (cal. δ_c) of each atom was scaling corrected from the calculated shielding tensor (σ) by using the formula $\delta_c = (186.5242 - \sigma)/1.0533$, using the scaling factors reported by Lodewyk, *et al.*¹⁵ The differences $\Delta\delta$ were determined by subtracting the experimental chemical shifts (δ_{exp}) from the calculated chemical shifts (δ_{cal}). The results were evaluated in terms of the mean absolute deviation (MAE), and root mean square deviation (RMSD).

Antibacterial assay

The antibacterial activity of compound **1** was evaluated against six pathogenic bacteria, *Agrobacterium tumefaciens*, *Bacillus subtilis*, *Pseudomonas lachrymans*, *Ralstonia solanacearum*, *Staphylococcus hemolyticus*, and *Xanthomonas vesicatoria*, as described previously.³

DPPH radical scavenging assay

The DPPH radical scavenging activity of compound **1** was determined based on the reduction of 1,1-diphenyl-2-picrylhydrazyl (DPPH) as reported previously.⁸ Butylated hydroxytoluene (BHT) was used as the positive control.

AchE inhibitory assay

The AchE inhibitory bioassay was performed using the method described by Meng *et al.*²⁸ Briefly, the assay was performed in 96-well plates, and in each well phosphate buffered saline (PBS, pH 7.7) solution (160 μL), AChE solution (1.0 U mL⁻¹, 10 μL), 5,5'-dithio-bis-(2-nitrobenzoic acid) solution (7.5 mM in PBS, 10 μL), and the tested compound of various concentrations (dissolved in 30% of DMSO-PBS, 10 μL) were added. After pre-incubation

at 30 °C for 15 min, 10 µL of acetylthiocholine iodide (10 mM in PBS) was added. The incubation was maintained for 30 min, then the absorbance was measured at 405 nm using a microplate spectrophotometer. Tacrine hydrochloride was used as the positive control, while PBS as the negative control. The percentage (%) of inhibition was determined using the following expression: $[(A_c - A_t)/A_c] \times 100$, where A_c is the absorbance of the negative control ($n = 3$), while A_t is that of the tested compounds ($n = 3$). The median inhibitory concentration (IC_{50}) was calculated using the linear regression between the inhibitory probability and concentration logarithm.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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