### ORIGINAL ARTICLE

# Blockage of the early step of lankacidin biosynthesis caused a large production of pentamycin, citreodiol and *epi*-citreodiol in *Streptomyces rochei*

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In our effort to find the key intermediates of lankacidin biosynthesis in *Streptomyces rochei*, three UV-active compounds were isolated from mutant FS18, a gene disruptant of *lkcA* encoding a non-ribosomal peptide synthetase (NRPS)-polyketide synthase (PKS) hybrid enzyme. Their structures were elucidated on the basis of spectroscopic data of NMR and MS. Two compounds of a higher mobile spot on silica gel TLC ( $R_f = 0.45$  in CHCl<sub>3</sub>-MeOH = 20:1) were determined to be an epimeric mixture of citreodiol and *epi*-citreodiol at the C-6 position in the ratio of 2:1. In contrast, the compound of a lower mobile spot ( $R_f = ~0$  in CHCl<sub>3</sub>-MeOH = 20:1) was identical to a 28-membered polyene macrolide pentamycin. The yields of citreodiols and pentamycin in FS18 were 5- and 250-fold higher compared with the parent strain. Introduction of a second mutation of *srrX*, coding a biosynthetic gene of the signaling molecules SRBs, into mutant FS18 did not affect the production of three metabolites. Thus, their production was not regulated by the SRB signaling molecules in contrast to lankacidin or lankamycin. *The Journal of Antibiotics* advance online publication, 3 December 2014; doi:10.1038/ja.2014.160

#### INTRODUCTION

The filamentous soil bacteria *Streptomyces* species produce a vast array of secondary metabolites including antibacterial, antifungal, anticancer, antiparasitic, herbicidic and immunosuppressant agents. Recent development of the genome sequencing strategy revealed that *Streptomyces* strains usually contain over 20 secondary metabolite biosynthetic gene clusters, many of which are cryptic and silent under laboratory culture condition.<sup>1</sup> Hence, *Streptomyces* genome is well considered to be an utmost important reservoir of natural products for new pharmaceutical applications.

Streptomyces rochei 7434AN4 produces two structurally unrelated polyketide antibiotics, lankacidin and lankamycin (Figure 1), and carries three linear plasmids, pSLA2-L, -M and -S.<sup>2</sup> Lankacidin and lankamycin inhibit peptide synthesis synergistically by binding to the neighboring sites in the large ribosomal subunit,<sup>3,4</sup> and their biosyntheses are strictly controlled by the small diffusible signaling molecules termed SRBs (Streptomyces rochei butenolides) in S. rochei.<sup>5</sup> In particular, lankacidin is a unique 17-membered carbocyclic polyketide with various biological activities.<sup>6</sup> Nucleotide sequencing of the largest plasmid pSLA2-L (210 614 bp) together with extensive mutational analysis revealed that the lankacidin biosynthetic gene (lkc) cluster was located on pSLA2-L.7-10 The lkc cluster contains a non-ribosomal peptide synthetase (NRPS)/polyketide synthase (PKS) hybrid gene (lkcA), three multidomain PKSs (lkcC, lkcF, lkcG), discrete dehydratase (lkcB) and acyltransferase (lkcD) genes, an amine oxidase gene (lkcE) and a gene cluster for pyrroloquinoline quinone (PQQ)

biosynthesis (*lkcK-lkcO*). Recently, involvement of an additional dehydratase activity in lankacidin biosynthesis was reported by Dickschat *et al.*<sup>11</sup>

It is noteworthy that the *lkc* cluster contains only five ketosynthase domains, although eight rounds of condensation are required for the construction of the lankacidin skeleton, suggesting the modulariterative mixed polyketide biosynthesis of lankacidin. We carried out extensive mutations on the *lkc*-PKS genes to isolate the key intermediates supporting this hypothesis.<sup>8,10</sup> Although we have not yet succeeded to isolate such intermediates, we noticed that the *lkcA* disruptant FS18 produced large amounts of UV-active compounds compared with the parent strain. We reported here the isolation and structural elucidation of citreodiol (**5a**), *epi-c*itreodiol (**5b**) and pentamycin (**6**) from this mutant. In addition, we also describe that the production of the three metabolites was not regulated by the SRB signaling molecules based on the result of an additional mutation of *srrX*.

#### RESULTS

#### Metabolites from the lkcA disruptant FS18

The *lkcA* mutant, termed FS18, exhibited complex metabolic changes in comparison with the parent and other *lkc*-PKS mutants. Strain FS18 failed to produce lankacidin and its derivatives due to the mutation in the early biosynthetic step of lankacidin (Figure 2). As for the lankamycin production, FS18 produced a similar level of lankamycin  $(8.2 \text{ mg} \text{l}^{-1})$  compared with the parent strain 51252 (8.5 mg l<sup>-1</sup>).

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Received 23 June 2014; revised 23 October 2014; accepted 6 November 2014

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Figure 1 Structures of lankacidin C (1), lankacidinol (2), lankacidinol A (3), lankamycin (4), citreodiol (5a), *epi*-citreodiol (5b) and pentamycin (6) isolated from *Streptomyces rochei* strains 51252 (parent) and FS18 (Δ*lkcA*). Ac, acetyl; Me, methyl.



**Figure 2** Analysis of the metabolites produced by strain FS18. (a) HPLC chromatograms acquired using photodiode array detector. (I) strain 51252 (parent), (II) strain FS18 ( $\Delta/kcA$ ). The crude extract was applied on a COSMOSIL 5C18-MS-II column (4.6×250 mm) and eluted with a mixture of acetonitrile-10 mm sodium phosphate buffer (pH 8.2) (3:7, v/v) at a flow rate of 1.0 ml min<sup>-1</sup>. (b) The UV-visible spectra of compounds 1-3, 5a, 5b and 6. (c) Silica gel TLC analysis of metabolites from 51252 (parent), FS18 ( $\Delta/kcA$ ) and KA78 ( $\Delta/kcA\Delta srrX$ ) when irradiated with UV lamp at 254 nm (I) and stained with a mixture of CHCl<sub>3</sub>-MeOH (15:1, v/v). (d) Bioassay of crude extracts against *Aspergillus niger*. The plate was incubated at 28 °C for 5 days.

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**Figure 3** Scheme of a protection-deprotection strategy for purification of compounds **5a** and **5b**. Treatment: (a) carbonyldiimidazole, toluene; (b) separation by silica gel chromatography; (c) 0.5 M NaOH, THF-MeOH (1:1, v/v).

Remarkably, this strain accumulated the large amounts of UV-active compounds (5 and 6) (Figures 2a and c) and exhibited significant antifungal activity against *Aspergillus niger* (Figure 2d).

The UV-active compound 5, a spot with strong UV-absorbance on TLC ( $R_f = 0.45$  in CHCl<sub>3</sub>-MeOH = 20:1), was successively purified by Sephadex LH20 with MeOH and silica gel chromatography with  $CHCl_3$ -MeOH = 50:1-30:1. This compound was detected at 5.9 min on HPLC with a maximum absorbance at 267 nm, which differed from that of lankacidin antibiotics ( $\lambda_{max} = 230 \text{ nm}$ ) (Figures 2a and b). High-resolution positive ESI-MS gave a distinct molecular ion peak at 237.1096 [M+Na]<sup>+</sup> (calcd. for C<sub>11</sub>H<sub>18</sub>O<sub>4</sub>Na, 237.1097). However, 22 carbon signals were observed in its <sup>13</sup>C-NMR spectrum, which suggests that this compound is a mixture of inseparable isomers in the ratio of 2:1 (5a:5b). To separate 5a and 5b, a protection-deprotection strategy was carried out using carbonyldiimidazole in toluene according to Shizuri et al.<sup>12</sup> as shown in Figure 3. The first step quantitatively vielded a mixture of carbonate derivatives 7a and 7b, which were then separated by silica gel chromatography ( $R_f = 0.55$  for 7a and 0.50 for 7b in hexane-EtOAc = 2:1). Deprotection of the carbonate group by treatment with 0.5 M NaOH in THF-MeOH (1:1, v/v) afforded diastereomerically pure 5a and 5b. From their spectral data, compounds 5a and 5b were identical to citreodiol (5a; Figure 1) and epicitreodiol (5b), both of which were originally isolated from Penicillium citreoviride B.12 As no complete set of NMR assignments has been reported, the <sup>1</sup>H and <sup>13</sup>C spectral data were summarized in Table 1. The optical rotation values of **5a** and **5b** were  $[\alpha]_D^{25} = -4.8$  (c = 0.21, CHCl<sub>3</sub>) and  $[\alpha]_D^{26} = +6.1$  (c = 0.16, CHCl<sub>3</sub>), respectively. All spectral data of 5a and 5b were identical to the reported values for synthetic (6S,7S)(+)-citreodiol and (6R,7S)(-)-epi-citreodiol, except for their opposite rotation values;  $[\alpha]_D^{22} = +4.3$  (c = 1.0, CHCl<sub>3</sub>) for (+)-citreodiol and  $[\alpha]_{D}^{21} = -7.1$  (*c* = 2.3, CHCl<sub>3</sub>) for (-)-*epi*-citreodiol.<sup>12-14</sup> Thus, compounds 5a and 5b were elucidated to be (6R,7R,2E,4E)-6,7-dihydroxy-2,6-dimethylocta-2,4-dienoic acid methyl ester and (6S,7R,2E,4E)-6,7-dihydroxy-2,6-dimethylocta-2,4-dienoic acid methyl ester, respectively. This is the first discovery of 5a and 5b from Streptomyces species. Average yield of 5 in FS18 was calculated to be  $1.2 \text{ mg} \text{l}^{-1}$  from the HPLC peak intensity, which was fivefold larger compared with the parent strain 51252.

Compound **6** with strong UV absorbance was detected as a bluegray spot on TLC when baked after anisaldehyde-H<sub>2</sub>SO<sub>4</sub> spraying ( $R_{\rm fS} = \sim 0$  in CHCl<sub>3</sub>-MeOH = 20:1 and 0.44 in CHCl<sub>3</sub>-EtOAc-MeOH-H<sub>2</sub>O = 22:22:44:11), and appeared as a distinct peak at 28.8 min on HPLC (Figures 2a and c). Its UV spectrum ( $\lambda_{\rm max}$  at 322, 338, 356 nm) (Figure 2b) was characteristic of methylpentaenes such as filipin and

Table 1 NMR data of citreodiol (5a) and epi-citreodiol (5b) in CDCl<sub>3</sub>

	Citreodiol ( <b>5a</b> )		epi-Citreodiol ( <b>5b</b> )	
Position	δC (Mult.)	δH (Mult., J in Hz)	δC (Mult.)	δH (Mult., J in Hz)
1	168.8 (s)	_	168.9 (s)	_
2	127.6 (s)	_	127.4 (s)	_
3	137.4 (d)	7.19 (d, 11)	137.6 (d)	7.21 (d, 11)
4	124.8 (d)	6.68 (dd, 15 and 11)	125.1 (d)	6.66 (dd, 15 and 11)
5	145.1 (d)	6.10 (d, 15)	142.9 (d)	6.12 (d, 15)
6	75.7 (s)	_	75.6 (s)	_
7	73.1 (d)	3.70 (q, 6.4)	74.1 (d)	3.71 (q, 6.4)
8	16.9 (q)	1.18 (d, 6.4)	18.1 (q)	1.18 (d, 6.4)
2-Me	12.8 (q)	1.98 (d, 1.0)	12.8 (q)	1.98 (d,1.5)
6-Me	22.0 (q)	1.28 (s)	24.8 (q)	1.34 (s)
OMe	51.9 (q)	3.76 (s)	51.9 (q)	3.76 (s)

Spectra were recorded at 500 MHz for <sup>1</sup>H and 125 MHz for <sup>13</sup>C.

The residential solvent signal ( $\delta C = 77$ ) and internal standard tetramethylsilane ( $\delta H = 0$ ) were used as a reference.

pentamycin.<sup>15,16</sup> This compound was successively purified by Sephadex LH20 with MeOH and a preparative HPLC with acetonitrile-10 mM sodium phosphate buffer (pH 8.2). Its molecular formula was determined by high-resolution positive ESI-MS to be C35H58O12 (observed *m/z* of 693.3810, calcd. for C<sub>35</sub>H<sub>58</sub>O<sub>12</sub>Na [M+Na]<sup>+</sup> =693.3821). In its <sup>13</sup>C NMR, 35 carbons were classified into 3 methyl, 9 methylene, 21 methine and 2 quaternary carbons. Methine carbons were further classified into 9 pentaene carbons and 12 deshielded carbons. All spectral data (Supplementary Table S1) agreed well with the reported data of pentamycin (= fungichromin) (6, Figure 1).<sup>17,18</sup> Pentamycin is a 28-membered pentaene macrolide with a strong antifungal activity,16 suggesting that the observed inhibitory zone against A. niger in Figure 2d was caused by the action of 6. Average yield of 6 in FS18 was 5.0 mg  $l^{-1}$ , while the relative yield in strain 51252 was estimated to be  $0.02 \text{ mg} \text{l}^{-1}$  from the calibration curve of 6. Hence, mutation in the early step of lankacidin biosynthesis showed 250-fold improvement in pentamycin production.

## Synthesis of pentamycin, citreodiol and *epi*-citreodiol was not regulated by the SRB signaling molecules

In many *Streptomyces* species, antibiotic production is strictly controlled by the small diffusible signaling molecules such as A-factor for streptomycin production in *Streptomyces griseus*<sup>19</sup> and virginia butanolides for virginiamycin production in *Streptomyces virginiae*.<sup>20</sup> Two signaling molecules, SRB1 and SRB2, were isolated from *S. rochei* and their structures of SRB1 and SRB2, were determined to be (1'*R*)2-(1'-hydroxyl-6'-oxo-8'-methylnonyl)-3-methyl-4-hydroxybut-2-en-1,4-olide and (1'*R*)2-(1'-hydroxyl-6'-oxo-8'-methyldecyl)-3-methyl-4-hydroxybut-2-en-1,4-olide, respectively.<sup>5</sup> SRB1 and SRB2 induced lankacidin and lankamycin production at 40 nM concentration.<sup>5</sup> It is noteworthy that the biosynthetic gene clusters of lankacidin and lankamycin as well as their regulatory genes including *srrX*, which encodes a synthetic enzyme for SRBs, are located on pSLA2-L in *S. rochei*.

To analyze whether the synthesis of compounds **5a**, **5b** and **6** is regulated by the SRB signaling molecules, a second mutation of *srrX* was introduced into strain FS18. The *Bam*HI site near the 5'-end of *srrX* was filled-in by Klenow enzyme (Figure 4a), which caused a frame shift in *srrX*. Plasmid pKAR3069 containing the mutated *srrX* gene on the *Streptomyces-E. coli* shuttle vector pRES18<sup>21</sup> was transformed into strain FS18 to give a *lkcA-srrX* double mutant KA78. The frame-shift



**Figure 4** Construction of the *lkcA-srrX* double mutant KA78 and analysis of its metabolites. (a) Construction of the *lkcA-srrX* double mutant KA78. Ec, *Eco*RI; Ba, *Bam*HI, Hi, *Hin*dIII; *tsr*, thiostrepton resistant gene. (b) Southern blot analysis of total DNA. Lane 1,  $\lambda/HindIII$ ; lane 2, strain FS18/*Bam*HI; lane 3, strain KA78/*Bam*HI. (c) HPLC chromatograms of metabolites from strains 51252, FS18, KA78 and KY85 (the *srrX* single mutant) monitored at 267 nm (I) and at 338 nm (II). The crude extract was applied on a COSMOSIL 5C18-MS-II column (4.6 × 250 mm) and eluted with a mixture of acetonitrile-10 mm sodium phosphate buffer (pH 8.2) (3:7, v/v) at a flow rate of 1.0 ml min<sup>-1</sup>.

mutation on *srrX* was confirmed by Southern hybridization analysis. As shown in Figure 4b, the 5.6-kb *Bam*HI and 1.9-kb *Bam*HI-*Eco*RI fragments in strain FS18 were replaced by the 7.5-kb *Bam*HI-*Eco*RI fragment in KA78. Metabolites produced by strain KA78 were analyzed by TLC, HPLC and bioassay against *A. niger* (Figures 2 and 4). This strain produced **5a**, **5b** and the antifungal agent **6** at a similar level to the single *lkcA* mutant FS18 (Figure 4c), while failed to accumulate lankacidins **1-3** or lankamycin **4** (Figure 2c). The single *srrX* mutant could not produce compounds **5a**, **5b** and **6** as well as **1-4**. This result showed that production of three compounds (**5a**, **5b**, **6**) was not regulated by the SRB signaling molecules.

#### DISCUSSION

In this study, we analyzed complex metabolic changes and the structures of citreodiol (**5a**), *epi*-citreodiol (**5b**) and pentamycin (**6**) overproduced by the *lkcA* disruptant FS18 of *S. rochei*. The LkcA protein plays an important role in both recognition of glycine as a starter unit and its condensation with a malonyl CoA extender unit in the early step of lankacidin biosynthesis.<sup>8</sup> Altering the metabolic flux of fatty acid precursors sometimes causes an enhanced production of polyketides in the producing microorganisms.<sup>22</sup> In the case of the polyether antibiotic nanchangmycin in *Streptomyces nanchangensis*, its titer was improved threefold by gene inactivation of the

uncharacterized PKS clusters.<sup>23</sup> For heterologous expression of secondary metabolite gene clusters, Komatsu et al.24 constructed versatile model host Streptomyces avermitilis SUKA17, whose chromosomal region including principal biosynthetic genes for avermectin and oligomycin was extensively deleted. Consequently, expression of the introduced oxytetracycline genes in the SUKA17 transformant was increased to yield a fivefold titer compared with the wild-type transformant. However, the overproduction of pentamycin and citreodiols in S. rochei seems different from these cases, because it occurred only in the *lkcA* mutant. None of the other genes in the *lkc* cluster, including three ketoreductase domains (lkcC-KR, lkcF-KR1 and lkcF-KR2), a C-methyltransferase gene (lkcC-MT), discrete acyltransferase (lkcD) and dehydratase (lkcB) genes, caused an overproduction when they were disrupted. These results suggest that the improved production of 5a, 5b and 6 in FS18 is not related to the cease of lankacidin production, though they are synthesized from a common precursor, malonyl CoA. The molecular mechanism of complex metabolic changes in FS18 remains to be clarified.

Citreodiol (5a) and epi-citreodiol (5b) were originally isolated from ascomycetous fungi Penicillium citreoviride B with a concomitant production of citreoviridin, a potent inhibitor of ATP synthesis and ATP hydrolysis.<sup>12</sup> Some of the secondary metabolic pathways are common in bacteria and fungi; for example, cephalosporins (a member of  $\beta$ -lactam antibiotics) are produced by both actinomycetes (Streptomyces clavuligerus) and fungi (Acremonium chrysogenum).<sup>25</sup> Biosynthetic genes for citreodiols have not been identified in P. citreoviride B or S. rochei. Comparative study will provide intriguing insights how secondary metabolite clusters have evolved among prokaryotic Streptomyces and eukaryotic fungi. Compounds 5a and **5b** were produced even by the pSLA2-L-deficient strain UV1-2,<sup>2</sup> harboring pSLA2-M and -S (data not shown). Sequence analysis revealed that pSLA2-M and -S have no structural genes for polyketide synthesis (accession numbers; AB597522 and AB905437, respectively).<sup>26</sup> Thus, the biosynthetic genes for 5a and 5b may be located on the S. rochei chromosome.

Pentamycin (6) was originally isolated from Streptomyces cellulosae ATCC12625 (as fungichromin)<sup>27</sup> and Streptomyces penticus (as pentamycin),<sup>16</sup> and its absolute configuration was determined by spectral comparison of the degradation products and partial synthesis.<sup>28,29</sup> A trace amount of pentamycin was also isolated from a lankacidinproducing strain of Streptomyces rochei var. volubilis.<sup>30</sup> Filipin III, a 14-deoxy congener of pentamycin, is the most studied pentaene antibiotic from biological<sup>31,32</sup> and biosynthetic<sup>33</sup> aspects. Its biosynthetic gene cluster (pte) in Streptomyces avermitilis spans over 80 kb and contains 13 genes.<sup>34</sup> Homologous genes with the *pte* cluster were not found on pSLA2-L, -M and -S, indicating the chromosomal location of the pentamycin gene (pem) cluster. Indeed, the pem cluster was identified in the contig sequences of strain 7434AN4, which contains 15 genes with an additional P450 monooxygenase gene pemI and a ferredoxin gene pemJ (data not shown). Further studies on the pem cluster are in progress in our laboratory.

Mutation of the *srrX* gene in strain FS18 revealed that the production of citreodiol, *epi*-citreodiol and pentamycin is not regulated by the SRB signaling molecules. In *S. rochei*, the signaling molecules/receptor system, SRBs/SrrA, strictly controls lankacidin and lankamycin production.<sup>35–37</sup> All of the genes responsible for biosynthesis and regulation of the two antibiotics are coded on the linear plasmid pSLA2-L. *Streptomyces coelicolor* A3(2) contains two signaling molecule/receptor systems, SCB1/ScbR<sup>38</sup> and methylenomycin furan (MMF)/MmfR.<sup>39</sup> The former regulates the cryptic modular PKS gene cluster (*kas*; SCO6273-SCO6288) coded on the chromosome, while

the latter does the methylenomycin biosynthetic genes coded on the linear plasmid SCP1. Hence, it is plausible that production of the three metabolites is controlled by an unidentified signaling molecule/ receptor system coded on the *S. rochei* chromosome. Together with the synergistic action of lankacidin and lankamycin to the neighboring sites on ribosome, the location of their biosynthetic clusters and regulatory genes on the linear plasmid pSLA2-L show ingenious strategies of *Streptomyces* in genome evolution and environmental adaptation.

#### MATERIALS AND METHODS

#### Strains and culture conditions

S. rochei wild-type strain 7434AN4 and strain 51252 that carries only pSLA2-L were described previously.<sup>2,7</sup> Strain FS18, a disruptant of the NRPS/PKS hybrid gene *lkcA*, was constructed previously (Supplementary Figure S1).<sup>8</sup> In strain FS18, the 1.6-kb kanamycin cassette was introduced into the *Bam*HI site (nt 33 166 of pSLA2-L) on the *lkcA* gene (nt 31 013–35 881 complement of pSLA2-L). YEME liquid medium (0.3% yeast extract, 0.5% polypeptone, 0.3% malt extract, 1% D-glucose and 34% sucrose) was used for preparation of *S. rochei* protoplasts.<sup>40</sup> Protoplasts were regenerated on R1M agar medium.<sup>41</sup> YM medium (0.4% yeast extract, 1.0% malt extract and 0.4% D-glucose, pH 7.3) was used for antibiotic production. Tryptic soy broth (TSB) medium was used for preparation of *Aspergillus niger* spore suspension and bioassay. *Escherichia coli* XL1-Blue (Agilent Technologies, Santa Clara, CA, USA) was used for construction of targeting plasmids. *E. coli* strains were grown in Luria Bertani (LB) medium supplemented with ampicillin (100 µg ml<sup>-1</sup>) for plasmid maintenance when necessary.

#### Detection and isolation of metabolites

Detailed procedures were described in Supplementary Information.

#### Construction of the *lkcA-srrX* double mutant KA78

A 1.8-kb PCR fragment containing the *srrX* gene was amplified using cosmid C7 (nt 143 101–184 445 of pSLA2-L)<sup>7</sup> as a template and a pair of primers, srrX-f1 (5'-GACGAATTCGTGAACGCGCCGGCCACCAG-3', where italics indicate an *Eco*RI site) and srrX-r1 (5'-CTGAAGCTTGTAGCCGCG CTCGTTGCGCC-3', where italics indicate a *Hin*dIII site). This fragment was digested with *Eco*RI and *Hin*dIII and cloned into pUC19 to give pKAR3067. The *Bam*HI restriction site near the 5'-end of *srrX* was eliminated by filling-in and self-ligating reactions to give pKAR3068, the vector of which was replaced by *E. coli-Streptomyces* shuttle vector pRES18<sup>21</sup> to give a targeting plasmid, pKAR3069.

This plasmid was transformed into protoplasts of *S. rochei* FS18. Thiostrepton-resistant transformants were subjected to sequential cultivation in YM liquid medium supplemented with thiostrepton  $(10 \,\mu g \, ml^{-1})$  to give the plasmid-integrated (single crossover) strain, which was then cultured in the absence of thiostrepton to give the double crossover strain. Mutation was confirmed by Southern hybridization using a DIG DNA Labeling and Detection Kit (Roche Diagnostics, Germany) according to the manufacturer's protocol.

#### Antifungal activity

Aspergillus niger was used as test microorganism. A spore suspension of *A. niger* was obtained by washing a well-grown Petri dish culture (90 $\emptyset \times 8$  mm) with a sterile aqueous solution containing 1% Tween 20 (0.5 ml). The resulting suspension was diluted with sterile water (10 ml) and filtered through a cotton plug. The bioassay plate was composed of two layers; the bottom layer contained TSB medium with 1.5% agar, while the top agar contained TSB-agar (0.8%) supplemented with 1% spore suspension of *A. niger*. Crude extracts from 100-ml broth (40 µl) were dropped onto paper disks (diameter; 0.8 cm), put on the bioassay plates and incubated at 28 °C for 5 days. The relative yield of **6** in strain 51252 was estimated by the calibration curve of inhibition zones obtained by purified **6**.

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#### ACKNOWLEDGEMENTS

We thank Professor John C. Vederas (University of Alberta) for a kind gift of natural pentamycin. We thank Mrs. Tomoko Amimoto (Natural Science Center for Basic Research and Development, Hiroshima University) for the measurement of high-resolution mass spectra. This work was supported by a Grants-in-Aid for Scientific Research on Innovative Areas from the Ministry of Education, Culture, Sports, Science and Technology of Japan (MEXT), and a Charitable Trust Araki Medical and Biochemical Research Memorial Fund.

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