

Stemphones, Novel Potentiators of Imipenem Activity against Methicillin-resistant *Staphylococcus aureus*, Produced by *Aspergillus* sp. FKI-2136

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Abstract A fungal strain FKI-2136 identified as genus *Aspergillus* was found to produce potentiators of imipenem activity against methicillin-resistant *Staphylococcus aureus* (MRSA). Two new compounds designated stemphones B and C were isolated along with a structurally related known compound cochlioquinone D from the fermentation broth of the producing strain by solvent extraction, silica gel column chromatography and preparative HPLC. These compounds have a common tetracyclic quinone skeleton. Stemphone C potentiated imipenem activity against the MRSA 512 fold by decreasing MIC value of imipenem from 16 $\mu\text{g/ml}$ to 0.03 $\mu\text{g/ml}$.

Keywords stemphones, imipenem potentiator, methicillin-resistant *Staphylococcus aureus*, MRSA, fungal metabolite, tetracyclic quinone

Introduction

Methicillin-resistant *Staphylococcus aureus* (MRSA) is known as a major nosocomial pathogen which has also developed resistance to many other antibiotics. Moreover, MRSA has been reported to acquire resistance to the last-resort antibiotic, vancomycin [1, 2]. These facts suggest that *S. aureus* would fully acquire resistance to vancomycin

in near future. It is therefore increasingly important and necessary to find new antimicrobial agents and to devise new measures that are effective against MRSA infection.

Based on the new concept of “anti-infective drugs” developed by Ōmura [3], various potentiators of antifungal miconazole activity against *C. albicans* were discovered by our group [4–8]. It is expected that potentiators of imipenem activity against MRSA would, likewise, have application to the development of therapeutics against MRSA infection. During our screening for imipenem potentiators against MRSA from microbial origin, three structurally related active compounds were isolated from the culture broth of a fungal strain FKI-2136. One was identified as cochlioquinone D originally reported as an inhibitor of the root growth of the host plant seedlings such as ryegrass and rice [9], but the others were found to be new compounds designated stemphones B and C (Fig. 1). Stemphone (renamed stemphone A in this paper) previously reported as an inhibitor of the growth of *Bacillus megaterium* and *Sarcina lutea* [10] was not detected in the culture broth of strain FKI-2136. In this paper, the taxonomy of the producing strain, fermentation, isolation, structure elucidation and biological properties of stemphones are described.

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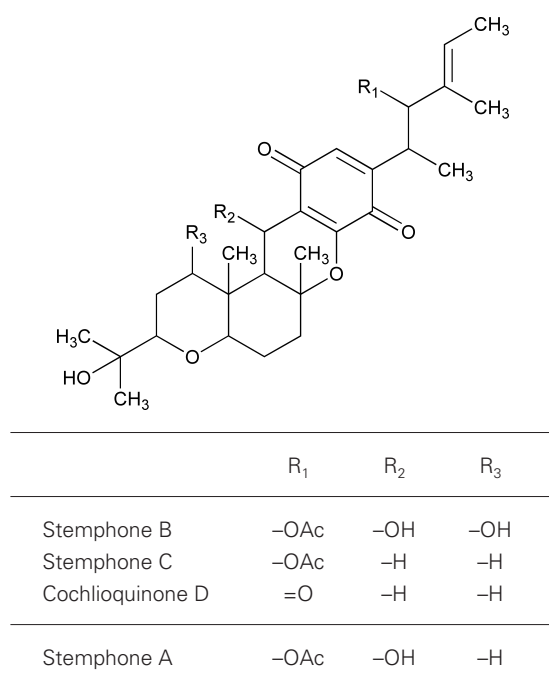


Fig. 1 Structures of stemphonones A, B and C and cochlioquinone D.

Materials and Methods

Materials

The following materials were purchased from commercial sources: Cloxacillin (ICN Biomedicals), cefazolin (Wako), vancomycin (Wako), tetracycline (Wako), ciprofloxacin (Wako), sulfate (Meiji Seika) and imipenem (Banyu Pharmaceutical). Erythromycin was isolated from microbial sources by our group.

General Experimental Procedures

Fungal strain FKI-2136 was isolated from a soil sample collected at Isigakijima Island, Okinawa Prefecture, Japan. This strain was used for production of stemphonones. Kieselgel 60 (E. Merck) and SSC-ODS-7515-12 (Senshu Sci.) were used for silica gel and octadecyl silyl (ODS) column chromatography, respectively. HPLC was carried out using an L-6200 system (Hitachi). For detection of stemphonones in culture broths, the samples (ethyl acetate extracts) dissolved in methanol were analyzed by an HP1100 system (Hewlett Packard) under the following conditions: column, Symmetry (2.1×150 mm, Waters); flow rate, 0.2 ml/minute; mobile phase, a 20-minute linear gradient from 60% CH₃CN to 100% CH₃CN containing 0.05% H₃PO₄; detection, UV at 210 nm. Under these conditions, stemphonones B, C and cochlioquinone D were

eluted as peaks with retention times of 4.7, 8.3 and 6.7 minutes, respectively.

UV spectra were recorded on a spectrophotometer (DU640, Beckman). IR spectra were recorded on a Fourier transform infrared spectrometer (FT-710, Horiba). Optical rotations were measured with a digital polarimeter (DIP-370, JASCO). EI-MS spectra and HREI-MS spectra were recorded on a mass spectrometer (JMS-AX505HA, JEOL). The various NMR spectra were collected with a spectrometer (XL-400, Varian).

Taxonomic Studies of the Producing Fungus

For the taxonomic studies of fungus, Czapeck yeast extract agar (CYA), malt extract agar (MEA), Czapeck yeast extract agar with 20% sucrose (CY20S) were used. Taxonomic studies and identification were conducted according to the procedures described by Klich [11]. Morphological observations were done under a light microscope (Vanox-S AH-2, Olympus) and a scanning electron microscope (JSM-5600, JEOL). Color names and hue numbers were determined according to the Color Harmony Manual [12].

Assay for Potentiating Activity of Imipenem against MRSA

MRSA K-24, a clinical isolate, was used and anti-MRSA activity was measured by two methods. 1) Paper disk method; MRSA was cultured in Mueller-Hinton broth (MHB, Difco) at 37°C for 20 hours and adjusted to 1×10⁸ CFU/ml. The inoculum (100 μl) was spread on MHA medium in a plate (10×14 cm, Eiken Kizai) containing MHB and 1.5% of agar (Shimizu Shokuhin) with or without imipenem (10 μg/ml), whose concentration has no effect on growth of MRSA. Paper disks (Advantec) containing various amounts of a sample were placed on the MHA plate and incubated at 37°C for 20 hours. Anti-MRSA activity was expressed as the diameter (mm) of the inhibitory zone on the MHA plate. 2) Liquid microdilution method [13]; after MHB (85 μl) was added to each well of a 96-well microplate (Corning), a sample dissolved in methanol (5 μl) was added to the final concentration as indicated (4 μg/ml for stemphone C as described below). Then imipenem, or other antimicrobial agents dissolved in distilled water (5 μl), were added to each well to a final concentration of 0.15 to 512 μg/ml. Finally, MRSA (5 μl) was added at a concentration of 1×10⁷ CFU/ml. Microplates were incubated at 37°C for 20 hours without shaking. MIC is defined as the lowest concentration of an antimicrobial agent where MRSA cannot grow.

Other Biological Assays

Antimicrobial activity against 15 species of microorganisms was measured by the paper disk method. The microorganisms were as follows; *Bacillus subtilis* ATCC 6633, *Staphylococcus aureus* FDA 209P, *Micrococcus luteus* PCI 1001, *Mycobacterium smegmatis* ATCC 607, *Escherichia coli* NIHJ, *E. coli* NIHJ-2 IFO 12734, *Pseudomonas aeruginosa* IFO3080, *Xanthomonas campestris* pv. *oryzae*, *Bacteroides fragilis* ATCC 23745, *Acholeplasma laidlawii* PG8, *Pyricularia oryzae* KF 180, *Aspergillus niger* ATCC6275, *Mucor racemosus* IFO4581, *Candida albicans* and *Saccharomyces cerevisiae*. Media for microorganisms were as follows: GAM agar (Nissui Seiyaku) for *B. fragilis*; Bacto PPLO agar (Difco) supplemented with horse serum 15%, glucose 0.1%, phenol red 0.25% and agar 1.5% for *A. laidlawii*; Nutrient agar (Difco) for the other bacteria; a medium composed of glucose 1.0%, yeast extract 0.5% and agar 0.8% for fungi and yeasts. A paper disk (i.d. 6 mm, Advantec) containing 10 μ g of sample was placed on an agar plate. Bacteria except *X. oryzae* were incubated at 37°C for 24 hours. Yeasts and *X. oryzae* were incubated at 27°C for 24 hours. Fungi were incubated at 27°C for 48 hours. Antimicrobial activity was expressed as diameter (mm) of the inhibitory zone.

Results

Taxonomy of the Producing Fungus

Colonies on CYA were 60~65 mm diameter after 7 days at 25°C, floccose to velutinous, radically sulcate, white (a) to light ivory (2ca) in color. The reverse side was cover tan (2ge). Colonies on MEA were 40~45 mm diameter, floccose to velutinous, light ivory (2ca) to bamboo (2gc). The reverse side was biscuit ecru (2ec). Colonies on

CY20S were 65~70 mm diameter, floccose to velutinous, radically sulcate, white (a) to light ivory (2ca). The reverse side was light wheat (2ea). At 5°C and 37°C, no colonies were observed on CYA. Soluble pigment was produced on CY20S and light tan (3gc) in color. Conidiogenesis on each medium was moderate to abundant. Conidiophores arose from foot cells and were 175~1500 μ m long. They were almost rough-walled and hyaline to slightly ivory. Vesicles were globose to subglobose and 15~60 μ m in diameter. Aspergilla (as shown in Figs. 2A and 2B) were biseriate and formed metulae and phialides. Metulae covered the entire surface of the vesicle, were cylindrical and 6~12 \times 3~6 μ m in size. Phialides were flask-shaped and 5~10 \times 2~3 μ m in size. Conidia (as shown in Fig. 2C) were globose to subglobose with finely roughened walls and 2~4 μ m in size.

From the above characteristics, strain FKI-2136 was considered to belong to the genus *Aspergillus* and named *Aspergillus* sp. FKI-2136. The strain was deposited at the National Institute of Bioscience and Human-Technology, Agency of Industrial Science and Technology Japan, as NITE BP-83.

Fermentation

A slant culture of the strain FKI-2136 grown on LCA (glycerol 0.1%, KH_2PO_4 0.08%, K_2HPO_4 0.02%, $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$ 0.02%, KCl 0.02%, NaNO_3 0.2%, yeast extract 0.02%, agar 1.5%, pH 6.0) was used to inoculate a 50-ml tube containing 10 ml of the seed medium (glucose 2.0%, polypeptone 0.5%, $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$ 0.05%, yeast extract 0.2%, KH_2PO_4 0.1%, agar 0.1%, pH 6.0). The tube was shaken on a reciprocal shaker at 27°C for 2 days. A one-ml portion of the seed culture was then inoculated into a 500-ml Erlenmeyer flask containing 100 ml of the production medium (glucose 1.0%, soluble starch 2.0%, soy bean oil 2.0%, pharmamedia 1.0%, meat extract 0.5%, $\text{MgSO}_4 \cdot$

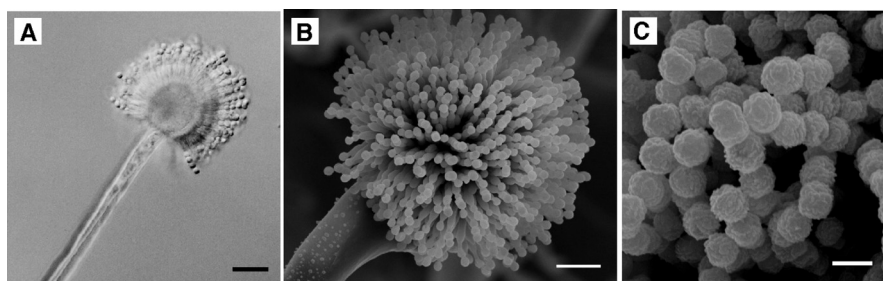


Fig. 2 Morphological observation of fungal strain FKI-2136.

(A) Conidial head under a light microscope (scale bar: 20 μ m).

(B) Biseriate aspergilla under a scanning electron microscope (scale bar: 10 μ m).

(C) Conidia under a scanning electron microscope (scale bar: 2 μ m).

7H₂O 0.1%, CaCO₃ 0.3%, trace salt solutions (FeSO₄·7H₂O 0.1%, MnCl₂·4H₂O 0.1%, ZnSO₄·7H₂O 0.1%, CuSO₄·5H₂O 0.1%, CoCl₂·6H₂O 0.1%) 1.0%, agar 0.1%, pH 6.0). The fermentation was carried out at 27°C for 11 days. Stemphones B and C and cochlioquinone D were detected in the culture broth from day 2 to day 3 after inoculation. Stemphone B and cochlioquinone D reached a maximal level (230 and 109 µg/ml, respectively) on day 5, and the titers gradually decreased until day 11. On the other hand, stemphone C reached a maximal level (1.1 mg/ml) on day 11.

Isolation

The 11-day old whole broth (1 liter) was centrifuged at 3000 rpm for 10 minutes. The mycelium was extracted with acetone (1.5 liters). After the acetone extracts were filtered and concentrated to remove acetone, the aqueous residue was extracted with ethyl acetate. The extracts were dried over Na₂SO₄ and concentrated *in vacuo* to dryness to yield crude materials (4.4 g). The materials were dissolved in CHCl₃, applied on a silica gel column (59 g), and eluted stepwise with 2:1 (v/v) of hexane-EtOAc solvent and 100:0, 100:1, 50:1, 10:1 and 1:1 (v/v) of CHCl₃-CH₃OH solvents (120 ml for each solvent). The 10:1 fraction (CHCl₃-CH₃OH) showing the activity was concentrated to give a red brown oily material (786 mg), which was subjected to a second silica gel column (50 g). The material was eluted with 100:1, 50:1, 20:1, 10:1 and 5:1 (v/v) of CHCl₃-CH₃OH solvents (12 ml×10 tubes for each solvent). The first active fraction from the 2nd tube of 100:1 to the 3rd tube of 50:1 (CHCl₃-CH₃OH) was collected and concentrated to give a red brown oil (274 mg). The oil containing enriched stemphone C was finally purified with preparative HPLC (column; PEGASIL ODS (Senshu Sci. Co.), 20×250 mm; solvent, 70% CH₃CN; detection, UV at 210 nm; flow rate, 6.0 ml/minute). Under these conditions, cochlioquinone D and stemphone C were eluted as peaks with retention times of 23 and 28 minutes, respectively (Fig. 3A). Each pooled fraction was concentrated *in vacuo* to dryness to give pure stemphone C (174 mg) and cochlioquinone D (42 mg) as yellow materials. The second active fraction from the 4th tube of 50:1 to the 3rd tube of 20:1 (CHCl₃-CH₃OH) containing stemphone B was also purified by HPLC (column; PEGASIL ODS, 20×250 mm; solvent, 55% CH₃CN, UV at 210 nm; flow rate, 9 ml/minute). Stemphone B was eluted as a peak with a retention time of 20 minutes (Fig. 3B). The fraction was concentrated *in vacuo* to dryness to yield pure stemphone B (58 mg) as a yellow material.

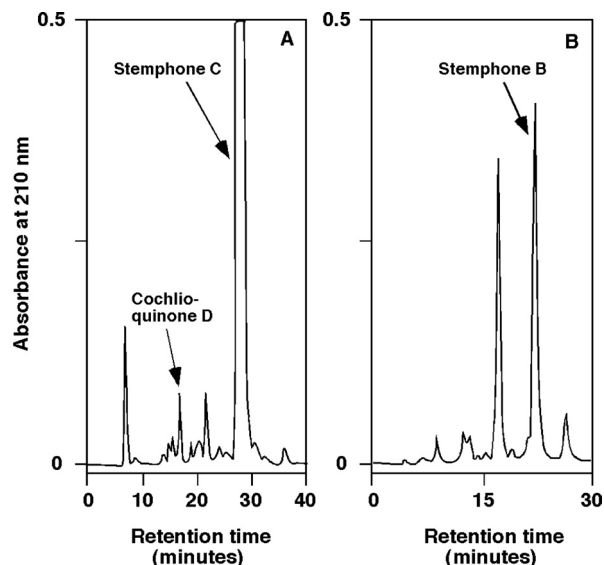


Fig. 3 Chromatographic profiles of isolation of stemphones by HPLC.

(A) Stemphone C and cochlioquinone D. Column, PEGASIL ODS 20×250 mm; solvent 70% aq acetonitrile; detection, UV at 210 nm; flow rate, 6.0 ml/minute; sample, 100 mg of active materials (obtained through second silica gel column chromatography) dissolved in 50 µl methanol was injected.

(B) Stemphone B. Column, PEGASIL ODS 20×250 mm; solvent 55% aq acetonitrile; detection, UV at 210 nm; flow rate, 9.0 ml/minute; sample, 100 mg of active materials (obtained through second silica gel column chromatography) dissolved in 50 µl methanol was injected.

Table 1 Physico-chemical properties of stemphones B and C

	Stemphone B	Stemphone C
Appearance	Yellow powder	Yellow powder
Molecular weight	546	514
Molecular formula	C ₃₀ H ₄₂ O ₉	C ₃₀ H ₄₂ O ₇
HREI-MS (<i>m/z</i>)		
Calcd.:	546.2835 (M ⁺)	514.2926 (M ⁺)
Found:	546.2829	514.2931
UV (MeOH) λ _{max} nm (ε)	206 (7748), 267 (5154), 383 (505)	207 (7604), 265 (7239), 398 (616)
[α] _D ²⁶	+144.0° (c=0.1, MeOH)	+94.6° (c=0.1, MeOH)
IR (KBr) ν _{max} (cm ⁻¹)	3440, 2935, 1727, 1644, 1602	3444, 2944, 1739, 1643, 1604

Physico-chemical Properties of Stemphones

Physico-chemical properties of stemphones are summarized in Table 1. They showed similar UV absorption maxima at 265 to 267 nm. The IR absorption at 3444~2935 and 1739~1602 cm^{-1} suggested the presence of hydroxy and carbonyl groups in the structures.

Structure Elucidation of Stemphone B

The molecular formula of stemphone B was determined to be $\text{C}_{30}\text{H}_{42}\text{O}_9$ on the basis of HREI-MS measurement. The ^{13}C NMR spectrum (in CDCl_3) showed 30 resolved signals,

which were classified into eight methyl carbons, three methylene carbons, two sp^3 methine carbons, two sp^2 methine carbons, five oxygenated sp^3 methine carbons, one sp^3 quaternary carbon, two oxygenated sp^3 quaternary carbons, four sp^2 quaternary carbons and three carbonyl carbons by analysis of the DEPT spectra. The ^1H NMR spectrum (in CDCl_3) displayed 41 proton signals, two of which were suggested to be hydroxy protons (δ 3.42 and δ 4.13). Taking the molecular formula into consideration, the presence of another hydroxy proton was suggested. The connectivity of proton and carbon atoms was established by

Table 2 ^1H and ^{13}C NMR chemical shifts of stemphones B and C and cochlioquinone D

Position	Stemphone B		Stemphone C		Cochlioquinone D	
	δ_{C}	δ_{H} (J Hz)	δ_{C}	δ_{H} (J Hz)	δ_{C}	δ_{H} (J Hz)
1	13.2 q	1.61 d, br (7.0)	13.1 q	1.59 d, br (7.0)	15.0 q	1.87 d (7.0)
2	125.3 d	5.56 q, br (7.0)	124.9 d	5.53 q (7.0)	138.6 d	6.86 q (7.0)
3	131.7 s	—	132.1 s	—	136.8 s	—
4	81.4 d	5.15 d (8.5)	81.5 d	5.14 d (8.5)	200.5 s	—
5	34.2 d	3.30 dq (8.5, 7.0)	33.5 d	3.34 dq (8.5, 7.0)	37.6 d	4.62 q (7.0)
6	148.8 s	—	148.0 s	—	146.5 s	—
7	181.0 s	—	181.3 s	—	181.4 s	—
8	151.5 s	—	152.3 s	—	152.1 s	—
9	117.8 s	—	117.8 s	—	118.4 s	—
10	188.5 s	—	187.1 s	—	186.8 s	—
11	132.6 d	6.47 s	132.4 d	6.48 d (1.0)	133.2 d	6.51 d (1.0)
12	62.3 d	4.90 d (11.0)	16.4 t	2.11 dd (19.0, 13.0) 2.52 dd (19.0, 5.0)	16.5 t	2.13 dd (19.0, 13.0) 2.52 dd (19.0, 5.0)
12-OH ^a	—	3.42, br	—	—	—	—
13	45.4 d	2.23 d (11.0)	46.6 d	1.43 dd (13.0, 5.0)	46.5 d	1.44 dd (13.0, 5.0)
14	83.8 s	—	80.5 s	—	80.6 s	—
15	37.1 t	1.88 m, 2.08 m	37.2 t	1.79 m, 2.18 m	37.2 t	1.80 m, 2.13 m
16	25.0 t	1.58 m, 1.86 m	25.2 t	1.63 m, 1.80 m	25.2 t	1.80 m
17	76.2 d	3.68 dd (13.0, 4.0)	84.0 d	3.12 dd (12.0, 4.0)	84.0 d	3.12 dd (12.0, 3.5)
18	40.8 s	—	35.5 s	—	35.5 s	—
19	70.7 d	4.10 t, br (3.0)	36.8 t	1.21 m, 1.86 m	36.8 t	1.20 m, 1.92 m
19-OH ^a	—	4.13, br	—	—	—	—
20	28.2 t	1.60 m, 1.96 m	21.3 t	1.48 m, 1.66 m	21.3 t	1.48 m
21	79.6 d	3.57 dd (13.0, 3.0)	84.9 d	3.21 dd (12.0, 3.0)	84.9 d	3.22 dd (12.0, 3.0)
22	71.7 s	—	71.8 s	—	71.8 s	—
23	23.8 q	1.14 s	23.8 q	1.16 s	23.8 q	1.17 s
24	26.4 q	1.20 s	26.1 q	1.18 s	26.1 q	1.19 s
25	13.1 q	0.98 s	12.2 q	0.88 s	12.2 q	0.89 s
26	21.3 q	1.32 s	20.8 q	1.28 s	20.8 q	1.27 s
27	16.9 q	1.04 d (7.0)	17.0 q	1.02 d (7.0)	16.4 q	1.27 d (7.0)
28	11.6 q	1.62 s	11.6 q	1.62 s, br	11.5 q	1.77 s, br
29	169.7 s	—	169.7 s	—	—	—
30	21.2 q	1.94 s	21.1 q	1.93 s	—	—

^a Exchangeable signals.

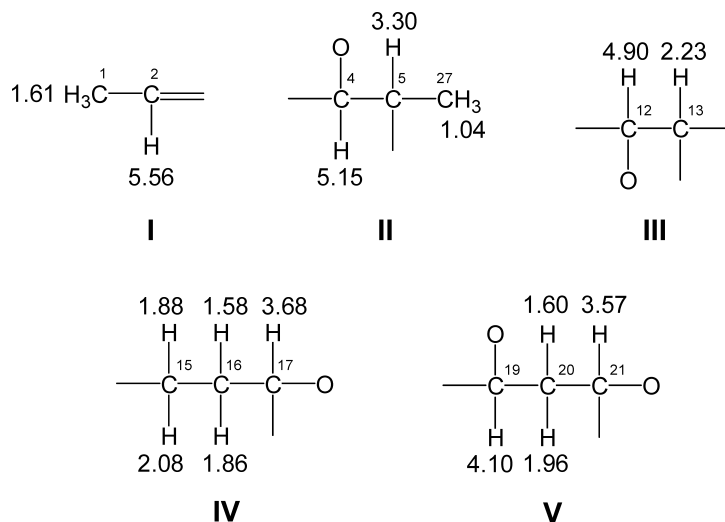


Fig. 4 Partial structures of stemphone B.

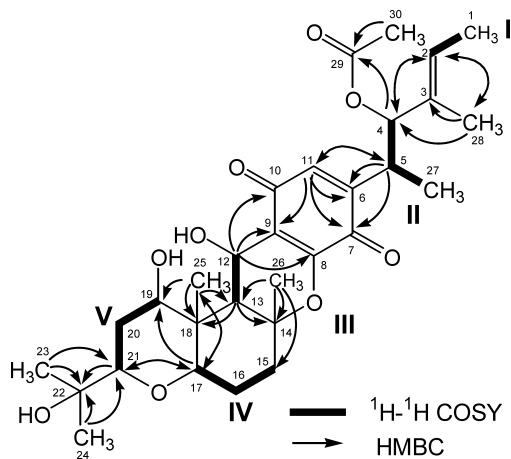


Fig. 5 ^1H - ^1H COSY and ^{13}C - ^1H HMBC experiments of stemphone B.

the ^{13}C - ^1H HMQC spectrum (Table 2). Analyses of the ^1H - ^1H COSY revealed the presence of five partial structures I to V as shown in Fig. 4. Furthermore, ^{13}C - ^1H long range couplings of 2J and 3J observed in the ^{13}C - ^1H HMBC spectrum (Fig. 5) gave the following linkages: 1) The cross peaks from H_3 -28 (δ 1.62) to C-2 (δ 125.3), C-3 (δ 131.7) and C-4 (δ 81.4), and from H-4 (δ 5.15) to C-2 showed the connection between the partial structures I and II. 2) The cross peaks from H-4 and H_3 -30 (δ 1.94) to C-29 (δ 169.7) showed that an acetoxy group was connected to the C-4 oxygenated methine carbon, indicating the presence of a side chain containing an acetoxy group and the partial structures I and II. 3) The cross peaks from H-5 (δ 3.30) to C-6 (δ 148.8), C-7 (δ 181.0) and C-11 (δ 132.6), from H-11 (δ 6.47) to C-5 (δ 34.2), C-6, C-7 and C-9 (δ 117.8),

and from H-12 (δ 4.90) to C-8 (δ 151.5), C-9 and C-10 (δ 188.5) showed the presence of a quinone skeleton, which was also supported by the chemical shifts of C-7 and C-10, and UV [10, 14, 15]. 4) The cross peaks from H_3 -26 (δ 1.32) to C-13 (δ 45.4), C-14 (δ 83.8) and C-15 (δ 37.1), from H-13 to C-14 and C-18 (δ 40.8), from H_3 -25 (δ 0.98) to C-13, C-17 (δ 76.2), C-18 and C-19 (δ 70.7), and from H-17 (δ 3.68) to C-19 and C-21 (δ 79.6) showed the connection from the partial structures III to V. Taking the degree of unsaturation into consideration, a tetracyclic structure containing the quinone ring should be formed as shown in Fig. 5. Further, the chemical shifts of C-8, C-14, C-17 and C-21 correspond to the values of oxygenated carbons, indicating that the tetracyclic structure contains two tetrahydropyran rings. 5) The cross peaks from H_3 -23 (δ 1.14) and H_3 -24 (δ 1.20) to C-21 and C-22 (δ 71.7), and the chemical shift of C-22 showed the presence of a 2-hydroxypropyl group. These results satisfied the degree of unsaturation and the molecular formula. The chemical shifts of stemphone B were comparable to those of stemphone A except for C-19 [15]. Taken together, the structure of stemphone B was elucidated as shown in Fig. 1. The geometry of the double bond between C-2 and C-3 was determined to be E configuration because stemphone B shares the same chemical shifts of stemphone A [15].

Structure Elucidation of Stemphone C

The molecular formula $\text{C}_{30}\text{H}_{42}\text{O}_7$ of stemphone C was two oxygen atoms smaller than stemphone B. The ^1H NMR spectrum of stemphone C was similar to stemphone B, but two signals corresponding to a hydroxy proton were missing. Correspondingly, two methylene carbon signals (δ 16.4 and δ 36.8) were observed instead of the two

oxygenated methine carbon signals for stemphone B by analysis of the ^{13}C NMR and DEPT spectrum. The position of the methylene carbons was determined by ^1H - ^1H COSY and HMBC experiments as shown in Fig. 6. The cross peaks from H₂-12 (δ 2.11, 2.52) to C-8 (δ 152.3), C-9 (δ 117.8), C-10 (δ 187.1) and C-18 (δ 35.5) indicated that the carbon (δ 16.4) was the C-12 position. The cross peaks from H-13 (δ 1.43), H-17 (δ 3.12) and H-20 (δ 1.48, 1.66) to C-19 (δ 36.8) indicated that the other one (δ 36.8) was the C-19 position. The chemical shifts of stemphone C were comparable to those of stemphone A except for C-12 [15]. Taken together, the structure of stemphone C was elucidated as shown in Fig. 1.

Biological Properties

Potential of Imipenem Activity against MRSA by Stemphones

By the paper disk assay, cochlioquinone D, and stemphones

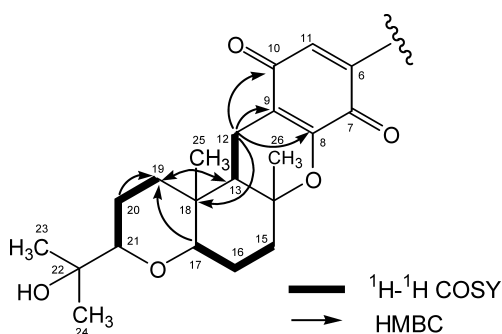


Fig. 6 ^1H - ^1H COSY and ^{13}C - ^1H HMBC experiments of stemphone C.

B and C showed almost no anti-MRSA activity at least up to $50\ \mu\text{g}/6\ \text{mm}$ disk, but showed anti-MRSA activity on MHA plates containing imipenem ($10\ \mu\text{g}/\text{ml}$) with inhibition zones of 9, 15 and 20 mm at $5\ \mu\text{g}/6\ \text{mm}$ disk, respectively. Next, the potentiating effect of stemphone C on the activity of other typical antimicrobial agents against MRSA was investigated by the liquid microdilution method. First, the effect of stemphone C itself was tested on the growth of MRSA in this method, resulting in an MIC value of $16\ \mu\text{g}/\text{ml}$. Therefore, the concentration of stemphone C was set up at $4\ \mu\text{g}/\text{ml}$ ($7.8\ \mu\text{M}$, one fourth of the MIC value, no effect on MRSA growth) to investigate the potentiating activity of the agents against MRSA. As summarized in Table 3, stemphone C markedly reduced the MIC value of imipenem from 16 to $0.03\ \mu\text{g}/\text{ml}$ and that of cloxacillin from 512 to $1.0\ \mu\text{g}/\text{ml}$, yielding 512-fold potentiation of their activity. Cefazolin activity against MRSA was also enhanced in combination with stemphone C. On the other hand, other antimicrobial agents showed almost no effect or slight potentiation (1~4-fold). These data suggested that stemphone C could potentiate β -lactam activity against MRSA.

Other Biological Activities

Stemphones B and C showed comparable antimicrobial activities against *B. subtilis*, *S. aureus*, *M. luteus*, and *A. laidlawii*, with almost the same inhibition zones of 9, 8, 14 and 9 mm at $5\ \mu\text{g}/6\ \text{mm}$ disk, respectively. On the other hand, cochlioquinone D showed a weak activity against only *M. luteus* with an inhibition zone of 8 mm at $5\ \mu\text{g}/6\ \text{mm}$ disk.

Table 3 MIC values of various antimicrobial agents against MRSA in the absence or presence of stemphone C

Antimicrobial agent	MIC ($\mu\text{g}/\text{ml}$)		Ratio (-/+ stemphone C)
	-	+ stemphone C*	
Imipenem	16	0.03	512
Cloxacillin	512	1	512
Cefazolin	64	4	16
Vancomycin	0.5	0.5	1
Streptomycin	2	0.5	4
Tetracycline	32	32	1
Erythromycin	>256	>256	1
Ciprofloxacin	64	32	2

* Concentration of stemphone C is $4\ \mu\text{g}/\text{ml}$.

Discussion

Stemphones and cochlioquinones, both of fungal origin, are structurally related by having a tetracyclic quinone skeleton in common. Compounds of stemphone family have been reported to show various biological activities including: nematocidal activity of cochlioquinone A [16]; inhibitory activity against diacylglycerol acyltransferase by cochlioquinones [17]; and inhibitory activity against cholesterol acyltransferase by epi-cochlioquinone [18]. In this report, we showed that stemphones B and C as well as cochlioquinone D exhibited activity as potentiator of β -lactam imipenem against MRSA. Among these compounds, stemphone C was the most potent followed by stemphone B. Cochlioquinone D showed only moderate activity. These results indicated that the acetoxyl residue at C-4 is important for the potentiating activity. The relationships of different biological activities exhibited by the stemphone family remain to be established.

Several natural products are known to potentiate β -lactam activity against MRSA. Polyoxotungstates were reported to show the activity possibly due to inhibition of penicillin binding proteins (PBPs) and β -lactamase [19]. Polyphenols such as epigallocatechin gallate isolated from tea [20], corilagin [21] and tellimagrandin I [22] were also reported to show the activity. They were considered to inhibit the function of PBP2' and suppress β -lactamase activity [23]. The potentiating activity of various β -lactams against MRSA by stemphone and the mechanism of the potentiation activity are under investigation and will be published in the near future.

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