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# Targeted isolation of sulfur-containing metabolites from *Lsr2*-deletion mutant strain of *Streptomyces roseosporus*†

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Deletion of the *Lsr2* gene in *Streptomyces roseosporus* up-regulated silent gene clusters and produced new secondary metabolites. An ultra-performance liquid chromatography quadrupole time of flight mass spectrometry (UPLC-QTOF-MS/MS) method was used to analyze metabolites of the mutant and wild-type strains, and recognize previously unreported sulfur-containing compounds based on their molecular formulas and fragmentation ions. The targeted isolation of unidentified compounds afforded six new sulfur-containing compounds, pyrismycins A–F (1–6), together with seven known analogues 7–13. Their cytotoxic effects were evaluated using four clinically relevant human cancer cell lines, gastric carcinoma SGC7901, breast carcinoma MDA-MB-231, lung carcinoma A549 and hepatocellular carcinoma HepG2. Compound 7 exhibited the most potent cytotoxicity with IC<sub>50</sub> values of 1.7, 5.8 and 6.3 μM against the SGC7901, HepG2 and MDA-MB-231, respectively.

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## Introduction

*Streptomyces* produce a wealth of chemically diverse secondary metabolites with a wide range of biological activities. These have been used as antibacterial, antifungal, anticancer, anti-parasitic and immunosuppressive agents.<sup>1</sup> Sulfur-containing metabolites are common across the *Streptomyces* kingdom and constitute a family of natural products which play an important role in the health care for humans.<sup>2,3</sup> Among the top 200 best-selling drugs in the current drug market, about 86% of them are sulfur and/or nitrogen containing compounds.<sup>4</sup>

Microbial genomics and bioinformatics analysis has revealed that many microorganisms have far greater potential to produce specialized metabolites than have been discovered by classic screening-based methods. Bioinformatics analysis of the genome of *S. roseosporus* disclosed 25 gene clusters, which include 14 NRPS or PKSs, three for terpene

biosynthesis, three for siderophore, two for bacteriocin, one ectoine, and two other gene clusters.<sup>5</sup> Relative to the limited number of secondary metabolites isolated from *S. roseosporus*, many gene clusters in this bacteria seemingly are not

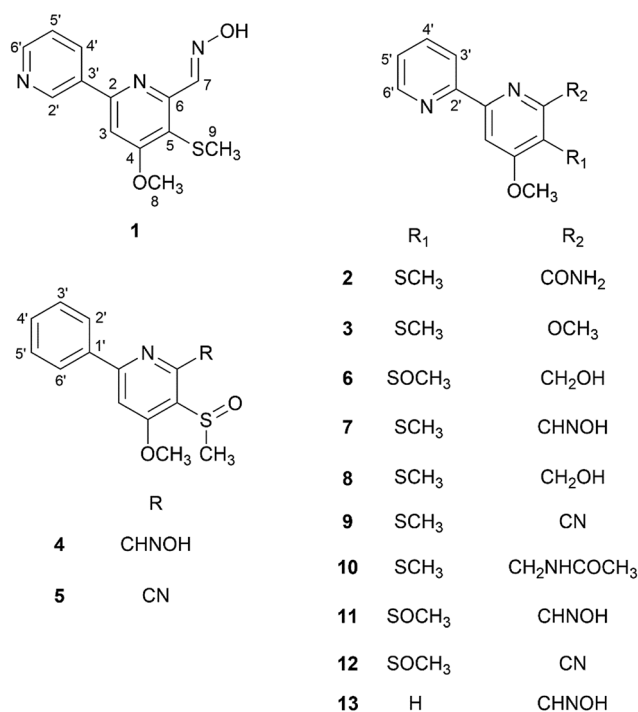


Fig. 1 Chemical structures of compounds 1–13.

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† Electronic supplementary information (ESI) available: ESI Experimental section; Table S1 (sulfur-containing metabolites detected and characterized in  $\Delta$ *SrLsr2* strain by UPLC-QTOF-MS and HR-MS/MS analysis); Table S2 (cytotoxic activities of 1–13 isolated from the  $\Delta$ *SrLsr2* strain of *S. roseosporus*); Fig. S1 (HR-MS/MS spectra and proposed fragmentation pathway of peaks 17 and 18 (1 and 2)); Fig. S2 (HPLC guided isolation of sulfur-containing metabolites); Fig. S3 (X-ray crystal structure of 10); NMR and HRMS spectra of compounds 1–6. CCDC 1533762. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7ra06482a



expressed, implying that there is additional metabolic potential in this species.

A number of approaches have been developed to activate and up-regulate silent or cryptic gene clusters to produce new compounds.<sup>6,7</sup> *Lsr2* is a small nucleoid-associated protein present in *S. roseosporus* that regulates the expression of many biosynthetic gene cluster and influences the organization of bacterial chromatin, but examples of influence the metabolites are less common.<sup>8,9</sup> Thus, we speculated that deletion of the *Lsr2* in *S. roseosporus* might activate or repress gene expression and produce novel compounds.

Herein, we report the isolation and structure elucidation of six new sulfur-containing compounds, pyrismycins A–F (1–6), with seven known analogues (7–13), from the *Lsr2*-deletion ( $\Delta Srlsr2$ ) mutant strain of *S. roseosporus*. In this research, UPLC-QTOF-MS/MS, a successful tool for dereplication purposes,<sup>10</sup> was used to identify sulfur-containing metabolites from the  $\Delta Srlsr2$  strain, with present in the wild-type (WT) strain, and then used to guide the isolation work. The cytotoxic effects of isolated compounds on four clinically relevant human cancer cell lines, SGC7901 gastric carcinoma, MDA-MB-231 breast carcinoma, A549 lung carcinoma and HepG2 hepatocellular carcinoma, were evaluated (Fig. 1).

## Results and discussion

UPLC-QTOF-MS analysis revealed 22 new peaks in the ethyl acetate extract of  $\Delta Srlsr2$  than were present in a similar extract of the WT strain (Fig. 2). This result indicated that deletion of the *Lsr2* gene increased the number of secondary metabolites produced by this mutant strain. 14 peaks were identified as potential sulfur-containing derivatives, as supported by the molecular formula and fragment ions, and compared with the *Streptomyces* compounds previously indexed in the SciFinder database (Table S1†).

Peak 9 ( $t_R = 13.75$  min) showed molecular ions at  $m/z$  291.0794  $[M + H]^+$  and its molecular formula was determined to

be  $C_{14}H_{14}N_2O_3S$  (Fig. 3). Tandem mass showed that the fragmentation of  $[M + H]^+$  precursor ion generated  $[M - OH]^+$  ion ( $m/z$  273.0695),  $[M - OH - CH_2]^+$  ion ( $m/z$  259.0539),  $[M - OH - OCH_3]^+$  ion ( $m/z$  243.0593) and fragment at  $m/z$  229.0446. The fragment ion  $m/z$  229.0446 further fragmented into fragment ion  $m/z$  204.0485 and 181.0759. The transition from fragment ion  $m/z$  156.0802 to 130.0660 indicated that it was a phenylpyridine derivative.<sup>11,12</sup> Such a molecular formula and the basic structure units have not been found previously in *Streptomyces* metabolites. Similar fragmentation was also observed for peak 15. Thus, both peaks 9 and 15 were targeted as potentially new metabolites.

Peak 17 ( $t_R = 20.31$  min), peak 18 ( $t_R = 20.43$  min) and peak 19 ( $t_R = 21.44$  min) shared a same molecular formula of  $C_{13}H_{14}N_3O_2S$  with nearly same molecular ions at  $m/z$  276.0797, 276.0794, and 276.0795  $[M + H]^+$ . They could be identified as isomeric compounds since their fragment pattern were different (Table S1†). The MS/MS spectra of peaks 18 and 17 are shown in Fig. S1,† with the molecular ions and the main fragment ions obtained under the high energy mode. The MS/MS spectra of peak 17 and peak 19 were nearly the same. The neutral loss of 33, 31, 28 and 26 mass units could account for the elimination of  $-SH$ ,  $-OCH_3$ ,  $-CO$  and  $-CN$ , respectively. The product ions at  $m/z$  157.0761 and 130.0652 were attributed to the characteristic fragment of dipyrindyl derivatives,<sup>12</sup> and the mass difference between peaks 17 and 18 was  $m/z$  182.0712 and 185.0707 units for cyano-substituted dipyrindyl and aldehyde-substituted dipyrindyl, respectively, suggesting the existence of new sulfur-containing compounds.<sup>12</sup>

The similar investigation was applied to all 22 new peaks. As a result, eight target peaks (5, 9, 15, 16, 17, 18, 20a and 21) were considered as unknowns of interest. Combined with the HPLC analysis on the subfractions (Fig. S2†) of every separation process, the extract was separated by column chromatography and semi-preparative HPLC to obtain six new sulfur-containing compounds, pyrismycins A–F (1–6), together with seven known

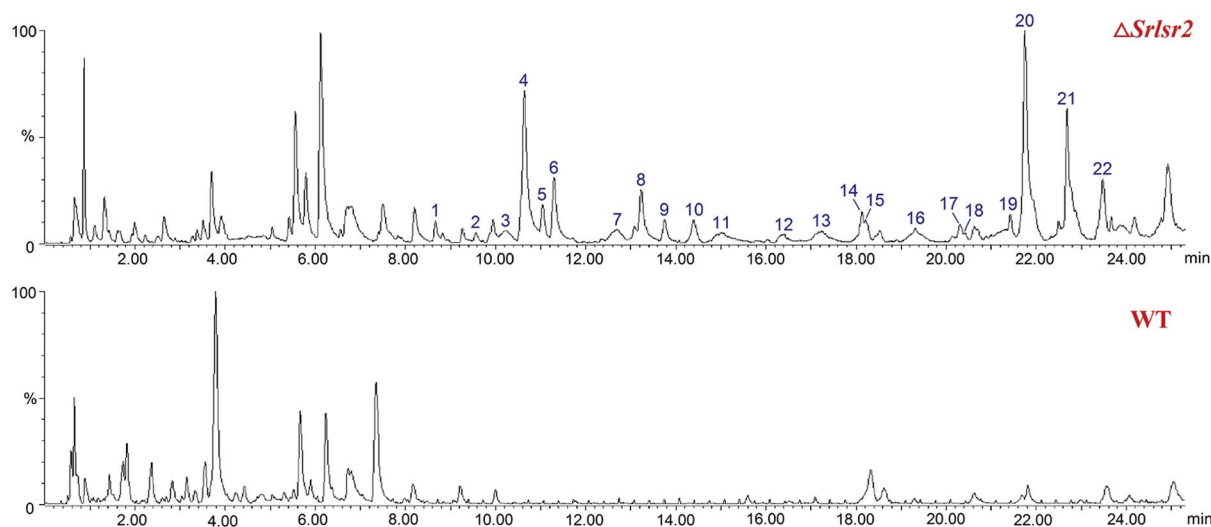


Fig. 2 UPLC-QTOF-MS profiles of the acetate extracts from  $\Delta Srlsr2$  (the upper diagram) and WT (the lower diagram) strains of *S. roseosporus*.



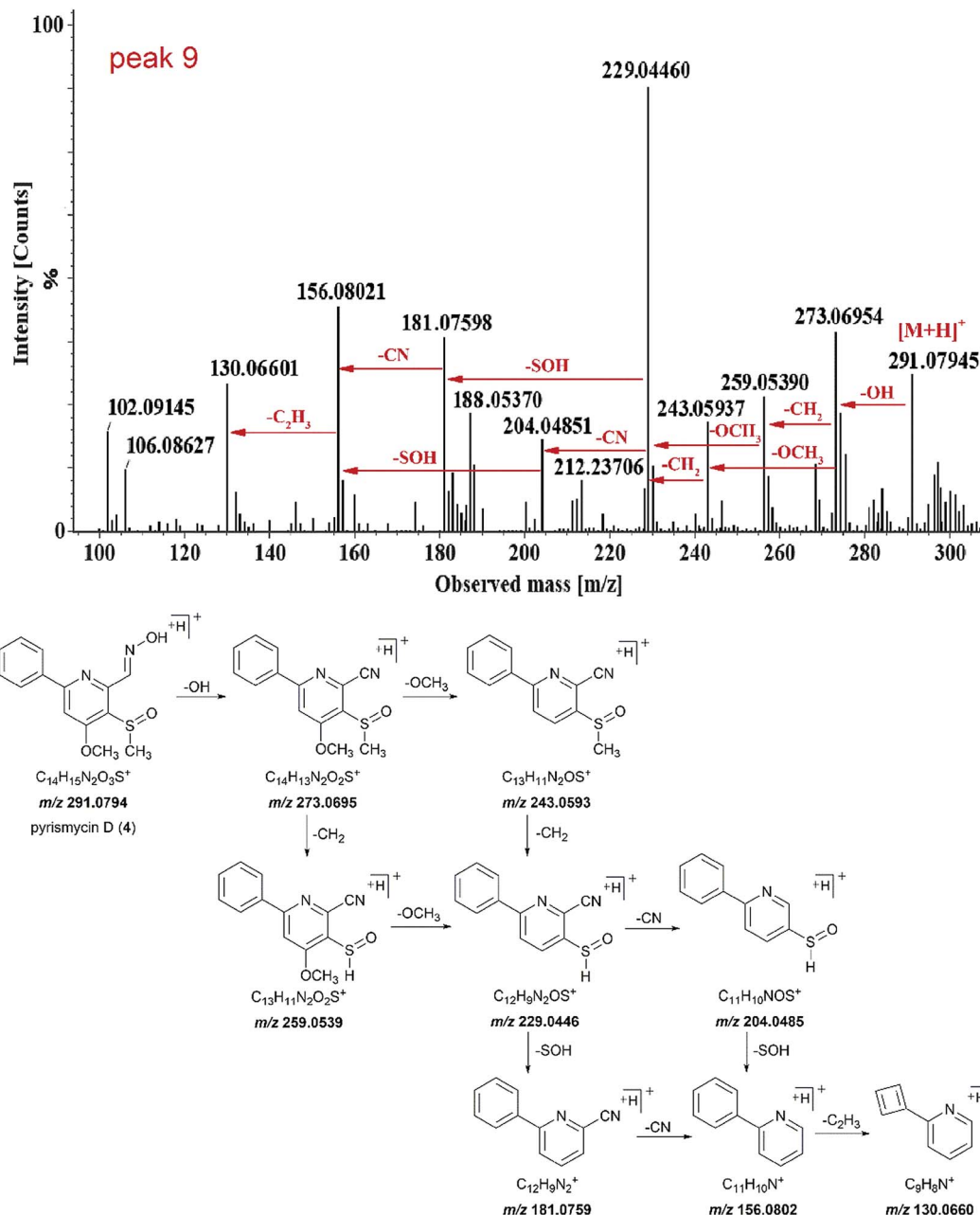


Fig. 3 HR-MS/MS spectra and proposed fragmentation pathway of peak 9 (4).

analogues 7–13. Their structures were elucidated by the means of HRESIMS, NMR data and single-crystal X-ray diffraction experiment.

Compound 1 (peak 17) was isolated as a yellow amorphous powder. Its molecular formula was assigned as  $C_{13}H_{13}N_3O_2S$  from HRESIMS peak at  $m/z$  276.0799  $[M + H]^+$  (calcd for 276.0807), which requires nine degrees of unsaturation. The  $^1H$  NMR spectrum showed signals for a proton spin system that included  $\delta_H$  9.33 (1H, d,  $J = 1.8$  Hz, H-2'), 8.66 (1H, dd,  $J = 4.8$ , 1.6 Hz, H-6'), 8.49 (1H, dd,  $J = 6.9$ , 4.8 Hz, H-5') and 7.56 (1H, ddd,  $J = 6.9$ , 1.8, 1.6 Hz, H-4') (Table 1). The above data along with the observation of the  $^{13}C$  NMR and DEPT resonances of one olefinic nonprotonated carbons at  $\delta_C$  133.6 (C-3') and four

olefinic methines carbons at  $\delta_C$  150.1 (C-6'), 148.0 (C-2'), 134.2 (C-5'), 128.9 (C-4') indicated the presence of a mono-substituted pyridine ring (Table 2).<sup>13</sup> Similarly, the  $^{13}C$  NMR and DEPT data further displayed four olefinic nonprotonated carbons at  $\delta_C$  166.8 (C-4), 154.6 (C-2), 153.0 (C-6), 120.0 (C-5) and one olefinic methine carbons at 103.9 (C-3), along with a singlet at  $\delta_H$  7.70 (1H, s, H-3), suggesting a tetra-substituted pyridine ring in 1.<sup>14</sup> The HMBC correlation between H-3 with C-3' was detected and confirmed the existence of 2,3'-dipyridyl moiety (Fig. 4). The  $^1H$  NMR data also showed signals for one methoxyl group and one methyl group at  $\delta_H$  4.08 (3H, s, H-8) and 2.33 (3H, s, H-9), respectively. The HMBC correlation between  $\delta_H$  4.08 with  $\delta_C$  166.8 indicated that the methoxyl group was substituted at C-4.



Table 1  $^1\text{H}$  NMR data of compounds 1–6 (400 MHz, in DMSO,  $\delta$  in ppm,  $J$  in Hz)

Position	1	2	3	4	5	6
3	7.70, s	8.03, s	8.10, s	7.68, s	7.93, s	8.03, s
7	8.72, s		3.91, s	8.77, s		4.87, d (8.8)
8	4.08, s	4.06, s	4.10, s	4.08, s	4.13, s	4.04, s
9	2.33, s	2.33, s	2.35, s	3.00, s	3.06, s	3.01, s
2'	9.33, d (1.8)			8.18, dd (7.7, 1.8)	8.19, dd (7.2, 2.4)	
3'		8.40, ddd (8.0, 1.2, 1.0)	8.31, ddd (8.0, 1.3, 1.2)	7.52, m	7.57, m	8.47, ddd (7.8, 1.3, 1.2)
4'	7.56, ddd (6.9, 1.8, 1.6)	7.94, ddd (8.0, 6.8, 1.6)	7.96, ddd (8.0, 6.0, 2.0)	7.52, m	7.57, m	7.96, ddd (7.8, 6.8, 1.6)
5'	8.49, dd (6.9, 4.8)	7.48, ddd (6.8, 4.8, 1.2)	7.51, ddd (6.0, 4.8, 1.3)	7.52, m	7.57, m	7.50, ddd (6.8, 4.9, 1.3)
6'	8.66, dd (4.8, 1.6)	8.70, ddd (4.8, 1.6, 1.0)	8.73, ddd (4.8, 2.0, 1.2)	8.18, dd (7.7, 1.8)	8.19, dd (7.2, 2.4)	8.71, ddd (4.9, 1.6, 1.2)
NOH	11.78, brs			11.93, brs		
–CONH <sub>2</sub>		7.98, d (1.77)	7.57, s			
7-OH						5.35, brs

Taking into consideration the presence of one sulfur atom in the molecular formula of **1** and the chemical shifts of H-9, C-9 ( $\delta_{\text{C}}$  17.7), and C-5, the existence of a methylthio group was established. The HMBC correlation between  $\delta_{\text{H}}$  2.33 with  $\delta_{\text{C}}$  120.0 indicated that the methylthio group was located at C-5 position. Eight degrees of unsaturation were occupied by two pyridyl groups. Thus, the remaining unsaturation suggested that **1** may possess a double bond, which was consistent with the NMR data of the downfield carbon chemical shift at  $\delta_{\text{C}}$  146.7 (C-7) and one N–OH proton signal at  $\delta_{\text{H}}$  11.78 (1H, brs). HMBC correlations of  $\delta_{\text{H}}$  8.72 (H-7) with  $\delta_{\text{C}}$  153.0 and 120.0 confirmed the presence of an oxime group at C-6. The geometry of the oxime group was established by the comparison of the chemical shift of the N–OH signal with those in the known bipyridine analogue, SF2738 A (**7**).<sup>14,15</sup> Thus, compound **1** was characterized as (*E*)-4-methoxy-5-(methylthio)-[2,3'-bipyridine]-6-carboxaldehyde oxime and named pyrismycin A.

Compound **2** (peak 18) was isolated as yellow amorphous powder. Its molecular formula was determined to be C<sub>13</sub>H<sub>13</sub>N<sub>3</sub>O<sub>2</sub>S based on the observation of an [M + Na]<sup>+</sup> ion at *m/z* 298.0619 (calcd for 298.0626) in its positive ion mode HRESIMS spectrum. Similar to the known compound **10**,<sup>15</sup> whose structure

was elucidated by a single-crystal X-ray diffraction experiment (Fig. S3<sup>†</sup>), **2** is also a 2,2'-dipyridyl derivative, as shown by the presence of a proton spin system including  $\delta_{\text{H}}$  8.70 (1H, ddd,  $J = 4.8, 1.6, 1.0$ , H-6'), 8.40 (1H, ddd,  $J = 8.0, 1.2, 1.0$ , H-3'), 7.94 (1H, ddd,  $J = 8.0, 6.8, 1.6$ , H-4') and 7.48 (1H, ddd,  $J = 6.8, 4.8, 1.2$ , H-5'), a singlet signal at  $\delta_{\text{H}}$  8.03 (1H, s, H-3) (Table 1), five olefinic nonprotonated carbons at  $\delta_{\text{C}}$  166.5 (C-4), 158.0 (C-2), 155.3 (C-2'), 154.2 (C-6) and 118.5 (C-5), and five olefinic methine carbons at  $\delta_{\text{C}}$  149.1 (C-6'), 137.2 (C-4'), 124.5 (C-5'), 120.9 (C-3') and 103.2 (C-3) (Table 2). The HMBC correlation of  $\delta_{\text{H}}$  8.03 with  $\delta_{\text{C}}$  155.3 further confirmed the existence of a 2,2'-dipyridyl skeleton (Fig. 4). A methoxyl group at  $\delta_{\text{H}}$  4.06 (3H, s, H-8) and  $\delta_{\text{C}}$  56.2 (C-8), a methylthio group at  $\delta_{\text{H}}$  2.33 (3H, s, H-9) and  $\delta_{\text{C}}$  17.4 (C-9) and an amide group at  $\delta_{\text{H}}$  7.98 (1H, d,  $J = 1.77$  Hz), 7.57 (1H, s) and  $\delta_{\text{C}}$  168.5 (C-7) were observed in NMR data of **2**. Cross peaks of  $\delta_{\text{H}}$  4.06 with  $\delta_{\text{C}}$  166.5 and  $\delta_{\text{H}}$  2.33 with  $\delta_{\text{C}}$  118.5 in HMBC indicated that the methoxyl group and methylthio group were located at C-4 and C-5, respectively. Thus, the amide group was substituted at C-6. **2** was characterized as 4-methoxy-5-(methylthio)-[2,2'-bipyridine]-6-carboxamide and named pyrismycin B.

Compound **3** (peak 5) was isolated as yellow amorphous powder. Its molecular formula was determined as C<sub>13</sub>H<sub>14</sub>N<sub>2</sub>O<sub>2</sub>S

Table 2  $^{13}\text{C}$  NMR data of compounds 1–6 (100 MHz, in DMSO,  $\delta$  in ppm)

Position	1	2	3	4	5	6
2	154.6, C	158.0, C	156.2, C	159.4, C	161.2, C	157.7, C
3	103.9, CH	103.2, CH	104.3, CH	104.1, CH	107.5, CH	103.3, CH
4	166.8, C	166.5, C	167.0, C	165.0, C	164.1, C	164.6, C
5	120.0, C	118.5, C	117.7, C	126.2, C	132.3, C	126.3, C
6	153.0, C	154.2, C	167.0, C	150.4, C	136.3, C	159.8, C
7	146.7, CH	168.5, C	52.5, CH <sub>3</sub>	146.1, CH	115.0, C	62.8, CH <sub>2</sub>
8	56.6, CH <sub>3</sub>	56.2, CH <sub>3</sub>	56.5, CH <sub>3</sub>	56.7, CH <sub>3</sub>	57.5, CH <sub>3</sub>	56.5, CH <sub>3</sub>
9	17.7, CH <sub>3</sub>	17.4, CH <sub>3</sub>	17.4, CH <sub>3</sub>	39.5, CH <sub>3</sub>	39.5, CH <sub>3</sub>	39.0, CH <sub>3</sub>
1'				137.5, C	130.5, C	
2'	148.0, CH	155.3, C	153.6, C	127.1, CH	127.4, CH	153.9, C
3'	133.6, C	120.9, CH	120.7, CH	128.6, CH	128.9, CH	121.2, CH
4'	128.9, CH	137.2, CH	137.8, CH	130.0, CH	130.7, CH	137.3, CH
5'	134.2, CH	124.5, CH	125.1, CH	128.6, CH	128.9, CH	124.9, CH
6'	150.1, CH	149.1, CH	149.6, CH	127.1, CH	127.4, CH	149.2, CH



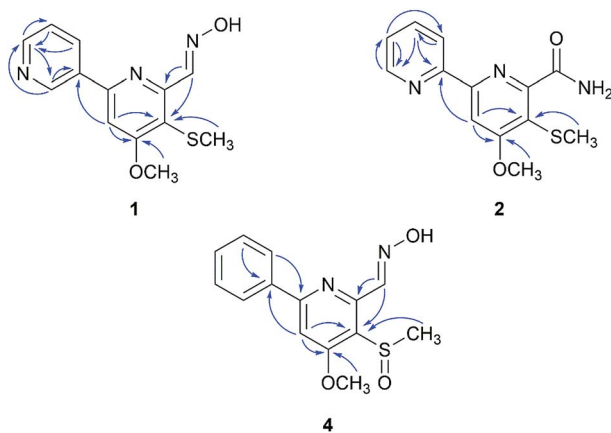


Fig. 4 Key HMBC correlations of 1, 2 and 4.

according to an  $[M + Na]^+$  ion at  $m/z$  285.0666 (calcd for 285.0674) in its positive ion mode HRESIMS spectrum. Comparison of the  $^{13}C$  and  $^1H$  NMR data of 3 with 2 revealed extensive similarities, except that the methoxy signals at  $\delta_H$  3.91 (3H, s, H-7) and  $\delta_C$  52.5 (C-7) in 3 was replaced by an amide group ( $\delta_H$  7.98 and 7.57,  $\delta_C$  168.5) in 2. The HMBC correlation of  $\delta_H$  3.91 with  $\delta_C$  167.0 (C-6) indicated that the methoxyl group was substituted at C-6. Thus, compound 3 was characterized as 4,6-dimethoxy-5-(methylthio)-2,2'-bipyridine and named pyrismycin C.

Compound 4 (peak 9) was isolated as yellow amorphous powder. The molecular formula was determined to be  $C_{14}H_{14}N_2O_3S$  from the  $[M + H]^+$  peak at  $m/z$  291.0793 (calcd for 291.0803) in its positive ion mode HRESIMS spectrum. The NMR data of 4 were similar to pyrisulfoxin A (11), previously isolated from *Streptomyces californicus* BS-75.<sup>16</sup> Both compounds shared the same tetra-substituted pyridine ring as revealed by the presence at  $\delta_C$  165.0 (C-4), 159.4 (C-2), 150.4 (C-6), 126.2 (C-5), 104.1 (C-3) (Table 2). Moreover, a methoxyl group at  $\delta_H$  4.08 (3H, s, H-8), a methylsulfoxide group at  $\delta_H$  3.00 (3H, s, H-9) and an oxime group at  $\delta_H$  8.77 (1H, s, H-7) (Table 1) were also analogous of 11 and substituted at C-4, C-5 and C-6 based on the HMBC correlations of  $\delta_H$  4.08 with  $\delta_C$  165.0,  $\delta_H$  3.00 with  $\delta_C$  126.2 and  $\delta_H$  8.77 (1H, s, H-7) with  $\delta_C$  150.4, respectively (Fig. 4). The difference lies on the mono-substituted pyridine ring in 11 and the mono-substituted phenyl ring in 4, as shown by the protons at  $\delta_H$  8.18 (each 1H, dd,  $J = 7.7, 1.8$  Hz, H-2', 6') and 7.51–7.53 (3H, m, H-3', 4', 5'), as well as the aromatic carbon signals at  $\delta_C$  137.5 (C-1'), 130.0 (C-4'), 128.6  $\times$  2 (C-3', 5') and 127.1  $\times$  2 (C-2', 6'). The HMBC correlations of  $\delta_H$  7.68 (1H, s, H-3) and 7.52 (each 1H, m, H-3'/4'/5') with C-1' and 8.18 (H-2') with C-2 confirmed the pyridine ring was connected at C-2 (Fig. 4). Thus, 4 was characterized as (*E*)-4-methoxy-5-(methylsulfinyl)-6-phenylpicolininaldehyde oxime and named pyrismycin D.

Compound 5 (peak 15) was isolated as yellow amorphous powder. Its HRESIMS data exhibited a  $[M + Na]^+$  ion at  $m/z$  295.0511 (calcd for 295.0517) corresponded to a molecular formula of  $C_{14}H_{12}N_2O_2S$  and revealed eleven degrees of unsaturation. The UV and NMR data features of compound 5 were very similar to those of 4, except for the substitution of a cyano

group ( $\delta_C$  115.0) at C-6 in 5, which was also supported by the existence of two nitrogen atom in its molecular formula, instead of an oxime group in 4. Thus, compound 5 was characterized as 4-methoxy-5-(methylsulfinyl)-6-phenylpicolinonitrile and named pyrismycin E.

Compound 6 (peak 16) was isolated as yellow amorphous powder. Its molecular formula was assigned as  $C_{13}H_{14}N_2O_3S$  on the basis of a  $[M + Na]^+$  ion at  $m/z$  301.0620 (calcd for 301.0623) in its positive ion mode HRESIMS spectrum. The  $^1H$  and  $^{13}C$  NMR data of 6 indicated it is an analogue of 11.<sup>14</sup> Comparison of the NMR data of 6 with those of 11 showed the absence of the oxime group, and the presence of a methylol group at  $\delta_C$  62.8 (C-7),  $\delta_H$  4.87 (2H, d,  $J = 8.8$  Hz, H-7) and 5.35 (1H, brs) in 6. This methylol group was positioned at C-6 based on the HMBC correlation of  $\delta_H$  4.87 with  $\delta_C$  159.8 (C-6). Thus, 6 was characterized as (4-methoxy-5-(methylsulfinyl)-[2,2'-bipyridin]-6-yl) methanol and named pyrismycin F.

The structures of compounds 7–13 were identified as the previously reported natural compounds, SF2738A (7, peak 19),<sup>14</sup> SF2738C (8, peak 6),<sup>13</sup> SF2738D (9, peak 4),<sup>15</sup> SF2738E (10, peak 11),<sup>15</sup> pyrisulfoxin A (11, peak 7),<sup>16</sup> pyrisulfoxin B (12, peak 14)<sup>16</sup> and caerulomycin A (13, peak 12),<sup>11</sup> by a comparison of their  $^1H$  and  $^{13}C$  NMR data with the literature.

All isolates (1–13) were evaluated for cytotoxicity against the SGC7901, MDA-MB-231, A549 and HepG2 cell lines (ESI Experimental section†). Compounds 7, 9 and 11 displayed measurable  $IC_{50}$  values, but others showed no cytotoxicity (Table S2†). The most potent one, SF2738A (7), showed cytotoxic activities against SGC7901 with  $IC_{50}$  value of 1.7  $\mu$ M, HepG2 with  $IC_{50}$  value of 5.8  $\mu$ M and MDA-MB-231 with  $IC_{50}$  value of 6.3  $\mu$ M, while pyrisulfoxin A (11) showed distinct cytotoxic activity for HepG2 with  $IC_{50}$  value of 7.2  $\mu$ M. A gross structure–activity relationships (SAR) for this class of compounds was discussed as follows: (1) the 2,2'-dipyridyl moiety may be crucial for the cytotoxicity since pyrismycin A (1), with a greatly similar structures with its isomer 7 except for the 2,3'-dipyridyl moiety, has no displayed deemed cytotoxic for any of the cell lines used. (2) The absence of oxime group in C-7 may remarkably reduce the cytotoxicity. Compound 3 with the methoxyl group in place of the oxime group of 7 significantly decreased the inhibitory activities, which also can be observed in the other analogues of 2 and 8–10. (3) As compared with 7, the presence of a methylsulfoxide group at C-5 in 11 may obviously improve the selectivity of the types of cancer cell lines.

## Experimental

### General experimental procedures

UV spectra were recorded on a Shimadzu UV-260 spectrophotometer (Shimadzu Corp., Japan) in absolute methanol (MeOH). IR spectra were record as KBr pellets on the Avatar 360 E.S.P spectrophotometer (Thermo Nicolet Co. Boston, MA, USA). NMR data were acquired with a Bruker Avance 400 spectrometer using solvent signals (DMSO;  $\delta_H$  2.50/ $\delta_C$  39.52) as reference. UPLC-QTOF-MS was operated using an Acquity UPLC system (Waters Corp., Milfoed, USA) with a QTOF-MS (XEVO-G2 QTOF, Waters MS Technologies, Manchester, UK), controlled by





MassLynx V4.1 software. Semipreparative reversed-phase HPLC separation was performed on a Shimadzu LC-6AD instrument packed with a YMS-pack ODS-A column (5  $\mu\text{m}$ , 250  $\times$  10 mm). X-ray diffraction was realized on a Bruker D8 QUEST X-ray single crystal diffractometer using Mo K $\alpha$  radiation ( $\lambda = 0.71073$ ). Sephadex LH-20 (GE Healthcare Bio-Sciences AB, Uppsala, Sweden), MCI Gel CHP 20/P120 (Mitsubishi Chemical Corp., Japan) and ODS-A 12 nm S-50  $\mu\text{m}$  (YMC Co., Ltd, Japan) were used for column chromatography, respectively. Human gastric carcinoma SGC7901, breast carcinoma MDA-MB-231, lung carcinoma A549 and hepatocellular carcinoma HepG2 were obtained from Shanghai Institutes for Biological Sciences, Chinese Academy of Sciences.

### Bacteria material

*S. roseosporus* *Lsr2*-deletion mutant strain ( $\Delta$ *Srlsr2*) and WT strain were obtained from Dr Guojian Liao, College of Pharmaceutical Sciences, Southwest University, China. The voucher specimens (no. 2016102901 and 2016102902) were deposited in the College of Pharmaceutical Sciences, Southwest University, China.

### Fermentation and isolation

The  $\Delta$ *Srlsr2* strain was grown on PDA plates at 28  $^{\circ}\text{C}$  for 6 days. Then the agar plate were cut into small pieces and inoculated into sterilized liquid potato dextrose broth medium in Erlenmeyer flasks for further incubation at 28  $^{\circ}\text{C}$  on a rotary shaker at 170 rpm for 6 days. The fermented whole broth was filtered to remove the mycelia, and the broth was then extracted four times with ethyl acetate to get about 60 L of extract solution. Then solvent was evaporated under reduced pressure to afford a crude residue (20.9 g). The extract was applied to an ODS column eluted with water, 30%, 40%, 50%, 60%, 70%, 80%, 100% MeOH and concentrated, affording 4 combined fractions (Fr.1–Fr.4). Fr.3 was then separately isolated on MCI gel column and eluted with MeOH–H<sub>2</sub>O to obtain 3 subfractions Fr.3.1 to Fr.3.3. Fr.3.1 was separated by semipreparative HPLC, using a gradient of 75–92% MeOH in H<sub>2</sub>O over 47 min, to give **3** (2.5 mg,  $t_{\text{R}} = 36$  min) and **9** (1.6 mg,  $t_{\text{R}} = 23$  min). Fr.3.2 was separated over Sephadex LH-20 column chromatography, using CH<sub>2</sub>Cl<sub>2</sub>–MeOH (1 : 3) as an eluent, to yield two subfractions (Fr.3.2.1 and Fr.3.2.2). Fr.3.2.1 was further purified using semipreparative HPLC, eluting with a MeOH–H<sub>2</sub>O (2 : 3) over 88 min, to give **4** (1.8 mg,  $t_{\text{R}} = 75$  min), **8** (3.5 mg,  $t_{\text{R}} = 32$  min) and **11** (2.5 mg,  $t_{\text{R}} = 36$  min). Similarly, Fr.3.2.2 was subjected to semipreparative HPLC, eluted with 33% MeOH in H<sub>2</sub>O over 63 min, to provide **5** (1.0 mg,  $t_{\text{R}} = 54$  min), **10** (2.7 mg,  $t_{\text{R}} = 27$  min), **12** (2.2 mg,  $t_{\text{R}} = 58$  min) and **13** (1.2 mg,  $t_{\text{R}} = 33$  min). Fr.3.3 was chromatographed by semipreparative HPLC, eluting with a step gradient of 32–60% MeOH in H<sub>2</sub>O over 93 min, giving **1** (2.0 mg,  $t_{\text{R}} = 64$  min), **2** (1.8 mg,  $t_{\text{R}} = 80$  min), **6** (2.1 mg,  $t_{\text{R}} = 39$  min) and **7** (3.0 mg,  $t_{\text{R}} = 85$  min).

### Structure characterization

**Pyrimycin A (1).** Yellow, amorphous powder; UV (MeOH)  $\lambda_{\text{max}}$  (log  $\epsilon$ ) 239 (3.9) nm; IR (KBr)  $\nu_{\text{max}}$  3431, 2929, 2872, 2364,

1631, 1568, 1377, 1300, 1232, 1128, 1018, 871, 790, 758, 692  $\text{cm}^{-1}$ ; <sup>1</sup>H and <sup>13</sup>C NMR data see Tables 1 and 2; HRESI (+) MS  $m/z$  276.0799 (calcd for 276.0807).

**Pyrimycin B (2).** Yellow, amorphous powder; UV (MeOH)  $\lambda_{\text{max}}$  (log  $\epsilon$ ) 285 (3.8) nm; IR (KBr)  $\nu_{\text{max}}$  3441, 3361, 2926, 2858, 1662, 1570, 1452, 1375, 1217, 1124, 1045, 854, 786, 738, 661  $\text{cm}^{-1}$ ; <sup>1</sup>H and <sup>13</sup>C NMR data see Tables 1 and 2; HRESI (+) MS  $m/z$  298.0619 (calcd for 298.0626).

**Pyrimycin C (3).** Yellow, amorphous powder; UV (MeOH)  $\lambda_{\text{max}}$  (log  $\epsilon$ ) 284 (3.6) nm; IR (KBr)  $\nu_{\text{max}}$  3433, 2933, 1724, 1647, 1635, 1598, 1438, 1369, 1224, 1033, 781, 692  $\text{cm}^{-1}$ ; <sup>1</sup>H and <sup>13</sup>C NMR data see Tables 1 and 2; HRESI (+) MS  $m/z$  285.0666 (calcd for 285.0674).

**Pyrimycin D (4).** Yellow, amorphous powder; UV (MeOH)  $\lambda_{\text{max}}$  (log  $\epsilon$ ) 247 (3.8) nm; <sup>1</sup>H and <sup>13</sup>C NMR data see Tables 1 and 2; HRESI (+) MS  $m/z$  291.0793 (calcd for 291.0803).

**Pyrimycin E (5).** Yellow, amorphous powder; UV (MeOH)  $\lambda_{\text{max}}$  (log  $\epsilon$ ) 243 (3.8) nm, 279 (3.7) nm; <sup>1</sup>H and <sup>13</sup>C NMR data see Tables 1 and 2; HRESI (+) MS  $m/z$  295.0511 (calcd for 295.0517).

**Pyrimycin F (6).** Yellow, amorphous powder; UV (MeOH)  $\lambda_{\text{max}}$  (log  $\epsilon$ ) 281 (4.1) nm; IR (KBr)  $\nu_{\text{max}}$  3441, 3346, 2987, 2926, 1564, 1421, 1271, 1217, 1028, 945, 881, 788, 686, 621, 540  $\text{cm}^{-1}$ ; <sup>1</sup>H and <sup>13</sup>C NMR data see Tables 1 and 2; HRESI (+) MS  $m/z$  301.0620 (calcd for 301.0623).

**X-ray crystallographic data for 10.** C<sub>15</sub>H<sub>17</sub>N<sub>3</sub>O<sub>2</sub>S, prism crystals from CHCl<sub>3</sub>/MeOH,  $M = 303.38$ , monoclinic,  $a = 11.2581(2)$  Å,  $b = 28.6822(10)$  Å,  $c = 19.2173(5)$  Å,  $\beta = 103.658(2)^{\circ}$ ,  $V = 6029.9(3)$  Å<sup>3</sup>,  $T = 293(2)$  K, space group  $P2_1/n$  (#14),  $Z = 4$ ,  $\mu$  (Mo K $\alpha$ ) = 0.223  $\text{mm}^{-1}$ , 27 243 reflections measured, 13 668 independent reflections ( $R_{\text{int}} = 0.0224$ ). The final  $R_1$  values were 0.0536 ( $I > 2\sigma(I)$ ), and the final  $wR_2$  ( $F^2$ ) values were 0.1460 (all data). The goodness of fit on  $F^2$  was 1.057. Crystallographic data were deposited with the Cambridge Crystallographic Data Centre (deposition code: CCDC 1533762).†

### UPLC-QTOF-MS conditions

Chromatographic separation was performed with a 2.1  $\times$  50 mm i.d., 1.7  $\mu\text{m}$  UPLC BEH C18 reversed-phase column and kept at a temperature of 40  $^{\circ}\text{C}$ . The mobile phase consisted of water (A) and acetonitrile (B). The linear gradient elution was performed as follows: 0–0.4 min, 5% B; 0.4–20 min, 5–45% B; 20–24 min, 45–65% B; 24–26 min, 65–76% B; 26–35 min, 76–100% B; 35–36 min, 100–0.5% B; 36–38 min, 0.5–0.5% B. A flow rate of 0.4  $\text{mL min}^{-1}$  was employed for elution, and the injected sample volume was set at 2  $\mu\text{L}$ . Mass spectrometry was recorded using a Xevo G2 QTOF equipped with an ESI source and MS full scanning was conducted in positive ion modes over the range  $m/z$  100–1000 Da, with a scan time of 0.2 s. The capillary voltages were set at 3000 V and the cone voltage was 40 V. Nitrogen gas was used both for the nebulizer and in desolvation. The desolvation and cone gas flow rates were 800 and 50  $\text{L h}^{-1}$ , respectively. The desolvation temperature was 450  $^{\circ}\text{C}$ , and the source temperature was 120  $^{\circ}\text{C}$ . The tandem MS data were acquired in the positive mode with a range of  $m/z$  100–1000 Da, and the low energy was set as 6 V, while the high energy was ramped from 20 to 40 V. The cone voltage was set as 40 V.



## Conclusions

The combination of UPLC-QTOF-MS/MS and HPLC analysis is an effective chemical dereplication method for structural analysis of compounds and rapid isolation of unknowns of interest in  $\Delta SrLsr2$  strain, whereas the conventional separation often requires a great deal of work to obtain the desired compounds. Pyrisulfoxins A–F (1–6), six sulfur-containing metabolites have not yet been described in the literature, and seven known analogues (7–13) were isolated from the  $\Delta SrLsr2$  strain. Among these compounds tested for cytotoxicity, SF2738A (7) showed the highest activity for SGC7901 cell lines, and pyrisulfoxin A (11) showed moderate cytotoxic for HepG2 cell lines. It was suggested that the 2,2'-dipyridyl moiety may be crucial for the cytotoxicity, and the absence of oxime group in C-7 may remarkably reduce the cytotoxicity.

This is the first report that deletion of the global regulator Lsr2 in *S. roseosporus* upregulated the expression of biosynthetic gene cluster and produced new compounds, pyrisulfoxins A–F (1–6). And this further demonstrated that deletion of the global regulator Lsr2 is an effective approach to produce new secondary metabolites.

## Conflict of interest

There are no conflicts of interest to declare.

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