

# A marine-derived *Streptomyces* sp. MS449 produces high yield of actinomycin X<sub>2</sub> and actinomycin D with potent anti-tuberculosis activity

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**Abstract** In the course of our screening program for anti-*Mycobacterium bovis bacillus* Calmette-Guérin (BCG) and anti-*Mycobacterium tuberculosis* H37Rv (MTB H37Rv) agents from our marine natural product library, a newly isolated actinomycete strain, designated as MS449, was picked out for further investigation. The strain MS449, isolated from a sediment sample collected from South China Sea, produced actinomycin X<sub>2</sub> and actinomycin D in substantial quantities, which

showed strong inhibition of BCG and MTB H37Rv. The structures of actinomycins were elucidated by nuclear magnetic resonance and mass spectrometric analysis. The strain MS449 was taxonomically characterized on the basis of morphological and phenotypic characteristics, genotypic data, and phylogenetic analysis. The 16S rRNA gene sequence of the strain was determined and a database search indicated that the strain was closely associated with the type strain of *Streptomyces avermitilis* (99.7 % 16S rRNA gene similarity). *S. avermitilis* has not been previously reported to produce actinomycins. The marine-derived strain of *Streptomyces* sp. MS449 produced notably higher quantities of actinomycin X<sub>2</sub> (1.92 mg/ml) and actinomycin D (1.77 mg/ml) than previously reported actinomycins producing strains. Thus, MS449 was considered of great potential as a new industrial producing strain of actinomycin X<sub>2</sub> and actinomycin D.

Caixia Chen, Fuhang Song, and Qian Wang contributed equally to this paper.

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## Introduction

Tuberculosis is a contagious disease, caused by *Mycobacterium tuberculosis* (MTB). MTB infection results in more than two million deaths per year and it is considered the leading cause of mortality in people co-infected with HIV/AIDS (World Health Organization 1999). About one third of the world's population has been infected with MTB and new infections occur at a rate of one individual per second (Dye et al. 1999). Tuberculosis is becoming a serious threat for disease control and its deadly synergy with HIV/AIDS and a surge in drug-resistant strains make it a primary objective for

our team in recent years to discover entirely novel agents that can markedly shorten the time to cure tuberculosis and simplify the treatment.

To date, a lot of work has been done in our lab to accelerate the lead discovery process (Song et al. 2010; Yu et al. 2010; You et al. 2010) and to improve the yields of some bioactive metabolites by biodiversity and taxonomy-guided natural product library construction (Ashforth et al. 2010), high throughput screening (Liu et al. 2010), and genetic engineering project (Zhang et al. 2005; Zhuo et al. 2010). Also, these remarkable efforts facilitated our screening process for anti-*Mycobacterium bovis bacillus* Calmette-Guérin (BCG) and anti-*M. tuberculosis* H37Rv (MTB H37Rv) compounds and enhanced the efficiency of the discovery of bioactive agents. The construction of a large marine natural product library (MNPL) (Bian et al. 2008) afforded a strong support for all screening work based on different high throughput screening models, such as synergic anti-fungal screening model (Zhang et al. 2007), anti-methicillin-resistant *Staphylococcus aureus*, anti-BCG, and anti-fungal screening models, and some other antimicrobial or antitumor screening models at the cellular and molecular levels.

We have developed a convenient high throughput screening model to identify inhibitors in an aerobic, logarithmic growth screen of BCG (Cowley and Av-Gay 2001). The BCG used here was transformed with green fluorescent protein (*GFP*) constitutive expression plasmid Puv3583 with direct readout of fluorescence as a measure of bacterial growth. In our screening for anti-BCG compounds from our MNPL, a number of positive strains were picked out and their secondary metabolites were analyzed by HPLC-diode array detector. These organisms proved to be a rich source of bioactive compounds. Strain MS449, which was isolated from a sediment sample collected from South China Sea, gained our special interest because of the potential activity against BCG and diversity of metabolites in its fermentation broth. The 16S rRNA gene sequence from the strain was determined and proved to have high identity (99.7 %) with *Streptomyces avermitilis*. As the type *S. avermitilis* strain which produces avermectins was available, we compared its features with MS449 and analyzed the HPLC-UV-Vis characteristic of its secondary metabolites and found that it evidently differed from the strain MS449 in morphological and metabolic characteristics. Therefore, MS449 was picked out for further investigation and resulted in three actinomycins with potent bioactivities.

Actinomycins are a family of chromopeptide lactone antibiotics, among which actinomycin D has been studied extensively and widely used clinically for the treatment of several types of malignant tumors, notably Wilms' tumor (Green 1997) and childhood rhabdomyosarcoma (Womer 1997). Since the discovery of actinomycin D, it has become an important tool in clinical, molecular, and cell biology

(Kurosawa et al. 2006) because of its unique structure and biological properties by intercalating into duplex DNA, which results in inhibition of DNA-dependent RNA polymerase and thus protein synthesis (Wadkins et al. 1998). It was reported to inhibit the coxsackievirus B3 (Sajjats et al. 2003) and was proposed for the treatment of AIDS because of its effectiveness in inhibiting the minus-strand transfer process in HIV-1 (Rill and Hecker 1996). By contrast, the anti-tuberculosis activity of actinomycins was not paid much attention, and little research has been reported. The literature review indicated that subsequent research mainly focused on the structure modification and biofunction of actinomycins with no more attention to actinomycin X<sub>2</sub> and X<sub>0β</sub> especially in the anti-tuberculosis aspect (Sun and Yang 2011). The strain with high production of actinomycin X<sub>2</sub>, actinomycin D, and relatively less actinomycin X<sub>0β</sub> may be valuable for researchers' investigation.

In this paper, we describe the marine-derived *Streptomyces* strain MS449, which exhibits potent activities against BCG and MTB H37Rv and is a stable high yield producer of actinomycin X<sub>2</sub>, actinomycin D, and actinomycin X<sub>0β</sub> even under non-optimized culture conditions. The strain MS449 was identified as *S. avermitilis* based on taxonomic experiments and it is a novel source for actinomycins.

## Materials and methods

### Microbial strains

An actinomycete strain MS449, used as the producing strain, was isolated from a sediment sample collected from South China Sea (longitude 110:22:47.058 and latitude 17:33:35.661) and grown on an GT medium agar slant consisting of soluble starch 2.0 %, L-asparagine 0.05 %, KNO<sub>3</sub> 0.1 %, K<sub>2</sub>HPO<sub>4</sub>·H<sub>2</sub>O 0.05 %, NaCl 0.05 %, and MgSO<sub>4</sub>·7H<sub>2</sub>O 0.05 % (pH 7.5) at 28 °C. MS449 was deposited at the China General Microbiological Culture Collection Center (CGMCC) with the accession number CGMCC No. 5452.

*Mycobacterium phlei* (ATCC 11758), used as an indicator organism during the isolation process, was grown on an agar plate consisting of peptone 1.0 %, malt extract 0.5 %, yeast extract 0.5 %, casein acid hydrolysate 0.5 %, beef extract 0.2 %, glycerol 0.2 %, Tween-80 0.005 %, MgSO<sub>4</sub>·7H<sub>2</sub>O 0.1 %, and agar 1.2 %, pH 7.2 at 37 °C. *M. bovis* BCG (Pasteur 1173P2) and *Mycobacterium tuberculosis* H37Rv (ATCC27294) were used here for bioactivity screening and assays.

*Enterococcus faecalis* HH22 (Shiojima et al. 1997), carrying bifunctional modifying enzyme genes in plasmid *PBE10*, was clinically isolated and resistant to gentamycin, penicillin, erythromycin, and tetracycline.

*Proteus mirabilis* PM631 SE (Neuwirth et al. 2001) was a clinical isolate producing TEM-3  $\beta$ -lactamase. *E. faecalis* HH22 and PM631 SE were preserved in the Institute of Medicinal Biotechnology, Chinese Academy of Medical Sciences & Peking Union Medical College. *E. faecalis* HH22 and PM631 SE, together with *Pseudomonas aeruginosa* (ATCC 27853), *Escherichia coli* (ATCC 25922), *S. aureus* (ATCC 29213), *Mycobacterium smegmatis* (ATCC 700044), and *M. phlei* (ATCC 11758), were used to test antimicrobial properties of the fermentation broth using paper disk-agar diffusion assay (Raahave 1974; Fosto et al. 2008).

#### Cultural and morphological properties of strain MS449

A set of cultural and phenotypic characteristics was examined using media and the International *Streptomyces* Project procedures recommended by Shirling and Gottlieb (1966). Mature aerial mycelium and substrate mycelium pigmentation were recorded on GT agar media following incubation at 28 °C for 28 days. Morphological features of spores and mycelia were observed by scanning electron microscopy (Quanta 200) after growth on GT, desiccation, and coating the cells with gold.

#### 16S rRNA gene sequence and phylogenetic analysis

Extraction of MS449 genomic DNA was done as described by the TINAamp Bacteria DNA Kit. Universal primers (27f: 5'-GAGAGTTTGATCCTGGCTCAG-3'; 1492r: 5'-CTACGGCTACCTGTTACGA-3') were used to amplify the 16S rDNA. PCR amplification [25  $\mu$ l final volume—0.4  $\mu$ l 20  $\mu$ M of each primer, 2.5  $\mu$ l 10 $\times$  buffer (TaKaRa, Dalian, China), 2.5  $\mu$ l 2.5 nM dNTP (TaKaRa), 2 U rTap polymerase (TaKaRa), and 1  $\mu$ l DNA template] of the 16S rDNA was performed on TaKaRa PCR Thermal Cycler with the initial denaturation at 94 °C for 5 min, 30 cycles of denaturation (94 °C, 1 min), annealing (55 °C, 1 min), and elongation (72 °C, 1 min 15 s), and a final elongation at 72 °C for 10 min. Multiple alignments with sequences of most closely related *Streptomyces* and calculations of levels of sequence similarity were carried out using CLUSTAL W (Thompson et al. 1994). The phylogenetic tree was constructed using the neighbor-joining method (Saitou and Nei 1987) by MEGA 4.0 (Tamura et al. 2007). The topology of phylogenetic tree was evaluated by bootstrap resampling method with 1,000 replicates (Felsenstein 1985).

#### Nucleotide sequence accession number

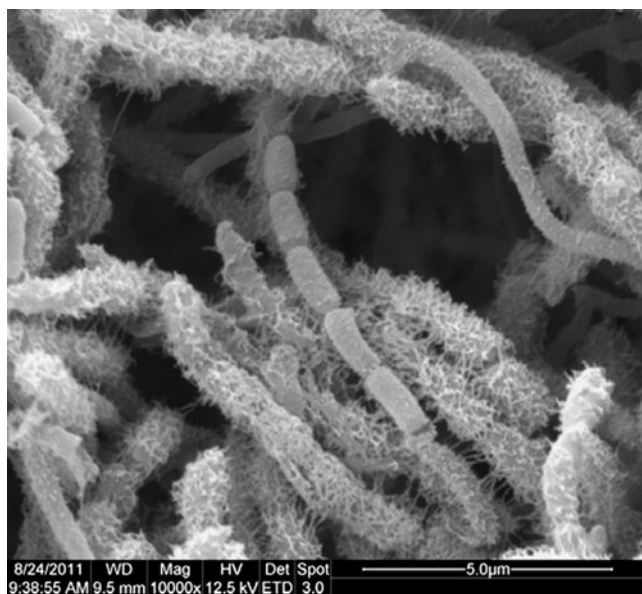
The nucleotide sequence of 16S rRNA gene reported in this article was assigned to the GenBank accession number JN104388.

#### Production of actinomycins by MS449 in different producing media

A stock culture of the strain was grown and maintained on a GT agar slant. The stock culture was transferred into 250-ml Erlenmeyer flasks containing 40 ml of the seed medium with the same components as the agar slant medium. The culture was incubated on a rotary shaker (220 rpm) at 28 °C for 96 h. Four milliliters of the seed culture was transferred into 250-ml Erlenmeyer flasks containing 40 ml of the producing medium. Three kinds of producing media were used: (a) M21 medium consisting of glucose 0.5 %, lactose 4 %, cottonseed protein 3 %, Bacto Peptone 0.5 %,  $K_2HPO_4$  0.05 %,  $MgSO_4 \cdot 7H_2O$  0.05 %, and KCl 0.03 % (pH 7.0); (b) 9A medium consisting of yeast extract 1.0 %, saccharose 5.0 %,  $K_2HPO_4$  1.0 %,  $NaNO_3$  0.3 %, KCl 0.05 %,  $MgSO_4$  0.05 %, and  $FeSO_4$  0.001 % (pH 6.5); and (c) MPG medium consisting of glucose 1.0 %, millet meal 2.0 %, cotton seed gluten meal 2.0 %, and MOPS 2.0 %, pH 7.0. The cultures were incubated at 28 °C for 17 days on a rotary shaker at 220 rpm. After incubation for 5 days, aliquots (2 ml) of the cultures were collected every 48 h and prepared for HPLC analysis as follows: After centrifugation (8,000 rpm, 3 min), the mycelia were extracted twice with acetone to give the organic fraction. The supernatants were added 0.5 ml HP-20 resins, rocked for 2 h on a shaker, and filtrated, and the resins were washed with acetone three times. The acetone fractions from the mycelia and the supernatants were combined and dried to give crude extracts. Each crude extract was then redissolved in 2 ml acetone, filtered through a 0.45- $\mu$ m membrane filter, and analyzed by RP-HPLC equipped with an Agilent XDB-C8, 5  $\mu$ m column (4.6 $\times$ 150 mm), and 70 % methanol in water as the mobile phase. External standard method was used in the quantitative analysis with actinomycin X<sub>2</sub> (Alexis), actinomycin D (Sigma), and actinomycin X<sub>0 $\beta$</sub>  (Alexis) as standards and 446 nm as the detection wavelength.



**Fig. 1** Colony characteristics of MS449 grown on GT agar at 28 °C for 28 days



**Fig. 2** Scanning electron micrograph of MS449 grown on GT agar at 28 °C for 28 days. Bar, 5 μm

#### Susceptibility test of fermentation broth with MPG medium

*P. aeruginosa*, *E. coli*, *S. aureus*, *M. smegmatis*, *M. phlei*, *E. faecalis* HH22, and PM631 SE were grown for 24 h at 37 °C in LB medium; 1 % (v/v) microorganism cultures were mixed with sterilized LB agar medium to make uniform microorganism suspensions for pouring plates before the medium solidified. Six-millimeter filter paper disks impregnated with testing samples were allowed to dry and then placed on the solidified agar plates and incubated at 37 °C overnight. The area of no growth around the disk was known as a “zone of inhibition”.

#### Purification and characterization of actinomycins

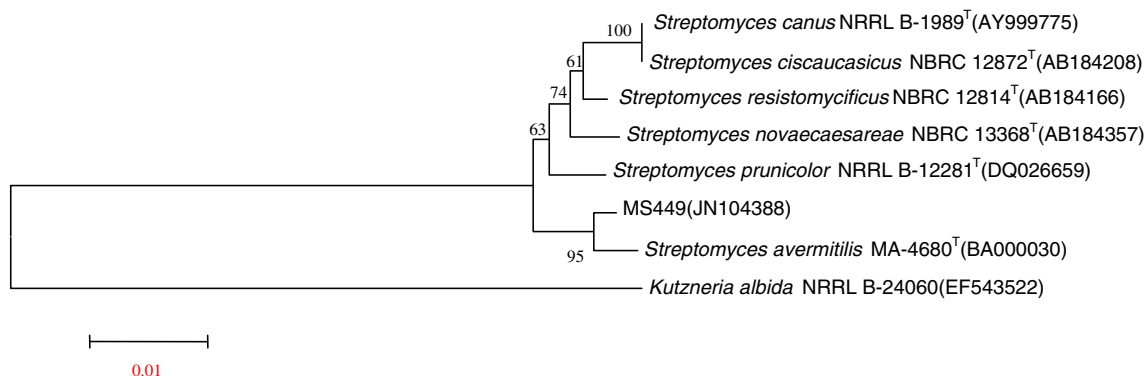
The fermentation broth of MS449 grown in MPG medium (200 ml) at 28 °C for 15 days was fractionated by centrifugation

to give the supernatant and mycelium. The mycelium was soaked with acetone and filtered. The filtrate was diluted to five times volume with the supernatant or water and then the mixture was charged on a column of HP-20 resins, which was washed with distilled water and then eluted with 50, 65, and 80 % acetone, respectively. The active fractions eluted with 65 % acetone were combined and removed acetone in vacuo, and then the water residue was extracted twice with ethyl acetate. The organic fractions were combined, dried (over Na<sub>2</sub>SO<sub>4</sub>), and concentrated under reduced pressure. The residue was then resuspended in acetone and gave a pure active constituent (**1**) using preparative RP-HPLC equipped with an Eclipse XDB-C18, 5 μm column (9.4×250 mm, Agilent), and 70 % MeOH in water as the mobile phase. The active fractions eluted with 80 % acetone from HP-20 resins were also separated into pure constituents using RP-HPLC and gave two pure compounds (**2** and **3**). Antibiotic activity of the culture broth was determined and the bioactive fractions were tracked by conventional paper disk-agar diffusion assay using *M. phlei*, and the zones of inhibition were measured after 18 h of incubation at 37 °C.

The actinomycins were characterized using spectroscopic analyses. Electrospray ionization mass spectra (ESI-MS) were recorded on a Bruker Esquire<sup>3000</sup> plus spectrometer and various nuclear magnetic resonance (NMR) spectra were recorded on a Varian Inova 600 MHz spectrometer and the solvent was acetone-*d*<sub>6</sub>.

#### Anti-BCG and MTB assay

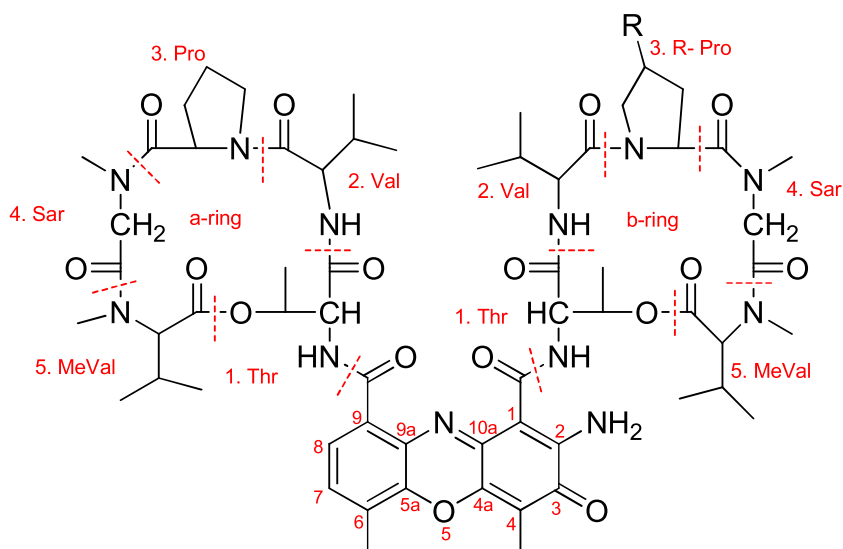
The anti-BCG and anti-MTB H37Rv assays utilize constitutive GFP expression (pUV3583c-GFP) with direct readout of fluorescence as a measure of bacterial growth (Cowley and Av-Gay 2001). The in vitro activity of compounds against BCG and MTB H37Rv was determined in a 96-well plate as previously described (Wang et al. 2010). BCG and MTB H37Rv were grown at 37 °C to mid-log phase in Middlebrook 7H9



**Fig. 3** Neighbor-joining phylogenetic tree of strain MS449 based on 16S rRNA gene sequence generated by Mega4.0. Numbers at nodes indicate levels of bootstrap support (percent) based on a neighbor-joining analysis

of 1,000 resampled datasets; only values >50 % are given. NCBI accession numbers are given in parentheses. Bar, 0.01 nucleotide substitutions per site. Also, *Kutzneria albida* was chosen as the outgroup



**Fig. 4** The structures of isolated compounds (1–3)

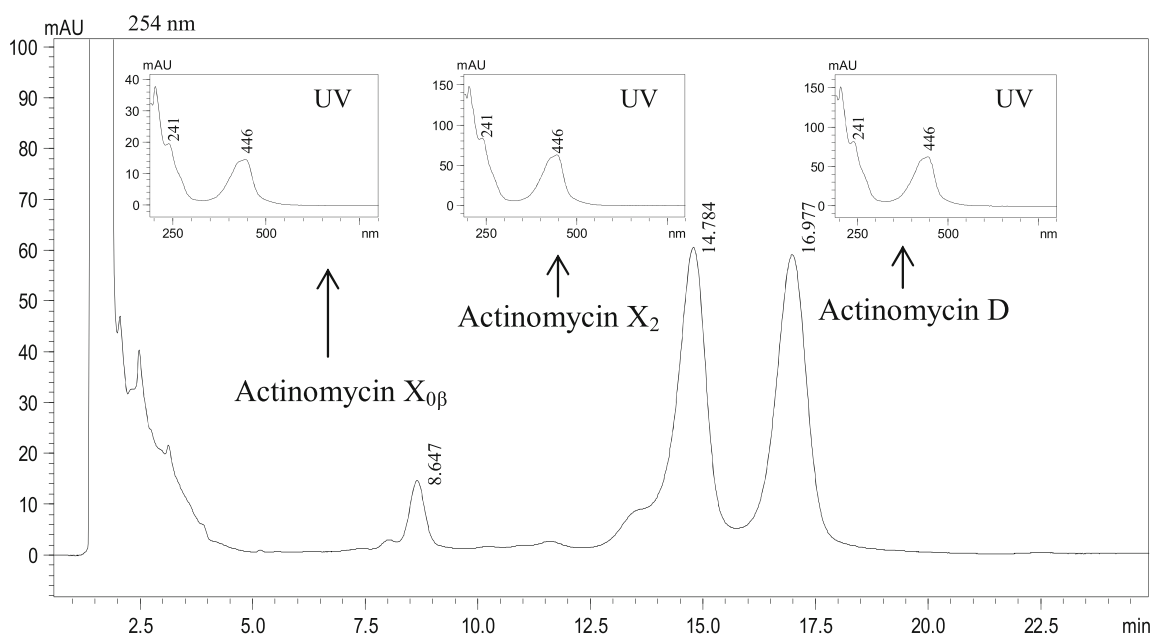
**1** (actinomycin X<sub>0β</sub>)    R= OH

**2** (actinomycin X<sub>2</sub>)    R= O

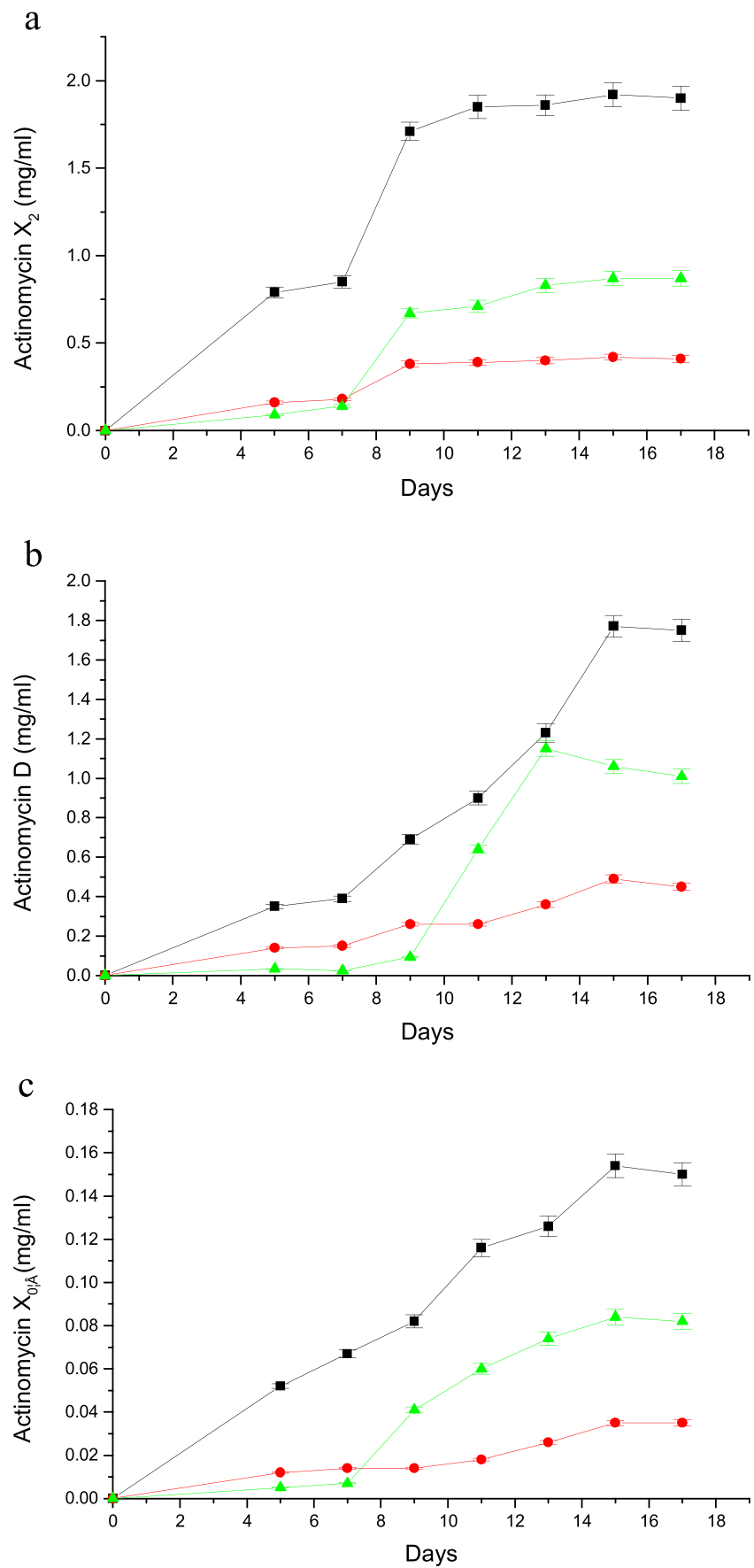
**3** (actinomycin D)    R= H

broth supplemented with 10 % OADC enrichment (Becton Dickinson), 0.05 % Tween-80, and 0.2 % glycerol. The cultures were then diluted to a bacterial suspension with OD<sub>600</sub> values of 0.025 with culture media. For primary screening,

aliquots (80 μl) of the bacterial suspension were added to each well of the clear flat-bottom 96-well microplates, followed by adding 2 μl of sample solutions (1 mg/ml). Isoniazid served as the positive control while DMSO served as the blank control.

**Fig. 5** HPLC analysis of crude extracts of MS449 cultured in MPG medium. Extracts were analyzed using Shimadzu 20A HPLC with an Agilent XDB-C8 column (5 μm, 4.6×150 mm), 70 % methanol as the mobile phase, 1.0 ml/min as the flow rate, and 254 nm as the detection wavelength

**Fig. 6** Time course production of actinomycin X<sub>2</sub> (a), actinomycin D (b), and actinomycin X<sub>0A</sub> (c) by MS449 cultured in MPG (square), M21 (circle), and 9A (triangle) media. Results represent the means  $\pm$  standard errors of the means (SEM) of three independent experiments



**Table 1** Susceptibility of the fermentation broth of MS449 to different microorganisms

Microorganisms	Diameters of inhibition zones (mm)
<i>Pseudomonas aeruginosa</i>	–
<i>Escherichia coli</i>	30.0
<i>Staphylococcus aureus</i>	17.0
<i>Mycobacterium smegmatis</i>	19.0
<i>Mycobacterium phlei</i>	20.0
<i>E. faecalis</i> HH22	17.5
PM631 SE	–
Negative control	–

– no activity was detected

The plates incubated at 37 °C for 3 days for BCG or 10 days for MTB H37Rv GFP fluorescence were measured with a multi-label plate reader (Perkin-Elmer Envision 2103) using the bottom read mode, with excitation at 485 nm and emission at 535 nm. Hits were defined as compounds that could inhibit more than 90 % of bacterial growth compared to the blank control at a concentration of 64 µg/ml or less.

For MIC determination of the compounds, bacterial suspensions were prepared and dispensed as described above, and compounds were serially diluted twofold from 25 to 0.19 µg/ml in each column. Isoniazid as positive control (800 to 6.25 ng/ml) and DMSO as blank control were also included in each plate. The plate was incubated at 37 °C for an appropriate time, and after incubation, the plate was bottom-read to record the GFP fluorescence. Here, MIC is defined as the minimum concentration of drug that inhibits more than 90 % of bacteria growth reflecting by fluorescence values. Anti-BCG and anti-MTB H37Rv assays were performed in Biosafety level (BSL) 2 and BSL3 settings, respectively.

## Results

### Characterization and identification of isolated strain MS449

A number of actinomycete strains were isolated from a sediment sample collected from South China Sea and investigated with small-scale fermentation. The strains with their crude extracts were deposited in our MNPL and evaluated for their ability to produce antibiotics by high throughput screening. Strain MS449 showed strong anti-BCG activity and was picked out to be further investigated. The colonial (Fig. 1) and morphological properties (Fig. 2) suggested that MS449 was a member of the genus *Streptomyces*. Substrate mycelium is yellow in color. Aerial mycelium, where produced, on medium such as GT is white to yellowish white with spinous surfaces. Soluble yellow pigments are produced. Aerial hyphae differentiate into short, straight chains of smooth-surfaced, cylindrical-

like spores (0.5–0.7 µm×1.5–2.0 µm) after 28 days of cultivation on GT plates.

The comparison between the 16S rRNA gene sequence of strain MS449 (1448 nt) and sequence in GenBank indicated that the novel isolate was closely related to the genus *Streptomyces* and has the highest similarity with *S. avermitilis* (99.70 %). Phylogenetic analysis based on 16S rRNA gene sequence analysis revealed that strain MS449 formed a distinct phylogenetic cluster with *S. avermitilis* in the phylogenetic tree (Fig. 3) with a bootstrap value above 95 %.

### Production of actinomycins by MS449 in different producing media

Actinomycin X<sub>0β</sub>, actinomycin X<sub>2</sub>, and actinomycin D (**1–3**) (Fig. 4) were detected in three crude extracts and the three major components were eluted by HPLC at 8.6, 14.7, and 16.9 min, respectively (Fig. 5). The relative proportion of the three identified antibiotics remained consistent in the three media. The highest actinomycin productivity was observed in MPG medium at 15 days (Fig. 6). The yields of actinomycin X<sub>0β</sub>, actinomycin X<sub>2</sub>, and actinomycin D from MPG medium were 0.15, 1.92, and 1.77 mg/ml, respectively.

### Purification and characterization of actinomycins produced by MS449

Active components were isolated from MPG medium culture broth (200 ml) and three bioactive compounds were obtained (**1–3**). Their structures were elucidated by UV, 1D NMR, 2D NMR (<sup>1</sup>H-<sup>1</sup>H COSY, <sup>1</sup>H-<sup>13</sup>C HSQC, <sup>1</sup>H-<sup>13</sup>C HMBC), and MS analysis as well as by comparison with the previously reported NMR data of actinomycins. All isolated compounds showed typical UV–Vis spectra with maximal absorbance at 241 nm (shoulder) and 446 nm similar to those of known actinomycins (Fig. 5). ESI-MS of isolated compounds revealed molecular ion peaks at *m/z* 1,271.3 [M+H]<sup>+</sup> for **1**, 1,269.1 [M+H]<sup>+</sup> and 1,291.1 [M+Na]<sup>+</sup> for **2**, and 1,255.1 [M+H]<sup>+</sup> and 1,277.1 [M+Na]<sup>+</sup> for **3** (Supplementary materials, Fig. S1), which were identical to those of actinomycin X<sub>0β</sub>, actinomycin X<sub>2</sub>, and actinomycin D, respectively. <sup>1</sup>H, <sup>13</sup>C (Supplementary materials, Table S1) and 2D NMR spectra of isolated compounds in acetone-*d*<sub>6</sub> showed a great similarity. From 1D, 2D NMR, and by comparison with the previously reported data (Yu and Tseng. 1992; Zhang et al. 2009; Lackner et al. 2000), the structures of **1**, **2**, and **3** were confirmed to be actinomycin X<sub>0β</sub>, actinomycin X<sub>2</sub>, and actinomycin D (Fig. 4).

### Susceptibility test of fermentation broth

Susceptibility of fermentation broth of MS449 in MPG medium against a variety of microorganisms was performed by

paper disk diffusion assay. The relevant comparisons between the zones of inhibition are shown in Table 1. The fermentation broth showed no activity against *P. aeruginosa* and PM631 SE. According to the test results, we choose one of the positive microorganisms, *M. phlei*, which also belongs to the genus *Mycobacterium*, as the indicator organism (Supplementary materials, Fig. S2) to track the bioactive components during the isolation process.

#### Anti-BCG and anti-tuberculosis activities of actinomycins

The pure compounds 1–3 all showed activities against BCG with MIC values of 0.25–0.5 µg/ml and MTB H37Rv with MIC values of 8.0, 1.0, and 8.0 µg/ml, respectively. Actinomycin X<sub>2</sub> showed the best activity against tuberculosis among these three compounds.

#### Discussion

MS449 is the first reported naturally occurring strain of *S. avermitilis* producing a notably high yield of actinomycin X<sub>2</sub> and actinomycin D. To the best of our knowledge, approximately 30 species of *Streptomyces* and *Micromonospora* were reported capable of producing actinomycins (Kurosawa et al. 2006; Praveen and Tripathi 2009). However, only a few strains were reported to produce relatively large quantities of one major component of actinomycins, such as *Streptomyces parvulus* (0.152 mg/ml actinomycin D under optimized conditions) (Praveen et al. 2008a), *Streptomyces griseoruber* (0.21 mg/ml actinomycin D) (Praveen and Tripathi 2009), a mutant strain of *Streptomyces sindensis* (0.85 mg/ml actinomycin D) (Praveen et al. 2008b), *Streptomyces* MITKK-103 (0.11 mg/ml actinomycin X<sub>2</sub>) (Kurosawa et al. 2006), *Streptomyces* spp. JAU4234 (0.62 mg/ml actinomycin X<sub>2</sub> under optimized conditions) (Xiong et al. 2008), and *Streptomyces nasri* strain YG62 (0.15 mg/ml actinomycin X<sub>2</sub>) (EI-Naggar 1998), whereas strain MS449 exhibited a unique ability to produce large quantities of actinomycin X<sub>2</sub> (2), actinomycin D (3), and actinomycin X<sub>0β</sub> (1) simultaneously with a production of 1.92, 1.77, and 0.15 mg/ml, respectively, under non-optimized conditions. Actinomycin X<sub>2</sub> production in MPG medium reached a sub-maximum peak after 9 days and increased slowly afterwards. The production of actinomycin D and actinomycin X<sub>0β</sub> in MPG increased markedly after 11 days. Maximum levels of actinomycin productivity were reached after 15 days (Fig. 6). Optimization of the culture conditions was expected to be done to exploit the full potential for industrial application of this strain.

Actinomycins, which were firstly discovered by Waksman and Tishler (1942), represent a family of chromopeptide antibiotics which differ solely in the peptide

portions of the molecules. To date, more than 30 analogues have been described. Actinomycins have become an important tool in molecular and cellular biology. Among actinomycins, actinomycin D has been extensively studied and used clinically. The exciting potentialities of actinomycin D have stimulated a great deal of research into its chemical, physical, and biological characteristics (Hollstein 1974; Henry 1985). Although little research about actinomycin X<sub>2</sub> and actinomycin X<sub>0β</sub> was reported, actinomycin X<sub>2</sub> showed better activity against MTB H37Rv than actinomycin D and actinomycin X<sub>0β</sub> in our research. It was also reported that actinomycin X<sub>2</sub> showed higher cytotoxicity toward HL-60 cells than actinomycin D (Kurosawa et al. 2006). It is affirmative that actinomycin X<sub>2</sub> will be more and more attractive. As a new source of actinomycin X<sub>2</sub>, MS449 is considered of great potential for industrial application.

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