

Figure S1. Confirmation of the integrity of the captured *spz* BGC, related to Figures 1 and 2 and Table 1. **A.** Restriction digest of pSMM before and after transformation in *E. coli* using KpnI and KpnI + SpeI enzymes. KpnI digest, expected sizes (kb) = 23.9, 11.1, 8.2, 4.9, 4.1, 2.7, 1.9, 1.5, 1.1, 1.0, 0.5. KpnI + SpeI digest, expected sizes (kb) = 22.9, 8.9, 8.2, 4.9, 4.1, 2.7, 2.1, 1.9, 1.5, 1.1, 1.0, 0.5. **B.** Restriction digest of pKDB01, expected sizes (kb): 18.0, 13.7, 8.3, 4.9, 2.0, 1.6, and pKDB02, expected sizes (kb): 17.0, 13.3, 8.3, 4.9, 2.0, 1.5, with EcoRI + Stul. * indicates digest is blocked due to overlapping *dcm* methylation. Std – sequenced plasmid digested with EcoRI + Stul, used as a control, expected sizes (kb) = 10.3, 3.2, 1.3. Plasmid backbone represented by blue rectangle.

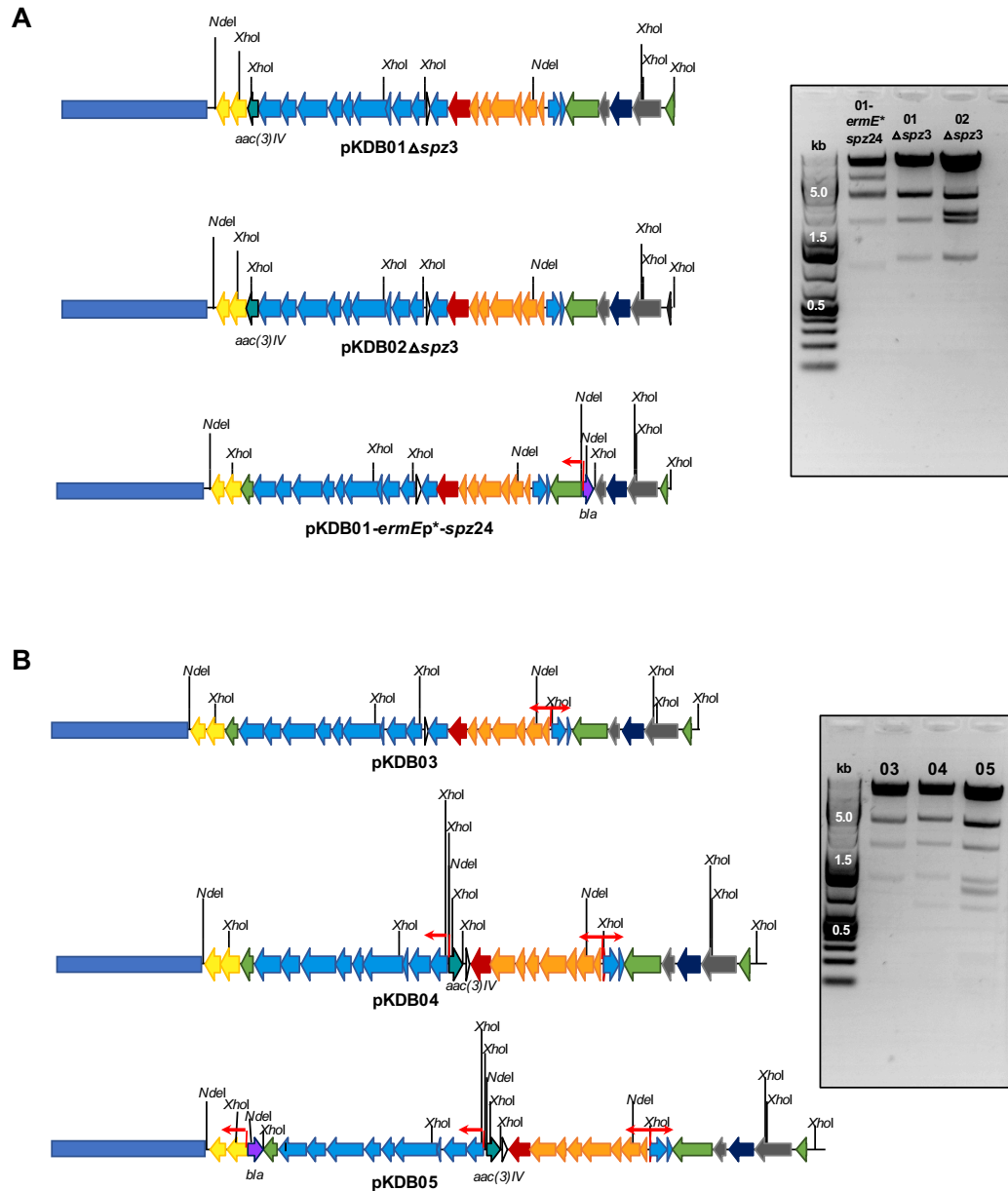
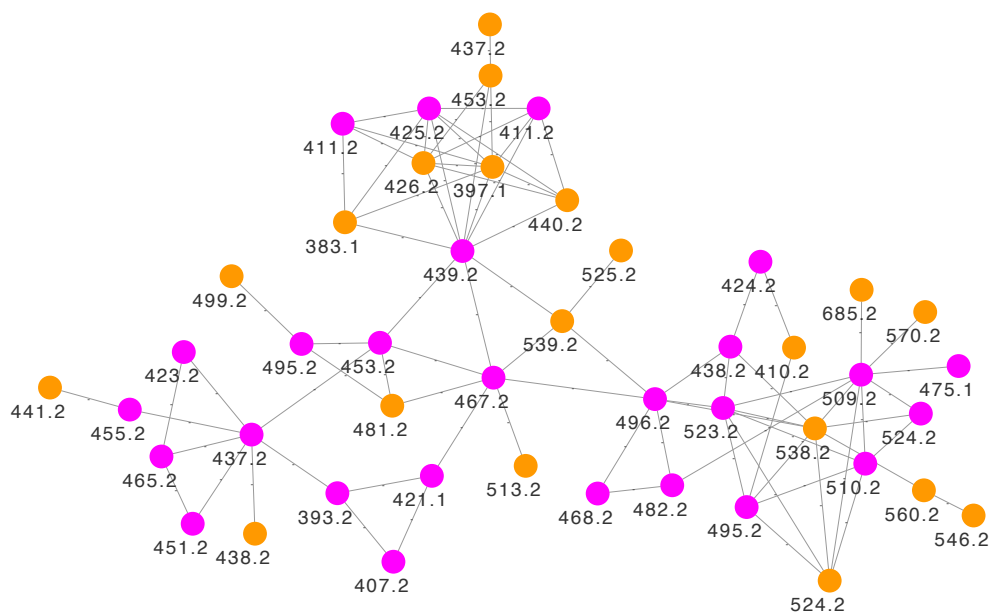


Figure S2. Engineering the *spz* BGC, related to Figures 2 and 3. A. Agarose gels of restriction digestions and corresponding maps of regulatory gene mutant plasmids. Enzyme pair *XhoI* + *NdeI* used for digestion. pKDB01 Δ *spz3*, expected sizes (kb): 10.6, 10.5, 8.7, 8.6, 3.2, 3.1, 1.9, 1.0, 0.1. pKDB02 Δ *spz3*, expected sizes (kb): 10.6, 10.5, 8.7, 8.6, 3.2, 2.1, 1.9, 1.0, 0.1. pKDB01-*ermE***p-spz24*, expected sizes (kb): 11.7, 10.5, 8.7, 5.2, 3.4, 3.2, 3.1, 1.9, 0.9, 0.2, 0.1. **B.** Restriction digest with *XhoI* + *NdeI* of refactored plasmids, agarose gel and corresponding plasmid map showing restriction sites: pKDB03, expected sizes (kb): 11.7, 10.5, 8.7, 7.6, 3.2, 3.1, 1.9, 1.1, 0.1; pKDB04, expected sizes (kb): 11.7, 10.5, 8.6, 7.5, 3.2, 3.1, 1.9, 1.1, 0.7, 0.3, 0.2, 0.1; pKDB05, expected sizes (kb): 10.9, 10.5, 8.7, 7.5, 3.2, 3.1, 1.9, 1.1, 0.9, 0.9, 0.7, 0.3, 0.3, 0.1. Plasmid backbone represented by blue rectangle.

A

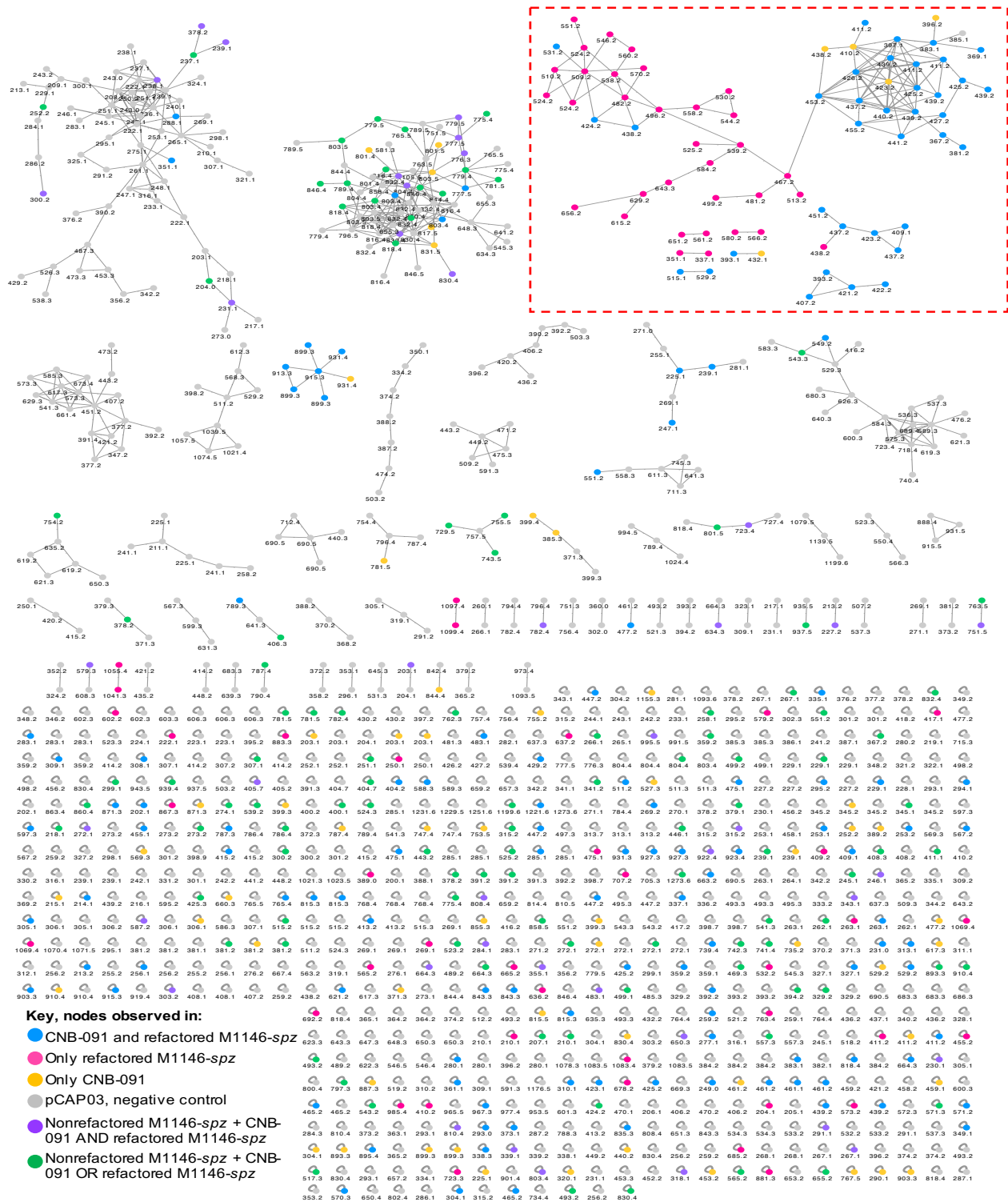
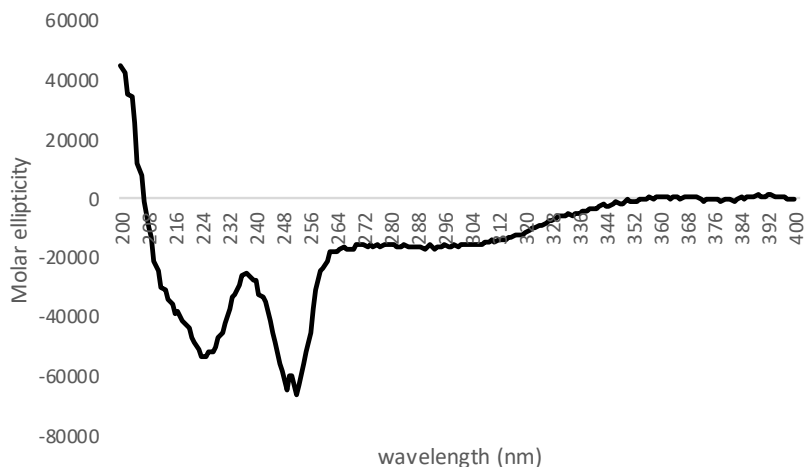


Figure S4. Molecular network showing production of streptopenazines, related to Figure 4. LCMS/MS data of ethyl acetate extracts from CNB-091, M1146-pCAP03, M1146-pKDB01 (non-refactored *spz* BGC) and M1146-pKDB03 (refactored *spz* BGC) were used for network generation. Nodes corresponding to streptopenazines highlighted in red square.

A**B**

```

Spz7_KR      ...-D1VYWAHL2PDT--VTPVEEILAAL3LDD4LVRAGKILHAGLSNFP5AWRV...
Ery1_KR      ...---LGGIGDDVPLSAVFHAAAT6LDD7GT-VDTLTGERIERASRAKVL...
SlnA1_KR     ...---LLDRIPEAHPLTGVFHAAGV8LDD9GM-VGALSAERLDAVLRPKTD...
RifA_KR      ...LEAVLRAIPA10EHPLTAVIHTAGV11LDD12GV-VTELT13PDRLATVRRPKVD...
AmphJ_KR     ...---LLASVPAEHPLTAVVHTAGV14LDD15GI-FPSLT16PDRLDSVMPKVD...
TlmH_KR     ...---VLAQIRSRGPIGGVVHAAGL17LDD18SI-LANMTPEQLHRVLR19SKVD...

```

Figure S5. Confirmation of stereochemistry at C-1' position of compound 18, related to Figure 1. A. Circular dichroism (CD) spectrum of streptopenazine G (**18**). **B.** Alignment of type I PKS KR domains and Spz7. Amino acid residues (LDD motif) defining Spz7 as type B KR are highlighted. Ery1 = erythromycin, SlnA1 = salinomycin, RifA = rifamycin, AmphJ = amphotericin, TlmH = thiolactomycin.

A**Sequence search results**[Show](#) the detailed description of this results page.

We found 2 Pfam-A matches to your search sequence (all significant)

[Show](#) the search options and sequence that you submitted.[Return](#) to the search form to look for Pfam domains on a new sequence.**Significant Pfam-A Matches**[Show](#) or [hide](#) all alignments.

Family	Description	Entry type	Clan	Envelope		Alignment		HMM		HMM length	Bit score	E-value	Predicted active sites	Show/hide alignment
				Start	End	Start	End	From	To					
AMP-binding	AMP-binding enzyme	Family	CL0378	19	441	19	440	1	422	423	287.3	1.6e-85	n/a	Show
AMP-binding_C	AMP-binding enzyme C-terminal domain	Domain	CL0531	449	525	449	525	1	76	76	43.6	4.3e-11	n/a	Show

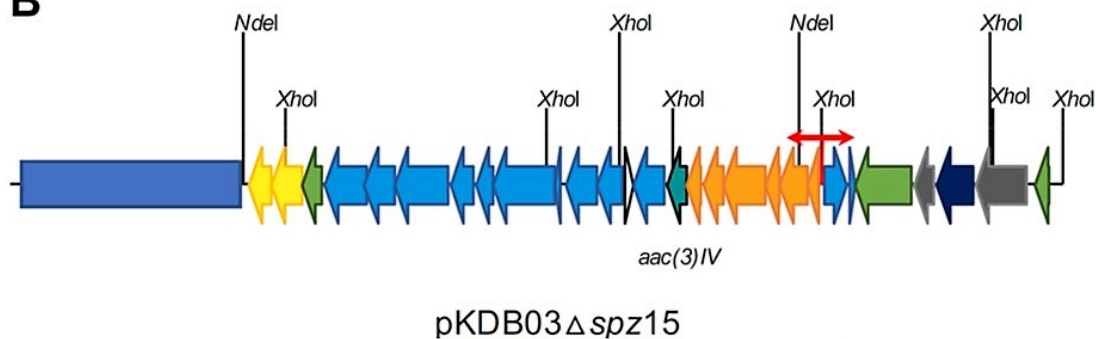
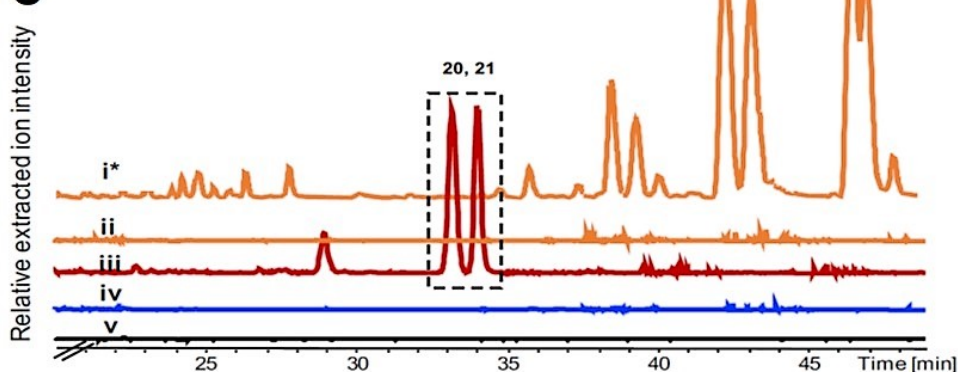
B**C**

Figure S6. Bioinformatic analysis and gene deletion of *spz15*, related to Figures 5 and 6. **A.** Protein family (Pfam) analysis of *Spz15*. Identified CL00378 AMP-binding domain which is characteristic for ANL superfamily of enzymes that includes adenylation domains. **B.** Restriction digest of pKDB03Δ*spz15* with *XhoI* and *NdeI* restriction enzymes. Expected sizes (kb): 11.7, 10.5, 7.5, 5.6, 3.2, 3.1, 2.3, 1.9, 1.1, 0.1. **C.** LCMS chromatograms: (i) Base Peak Chromatogram (BPC) of M1146-pKDB03Δ*spz15* *zoomed out 100x, (ii) Extracted Ion Chromatogram (EIC) (*m/z* 510.2, corresponding to compounds **20** and **21**) of M1146-pKDB03Δ*spz15*, (iii) M1146-pKDB03 EIC (*m/z* 510.2), (iv) CNB-091 EIC (*m/z* 510.2), and (v) M1146-pCAP03 EIC (*m/z* 510.2).

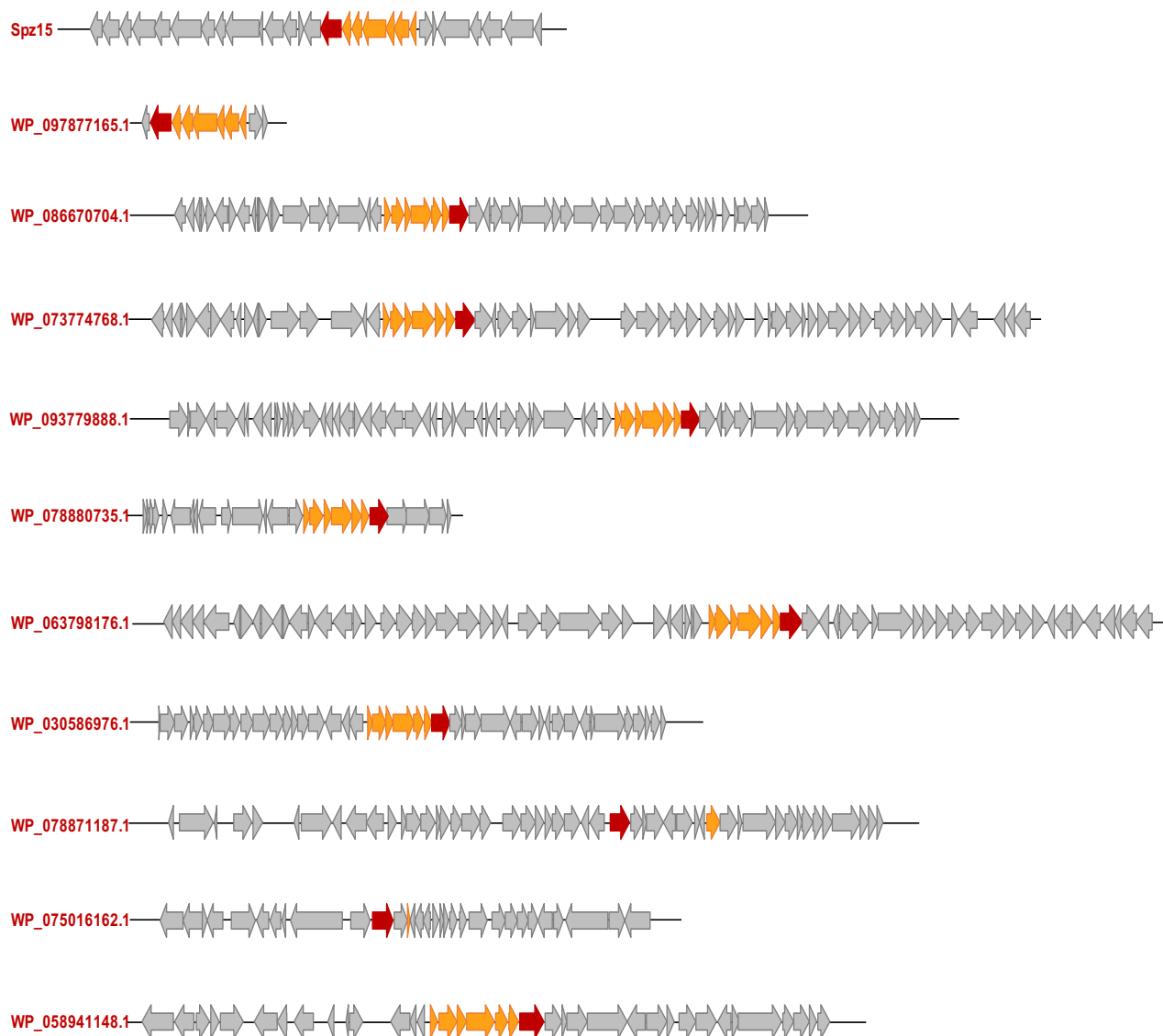


Figure S7. antiSMASH predicted gene neighborhoods of *phz*-associated discrete adenylation proteins, related to Figure 5. Red gene = adenylation enzyme homolog found through blastp, orange genes = phenazine biosynthesis homologs. Spz15 = from *Streptomyces* sp. CNB-091, WP_097877165.1 = from *Streptomyces* sp. ms184, WP_086670704.1 = from *Streptomyces albovinaceus*, WP_073774768.1 = from *Streptomyces* sp. TSRI0445, WP_093779888.1 = from *Streptomyces* sp. yr375, WP_078880735.1 = from *Kitasatospora purpeofusca*, WP_063798176.1 = from *Streptomyces* sp. 150FB, WP_030586976.1 = from *Streptomyces anulatus*, WP_078871187.1 = from *Streptomyces caatingaensis*, WP_075016162.1 = from *Streptomyces rubidus*, WP_058941148.1 = from *Streptomyces kanasensis*.

Table S1, related to Figures 2, 3 and Method Details. Promoter cassette sequences used for refactoring. Bold = restriction site, italic = added for scar, blue = FRT site, red = antibiotic resistance, green = promoter, purple = *actII-ORF4*.

Cassette	Sequence
<i>sp44-p21</i>	<p> CTCGAGGGTGAACCGATCTCCTCGTTAGGGTCAACCCAGACTTTACAACCCGCACAGCATGT TGTCAAAGCAGAGACGGTTTGAATGTGAACAGCCACTATCATATGTGCAGTTCGAAGTTCCTAT TCTCTAGAAAGTATAGGAACTTCGGTTCATGTGCAGCTCCATCAGCAAAGGGGATGATAAGTT TATCAACCACCGACTATTTGCAACAGTGCCGTTGATCGTGTATGATCGACTGATGTCATCAGCG GTGGAGTGCAATGTCGTGCAATACGAATGGCGAAAAGCCGAGCTCATCGGTGAGCTTCTCAAC CTTGGGGTTACCCCCGGCGGTGTGCTGCTGGTCCACAGCTCCTTCCGTAGCGTCCGGCCCT CGAAGATGGGCCACTTGACTGATCGAGGCCCTGCGTGTGCGTGGTCCGGGAGGGACG CTCGTCATGCCCTCGTGGTCAGGTCTGGACGACGAGCCGTTGATCCTGCCACGTCCGCCGT TACACCGGACCTTGAGTTGTCTTGACACATTCTGGCGCCTGCCAAATGTAAAGCGCAGCGC CCATCCATTTGCCTTGGCGGACGCGGGCCACAGGCAGAGCAGATCATCTCTGATCCATTGCC CCTGCCACCTCACTCGCTGCAAGCCCGGTGCCCGTGTCCATGAACTCGATGGGCAGGTAT TTCTCCTCGGCGTGGGACACGATGCCAACACGACGCTGCATCTTGCCGAGTTGATGGCAAAG GTTCCCTATGGGGTCCGAGACACTGCACCATTCTCAGGATGGCAAGTTGGTACGCGTCGAT TATCTCGAGAATGACCACTGCTGTGAGCGCTTGCCTTGGCGGACAGGTGGCTCAAGGAGAA GAGCCTTCAGAAGGAAGGTCCAGTCGGTCATGCCTTGTCTCGGTTGATCCGCTCCCGCGACAT TGTGGCGACAGCCCTGGGTCAACTGGGCCGAGATCCGTTGATCTTCTGCATCCGCCAGAGG CGGGATGCGAAGAATGCGATGCCGCTCGCCAGTCGATTGGCTGAGAAGTTCCTATTCTCTAGA AAGTATAGGAACTTCAAGCTTTGCTCGAGTGTGCGGGCTCTAACACGTCCTAGTATGGTAGGA TGAGCAATCTAGTCGAGCAACGGAGGTACGGACCATATG </p>
<i>actIp</i>	<p> CTCGAGGGTTCATGTGCAGCTCCATCAGCAAAGGGGATGATAAGTTTATCAACCACCGACTAT TTGCAACAGTGCCGTTGATCGTGTATGATCGACTGATGTCATCAGCGGTGGAGTGCAATGTC GTGCAATACGAATGGCGAAAAGCCGAGCTCATCGGTGAGCTTCTCAACCTTGGGGTTACCCCC GGCGGTGTGCTGCTGGTCCACAGCTCCTTCCGTAGCGTCCGGCCCTCGAAGATGGGCCACT TGGACTGATCGAGGCCCTGCGTGTGCGTGGTCCGGGAGGGACGCTCGTCTATGCCCTCG TGTCAGGTCTGGACGACGAGCCGTTGATCCTGCCACGTCCGCCGTTACACCGGACCTTGG AGTTGTCTTGACACATTCTGGCGCCTGCCAAATGTAAAGCGCAGCGCCCATCCATTTGCCTT GCGGACGCGGGCCACAGGCAGAGCAGATCATCTGATCCATTGCCCTGCCACCTCACTC GCCTGCAAGCCCGGTGCGCCGTGTCCATGAACTCGATGGGCAGGTACTTCTCCTCGGCGTGG GACACGATGCCAACACGACGCTGCATCTTGCCGAGTTGATGGCAAAGGTTCCCTATGGGGTG CCGAGACACTGCACCATTCTCAGGATGGCAAGTTGGTACGCGTCGATTATCTCGAGAATGAC CACTGCTGTGAGCGCTTGCCTTGGCGGACAGGTGGCTCAAGGAGAAGAGCCTTCAGAAGGA AGGTCCAGTCGGTCATGCCTTGTCTCGGTTGATCCGCTCCCGCGACATTGTGGCGACAGCCCT GGGTCAACTGGGCCGAGATCCGTTGATCTTCTGCATCCGCCAGAGGCGGGATGCGAAGAAT GATGCCGCTCGCCAGTCGATTGGCTGACATATGCCACTGCCTCTCGGTAATAATCCAGCAAA AATTAATCAGTGACGCTCGCTGCACTGATTAATTTTTGATCAATAGGAGATCGCTTGTGACGGC AAGCACATTGAAATCTGTTGAGTAGGCCTGTTATTGTGCGCCCCAGGAGACGGAGAATCTCGA CGGGGGCGCAGATGAGATTCAACTTATTGGGACGTGTCCATGTAATCACCGATGCGGGATGTG TAATTCGGCTTAAATCCTCGAAGGCGACCCAGCTCCTGGTGTGCTGCTCCTCAGGCGGCACG AGGTGGTGGGATCGGGGGTGTCTATCGAGGAGTTGTGGGCGGACCACCCGCCCGCAGCGC CATGACGACGCTGCAGACGTACGTGTACCACACCCGCCGGTGTGTTGGGGGAGCACCGGGTG ACGAGCGACGACCGGGAATTGGTCTGACCCAGCCGCGCCGGCTACTTCGCCCTGATCGACGA GGACGAATCGACGTCGCGGTGCGCCGAGCGTGTATCCGCACCGGCGCCGGCTGCTCGAG GAGAACCGGCTCGAGGAGGCGCTCGCCTCGTTGGACGCGGGACTGGATCTCTGGCGAGGCC CGGCGCTGTCCACCGTACCGTGC GGCCGGGTGCTCGAAAGCAATATCGCGCACCTGGAAGAG CTGCGGCTTTTTGGAATGCAGCTCCGTATCGACGCGAATTGGCGGCTGGGCAGAATAGGGCC GATGATTCGGAACTCCGGTCCCTGGTAATTTTCGCATCCGCTGAACGAGACCCTGCACGCCAA ACTGATGGGCGCGCTCTGTGAGATGGGCAGGCGCGCCGAGGCGCTGGAATCGTATCGGAATC TCCGGCGGATACTGTCCGACGAAGTGGGGTGGATCCGACGCCGGAATCCAGCGTATGAC ATGGAATTCACCGGTGAGAAGGTGCTCGTGTAGCACCGGTCCGTGAACCGTGGAGGCC CTATGTCTCTTAAGTGTTCCCCTCCCTGCCTCGTGGTCCCTCACGCGCTCAGCTTTGGGCGCC CGGCTCGAGCGGCGGTGCAAGGGAGATGGGGTGC CGCTGGACGCGGGCGCCGGTGGATCCG GCATCGAGGGGTCCCGTATCGGCCTTCGAGCCTCCTTCGAGCCACGGGGCCGACGATGACGA CGACCACCGGACGAACGCATC </p>

*ermE**p

ATGGGGACCTCCTGGGGTGC GTTGGACCGCTGGATCCTACCAACCGGCACGATTGTGCCAC
AACAGCATCGCGGTGCCACGTGTGGACCGCGTCGGTCAGATCCTCCCCGCACCTCTCGCCAG
CCGTCAAGATCGACCGCGTGCACCA**CATATG**TTACCAATGCTTAATCAGTGAGGCACCTATCT
CAGCGATCTGTCTATTTTCGTTTCATCCATAGTTGCCTGACTCCCCGTCGTGTAGATAACTACGAT
ACGGGAGGGCTTACCATCTGGCCCCAGTGTGCAATGATACCGCGAGACCCACGCTCACCGG
CTCCAGATTTATCAGCAATAAACAGCCAGCCGGAAGGGCCGAGCGCAGAAGTGGTCCTGCA
ACTTTATCCGCCTCCATCCAGTCTATTAATTGTTGCCGGGAAGCTAGAGTAAGTAGTTCCGCCAG
TTAATAGTTTGCGCAACGTTGTTGCCATTGCTACAGGCATCGTGGTGTACGCTCGTCGTTTGG
TATGGCTTCATTCAGCTCCGTTCCCAACGATCAAGGCGAGTTACATGATCCCCCATGTTGTGC
AAAAAGCGGTTAGCTCCTTCGGTCCCTCCGATCGTTGTCAGAAGTAAGTTGGCCGCAGTGTTA
TCACTCATGGTTATGGCAGCACTGCATAATTCTCTTACTGTGCATGCCATCCGTAAGATGCTTTTC
TGTGACTGGTGAGTACTCAACCAAGTCATTCTGAGAATAGTGTATCGGGCACCAGTTGCTC
TTGCCCGCGTCAATACGGGATAATACCGCGCCACATAGCAGAACCTTTAAAAGTGTCTCATCT
GGAAAACGTTCTTCGGGGCGAAAACCTCTCAAGGATCTTACCCTGTTGAGATCCAGTTTCGATG
TAACCCACTCGTGCACCCAACCTGATCTTCAGCATCTTTTACTTTACCAGCGTTTCTGGGTGAG
CAAAAACAGGAAGGCAAAATGCCGCAAAAAGGGAATAAGGGCGACACGGAAATGTTGAATAC
TCATACTCTTCCTTTTTCAATCATGATTGAAGCATTATCAGGGTTATTGTCTCATGAGCGGATA
CATATTTGAATGCTCGAG

Table S3, related to STAR Methods section “Bioactivity testing of streptophenazines and MIC determination” Minimum Inhibitory Concentrations (MICs) ($\mu\text{g/mL}$) of oxo-streptophenazine A (**9**), streptophenazine C (**13**), streptophenazine A (**16**), and streptophenazine Q (**20**).

Strain	Compound 9	13	16	20
Group A <i>Streptococcus</i>	>50	>50	>50	2.5
<i>Acinetobacter baumannii</i> 5075	>50	>50	>50	40
<i>Klebsiella pneumoniae</i> 1100	>50	>50	>50	>40
MRSA TCH1516	>50	>50	>50	40

Table S4, related to STAR methods. Plasmids and strains used in this work

Plasmids	Description	Sources
pCAP03	TAR cloning and broad-host-range heterologous expression vector; <i>CEN6-ARS4, oriT, traJ, pUC ori, Kan^r, Apra^r, pADH1, URA3, TRP1</i>	Tang <i>et al.</i> , 2014
pSMM	Derivative of pCAP03 with 48 kb captured <i>spz</i> cluster	This work
pKDB01	Derivative of pCAP03 with 37.5 kb captured <i>spz</i> cluster	This work
pKDB02	Derivative of pKDB01 without TetR regulatory gene (<i>spz28</i>)	This work
pKDB01 Δ <i>spz3</i>	Derivative of pKDB01 without <i>spz3</i> (LysR type regulatory gene)	This work
pKDB02 Δ <i>spz3</i>	Derivative of pKDB02 without <i>spz3</i> (LysR type regulatory gene)	This work
pKDB01- <i>ermE</i> * <i>p-spz24</i>	Derivative of pKDB01 with <i>ermE</i> * promoter in front of <i>spz24</i> (LuxR-type regulatory gene), <i>Kan^r, Amp^r</i>	This work
pKDB03	Derivative of pKDB01 with <i>sp44-p21</i> cassette	This work
pKDB04	Derivative of pKDB03 with <i>actIp</i> cassette	This work
pKDB05	Derivative of pKDB04 with <i>ermE</i> * <i>p</i> cassette	This work
pKDB03 Δ <i>spz15</i>	Derivative of pKDB03 with <i>spz15</i> (putative adenylation protein-encoding gene) deleted	This work
pCAP03- <i>ermE</i> * <i>p</i>	Derivative of pCAP03 containing <i>ermE</i> * <i>p</i> cassette between XhoI and NdeI, <i>Kan^r, Amp^r</i>	This work
pCAP03- <i>actIp</i>	Derivative of pCAP03 containing <i>actIp</i> cassette between XhoI and NdeI, <i>Kan^r, Apra^r</i>	This work
pCAP03- <i>sp44-p21</i>	Derivative of pCAP03 containing <i>sp44-p21</i> cassette between XhoI, NdeI, <i>Kan^r, Apra^r</i>	This work
pIJ790	λ -RED (<i>gam, bet, exo</i>), <i>cat, araC</i> , rep101 ^{ts} , oriR101, P araBAD	Gust, 2003
pUB307	Self-transmissible plasmid that mobilizes other plasmids <i>in trans</i> for DNA transfer into hosts: RP4, <i>neo</i>	Flett, 1997
Strains	Description	
<i>Streptomyces</i>		
<i>Streptomyces sp.</i> CNB-091	Native producer of streptopenazines	Trischman <i>et al.</i> , 1994
<i>Streptomyces coelicolor</i> M1146	Host strain for heterologous expression derived from <i>S. coelicolor</i> M145: Δ <i>act, Δred, Δcpk, Δcda</i> .	Gomez-Escribano and Bibb, 2011
<i>S. coelicolor</i> M1146-pCAP03	Heterologous host containing empty pCAP03 as a control	This work
<i>S. coelicolor</i> M1146-pSMM	Heterologous host containing pSMM (<i>spz</i> BGC captured in 48 kb DNA fragment)	This work
<i>S. coelicolor</i> M1146-pKDB01	Heterologous host with integrated pKDB01 (37.5 kb captured <i>spz</i> cluster)	This work
<i>S. coelicolor</i> M1146-pKDB02	Heterologous host with integrated pKDB02 (Δ <i>spz28</i>)	This work
<i>S. coelicolor</i> M1146-pKDB01 Δ <i>spz3</i>	Heterologous host with integrated pKDB01 Δ <i>spz3</i>	This work
<i>S. coelicolor</i> M1146-pKDB02 Δ <i>spz3</i>	Heterologous host with integrated pKDB02 Δ <i>spz3</i>	This work

<i>S. coelicolor</i> M1146-pKDB01- <i>ermE</i> * <i>p-spz24</i>	Heterologous host with integrated pKDB01- <i>ermE</i> * <i>p-spz24</i>	This work
<i>S. coelicolor</i> M1146-pKDB03	Heterologous host with integrated pKDB03	This work
<i>S. coelicolor</i> M1146-pKDB04	Heterologous host with integrated pKDB04	This work
<i>S. coelicolor</i> M1146-pKDB05	Heterologous host with integrated pKDB05	This work
<i>S. coelicolor</i> M1146-pKDB03 Δ <i>spz15</i>	Heterologous host with integrated pKDB03 Δ <i>spz15</i>	This work
<i>Escherichia coli</i>		
DH10B	F- <i>mcrA</i> Δ (<i>mrr-hsdRMS-mcrBC</i>), Φ 80 <i>lacZ</i> Δ M15, Δ <i>lacX74 recA1 endA1 araD139 Δ (<i>ara leu</i>)7697 <i>galJ galK rpsL nupG</i> λ-. Storage and maintenance</i>	
BW25113	K-12 derivative: Δ <i>araBAD</i> , Δ <i>rhaBAD</i> ,	Datsenko and Wanner, 2000
BT340	DH5 α /pCP20, containing FLP recombinase	Cherepanov and Wackernagel, 1995
ET12567	F- <i>dam13</i> ::Tn9, <i>dcm6</i> , <i>hsdM</i> , <i>hsdR</i> , <i>recF</i> ,143 <i>zjj-202</i> ::Tn10, <i>galK2</i> , <i>galT22</i> , <i>ara-14</i> , <i>pacY1</i> , <i>xyl-5</i> , <i>leuB6</i> , <i>thi-1</i> , <i>tonA31</i> , <i>rpsL136</i> , <i>hisG4</i> , <i>tsx-78</i> , <i>mtl-1</i> , <i>glnV44</i> . Donor strain for conjugation between <i>E. coli</i> and <i>Streptomyces</i> in triparental mating	MacNeil et al., 1992
Other		
Group A <i>Streptococcus</i>	Clinical isolate used for bioactivity assays	Nizet lab,UCSD
<i>Acetobacter baumannii</i> 5075	Clinical isolate used for bioactivity assays	Nizet lab,UCSD
<i>Klebsiella pneumoniae</i> 1100	Clinical isolate used for bioactivity assays	Nizet lab,UCSD
MRSA TCH1516	Clinical isolate used for bioactivity assays	Nizet lab,UCSD
<i>S. cerevisiae</i> VL6-48N	MAT α <i>trp1</i> - Δ 1 <i>ura3</i> - Δ 1 <i>ade2</i> -101 <i>his3</i> - Δ 200 <i>lys2 met14</i> <i>cir</i> ^o , TAR cloning	Kouprina and Larinova, 2016

Table S5, related to STAR methods. Primers used in this work.

Primer name	Sequence (5' to 3')	Purpose
Frag1_F	GAGTAGCAGCACGTTTCCTTATATGTAGCTTTCGACAT ATGCATGAGCTGTCTCCTGGTGTGGTGGGCAGG	TAR cloning <i>spz</i> BGC
Frag1_R	CGACCTGCCCCAACTCGACGGGCTGGAAGT	TAR cloning <i>spz</i> BGC
Frag2_F	CACGATGCCCAGCAGCAGTCCCATGTCGTGG	TAR cloning <i>spz</i> BGC
Frag2_R	GCATCCAGTGCAGTACAGCCTCGCCGAGCG	TAR cloning <i>spz</i> BGC
Frag3_F	CCAGGTAATCCTCCAAGTCTCGCCGGTCCG	TAR cloning <i>spz</i> BGC
Frag3_R	CTCATAAGGATGCCTTCTGCGGGTGTGGAGACC	TAR cloning <i>spz</i> BGC
Frag4_F	CGATCGCCGCTGACCGACTGGAGCAGG	TAR cloning <i>spz</i> BGC
Frag4_R	GCTGCCGCTGAACGCCAACGGGAAGGTGGAC	TAR cloning <i>spz</i> BGC
Frag5_F	GAGGGCGGTCAGGGCAGGATTCACGGAATGC	TAR cloning <i>spz</i> BGC
Frag5_R	GGAGGCAGGCGCTACTGGACGTTCCCTCATCC	TAR cloning <i>spz</i> BGC
Frag6_F	CCATCATCTTGAGCTCTTCGTGACGACCATG	TAR cloning <i>spz</i> BGC
Frag6_R	GAACCAGGAGATCGCCGACGGCTCGTCT	TAR cloning <i>spz</i> BGC
Frag7_F	CTCGGCCTGGAATCACTTCAGATGATGCGCCTG	TAR cloning <i>spz</i> BGC
Frag7_R	GTTCTGATCGCCAGTTCCTGCAGAGCGTG	TAR cloning <i>spz</i> BGC
Frag8_F	GTCCTCCGACGAACATCAGGATGGAGCCG	TAR cloning <i>spz</i> BGC
Frag8_R+TetR	CTCGGTTTGACGCCTCCCATGGTATAAATAGTGGCTC GAGGCACAGGACGGCATCGCCCGAGCTGAGC	TAR cloning <i>spz</i> BGC
Frag8_R-TetR	CTCGGTTTGACGCCTCCCATGGTATAAATAGTGGCTC GAGGGTCCGTCCGGTCCCAGGAGAACCAGCTGATC GGCTTGCGGAGAGACCGGGCGGGGATCAGCCGAGG AGGGCGCGGCGTGTACGCCAATCGACTGGCGAGC GGCAT	TAR cloning <i>spz</i> BGC
Δ<i>spz3</i>_F	GCTCGTCGCACCGTCGCCCGCTCCTGAACCGGCC GCGGGGAAGGACCACGGTTCATGTGCAGCTCCATCA GCAAAAG	Delete <i>spz3</i> (LysR)
Δ<i>spz3</i>_R	GTGAAGGCGACGAACAGGAAGTG	Delete <i>spz3</i> (LysR)
Δ<i>spz3</i>_seq_F	GATCGAGCCCGCCGACGTG	Confirm <i>spz3</i> deletion
Δ<i>spz3</i>_seq_R	GTCGGGAGCGTGCGGCACAGTGCGCACGTCGGGCA GGACTGGCACGGTTCATATGGGGACCTCCTGGGGTG CGTTGG	Confirm <i>spz3</i> deletion
Upreg-<i>spz24</i>_F	CCCGATCGTGACGTGCGCGCCGTGCGCCCTAGG CCCTCCGGGCGGACGCTCGAGCATTCAAATATGTAT CCGCTCATG	Insert <i>ermE</i> * promoter in front of <i>spz24</i> (LuxR)
Upreg-<i>spz24</i>_R	CAGCAGGCTGCTCTTGCCGACAC	Insert <i>ermE</i> * promoter in front of <i>spz24</i> (LuxR)
Upreg-<i>spz24</i>_seq_F	GATGCGCGGTATGTGGGAGCGC	Sequencing to confirm insertion
Upreg-<i>spz24</i>_seq_R	ATTATACATATGTGCAGTTCGAAGTTCCTATTCTCTAG AAAGTATAGGAATTCGGTTCATGTGCAGCTCCATCA GCAAAAG	Sequencing to confirm insertion
<i>aac(3)IV</i>+FRT_F	ATTATAAAGCTTGAAGTTCCTATACTTTCTAGAGAATA GGAACCTCTCAGCCAATCGACTGGCGAGCGGCATCG	Amplify <i>aac(3)IV</i> gene with FRT sites for cloning between pET28a NdeI and HindIII sites
<i>aac(3)IV</i>+FRT_R	GACGCCTCCCATGGTATAAATAGTGGCTCGAGGGTG AACCGATCTCCTCGTTAGGGTC	Amplify <i>aac(3)IV</i> gene with FRT sites for cloning between pET28a NdeI and HindIII sites
<i>sp44</i>_F	TCTAGAGAATAGGAACCTTCAACTGCACATATGATAG TGGCTGTTACATTTCGAACCGTCTCTG	Amplify <i>sp44</i> promoter with homology to pCAP03 and <i>aac(3)IV</i>
<i>sp44</i>_R	CTATTCTCTAGAAAGTATAGGAACCTTCAAGCTTGTCT CGAGTGTGCGGGCTCTAACACGTC	Amplify <i>sp44</i> promoter with homology to pCAP03 and <i>aac(3)IV</i>
<i>p21</i>_F		Amplify <i>p21</i> promoter with homology to pCAP03 and <i>aac(3)IV</i>

<i>p21_R</i>	AGCACGTTCTTATATGTAGCTTTGACATATGGTCC GTACCTCCGTTGCTCGACTAGAT	Amplify <i>p21</i> promoter with homology to pCAP03 and <i>aac(3)/IV</i>
<i>aac(3)/IV+FRT_sp44_pCAP03_F</i>	CAGAGACGGTTTGAATGTGAACAGCCACTATCATAT GTGCAGTTTGAAGTTTCTATTCTCTAGA	Amplify <i>aac(3)/IV</i> +FRT with homology to <i>sp44</i> promoters
<i>aac(3)/IV+FRT_p21_pCAP03_R</i>	ACGTGTTAGAGCCCCGACACTCGAGCAAAGCTTGAA GTTCTATACTTTCTAGAGAATA	Amplify <i>aac(3)/IV</i> +FRT with homology to <i>p21</i> promoter
<i>sp44+p21_cas_insert_F</i>	TGGTCGGCGTCTCTGTTGCTGAACTGATGCTCTGG GCGACGTCGGGCATCTCGAGGGTGAACCGATCTCCT CGTTAGG	Amplify <i>sp44-p21</i> cassette with homology sequences for targeted insertion into <i>spz</i> BGC
<i>sp44+p21_cas_insert_R</i>	CCGGGGACGTGGGTGCCGAGTCCGGCGAGGTAGGT GTTCTCGAACTTCATCATATGGTCCGTACCTCCGTTG CTCGAC	Amplify <i>sp44-p21</i> cassette with homology sequences for targeted insertion into <i>spz</i> BGC
<i>sp44+p21_cas_insert_seq_F</i>	GTGAACAGGAGATGCCGGGTG	Confirm <i>sp44-p21</i> cassette insertion
<i>sp44+p21_cas_insert_seq_R</i>	GCCTCGTACCAGCCTTCCTG	Confirm <i>sp44-p21</i> cassette insertion
<i>actI_p_F</i>	CCGGTCCGTGAACGCGGTGGAGCCCTATGTCTCTTA AGTGTTCCTCCCTCCCTGCC	Amplify <i>actI</i> promoter with homology to <i>actII-ORF4</i> and pCAP03
<i>actI_p_R</i>	CACGTTCTTATATGTAGCTTTGAGATGCGTTTCGTC CGGTGGTCGTCGTCA	Amplify <i>actI</i> promoter with homology to <i>actII-ORF4</i> and pCAP03
<i>actII-ORF4_F</i>	CCGCTCGCCAGTCGATTGGCTGACATATGCCACTGC CTCTCGGTAATAATCC	Amplify <i>actII-ORF4</i> with homology arms for assembly with <i>aac(3)/IV</i> and <i>act1p</i>
<i>actII-ORF4_R</i>	GGCAGGGAGGGGAACACTTAAGAGACATAGGGCTC CACCGCTTACGCGACCGG	Amplify <i>actII-ORF4</i> with homology arms for assembly with <i>aac(3)/IV</i> and <i>act1p</i>
<i>aac(3)/IV+actI_p_F</i>	GACGCCTCCCATGGTATAAATAGTGGCTCGAGGGTT CATGTGCAGCTCCATCAGCAAAG	Amplify <i>aac(3)/IV</i> with homology to <i>actII-ORF4</i> and pCAP03
<i>aac(3)/IV+actI_p_R</i>	GGATTTTACCGAGAGGCAGTGGCATATGTCAGCCAA TCGACTGGCGAGCGG	Amplify <i>aac(3)/IV</i> with homology to <i>actII-ORF4</i> and pCAP03
<i>actI_p_cas_insert_F</i>	GGGTCCGCGTCGTACGGAAGGTCAAGAATCTTCGGG TCGGCGGAAGCCACGATGCGTTCTCCGGTGGTCTG TCGTC	Amplify <i>actI_p</i> cassette with homology sequences for targeted insertion into <i>spz</i> BGC
<i>actI_p_cas_insert_R</i>	GTGCTCGAATGTCCCATACCCCAAGACGTAGAAGT TCTCTGGAGGAACGACTCGAGGGTTCATGTGCAGCT CCATCAG	Amplify of <i>actI_p</i> cassette with homology sequences for targeted insertion into <i>spz</i> BGC
<i>actI_p_cas_insert_seq_F</i>	CTCGAGCCAGTAGCGGGATC	Confirm <i>actI_p</i> cassette insertion within <i>spz</i> BGC
<i>actI_p_cas_insert_seq_R</i>	CATTGTGCACGGTCCACCG	Confirm <i>actI_p</i> cassette insertion within <i>spz</i> BGC
<i>ermE*p_F</i>	GTGCCTCACTGATTAAGCATTGGTAACATATGTGGTG CACGCGGTGATCTTGACGGCTG	Amplify <i>ermE*</i> promoter with homology arms for assembly with <i>bla</i> and pCAP03
<i>ermE*p_R</i>	CAGCACGTTCTTATATGTAGCTTTGAATGGGGACC TCCTGGGGTGCCTGGACC	Amplify <i>ermE*</i> promoter with homology arms for assembly with <i>bla</i> and pCAP03
<i>bla+ermE*p_F</i>	TGACGCCTCCCATGGTATAAATAGTGGCTCGAGCATT CAAATATGTATCCGCTCATGAGA	Amplify <i>bla</i> with homology to <i>ermE*p</i> and pCAP03 for assembly
<i>bla+ermE*p_R</i>	CAGCCGTCAAGATCGACCGCGTGCACCACATATGTT ACCAATGCTTAATCAGTGAGGCAC	Amplify <i>bla</i> with homology to <i>ermE*p</i> and pCAP03 for assembly
<i>ermE*p_cas_insert_F</i>	CAGGGTCGCCGTGACGGCGAACTGGAGGCGGGCGA GGTCGACGACGTCCACATGGGGACCTCTGGGGTG CGTTGG	Amplify <i>ermE*p</i> cassette with homology sequences for targeted insertion into <i>spz</i> BGC
<i>ermE*p_cas_insert_R</i>	CTTCCACAGCCACGCCCGCGCCCTCCTCGGCTGATC CCCGCCCGGTCTCTCTCGAGCATTCAAATATGTATCC GCTCATG	Amplify <i>ermE*p</i> cassette with homology sequences for targeted insertion into <i>spz</i> BGC
<i>ermE*p_cas_insert_seq_F</i>	GTGAGGTGCTCGTGCGCAG	Confirm <i>ermE*p</i> cassette insertion within <i>spz</i> BGC
<i>ermE*p_cas_insert_seq_R</i>	GTCCTGCCCGAGCAGTATGTCC	Confirm <i>ermE*p</i> cassette insertion within <i>spz</i> BGC
Δ <i>spz15_F</i>	ACGGGTTCCAGGCGGCCGGAGACGACGTCCTCGCG CAGCGCCCCCGGTCTCAGCCAATCGACTGGCGAG CGGCAT	Delete <i>spz15</i> (adenylation protein)
Δ <i>spz15_R</i>	GGCGGGTCTCGGCTCCAGCCCTGAGCCCGGGGCG ACTCTCCGCCTCCGGTTCATGTGCAGCTCCATCAG CAAAG	Delete <i>spz15</i> (adenylation protein)
Δ <i>spz15_seq_F</i>	GTCAGGGCAGGATTCACGGAATG	Confirmation of <i>spz15</i> deletion
Δ <i>spz15_seq_R</i>	GTACGAACTCCGTTTCGACAGGGTG	Confirmation of <i>spz15</i> deletion

Table S6, related to STAR methods. Primers used for RT-PCR.

Primer name	Sequence (5' to 3')	Primer name	Sequence (5' to 3')
<i>hrdB_F</i>	CCTCCGCCTGGTGGTCTCG	<i>hrdB_R</i>	AACTTGTAGCCCTTGGTGTAGTCGAAC
<i>spz1_F</i>	CGTCACCGAGAGCAGCCACAGCG	<i>spz1_R</i>	GTTGCTCCAAGCACTGCGACTGCC
<i>spz2_F</i>	CGATCAGGGAGAAGTTCACGACG	<i>spz2_R</i>	CTGCACTTCTGTTCGTGCGCCTT
<i>spz3_F</i>	GGACATACTGCTCGGGCAGGAC	<i>spz3_R</i>	CAGTTGGAGTACTTCTCGCGG
<i>spz4_F</i>	GCCTGATCAGCACCACGTCGG	<i>spz4_R</i>	GAACTTCGCGTACGACCTGTGCG
<i>spz5_F</i>	CGATCTGCTCGAAGTCGAACAGCC	<i>spz5_R</i>	GACACCATGTACGCGTCACTCATCC
<i>spz6_F</i>	GTGCAGAGTCTCCAGAACGTCGTGG	<i>spz6_R</i>	CACACCGCAGGTCGTGTTACCTTC
<i>spz7_F</i>	GCGAGGCTGTACTCGCACTGGATG	<i>spz7_R</i>	GTCTCCGAATACGCGCTGGGCAC
<i>spz8_F</i>	GAACGTGGTCACCGAATGTCCAG	<i>spz8_R</i>	GACCGAGCTCGACCTGTACTGCTTC
<i>spz9_F</i>	CTCGAAGAAGCTCTCGTCCGGTTCCG	<i>spz9_R</i>	CTTGAGGACTGGAAGCCGTACGGG
<i>spz10_F</i>	GTGGTCTGAACAGGTCGATCAGCTC	<i>spz10_R</i>	GAGAAGGATTCCTGACCCTCGTCCG
<i>spz11_F</i>	GTCTTGCTCCGCATCAGGAATTCC	<i>spz11_R</i>	GATCACCTCGACATGATCGTCGAC
<i>spz12_F</i>	CTCACGTGGTGCATGACCACCTGG	<i>spz12_R</i>	GACATCGTCGGCGTCTTCGAGAG
<i>spz13_F</i>	CGATGATCGCCTGGTGGGATCTC	<i>spz13_R</i>	GTGGTGGCGACGACGTCTGAAC
<i>spz14_F</i>	CGAGATGACGTACGTCCACCACGC	<i>spz14_R</i>	GCTTCCAGACGGCCTACCGGCATC
<i>spz15_F</i>	CACGTCCGCCAGGTGGTAGGTGAG	<i>spz15_R</i>	GAGGTGACGGTGTGGTTCTCGGTG
<i>spz16_F</i>	CTCCACCCTGTCTGAACGGAGTTC	<i>spz16_R</i>	GACCGCCGATATCACCGTCCCCTTTC
<i>spz17_F</i>	GCGAGAACATCCTGCTGCGCCAC	<i>spz17_R</i>	CTCCTATGTGGTTCGTGACGCCTTC
<i>spz18_F</i>	CGGTTGATCTCGACGACACCGAGC	<i>spz18_R</i>	CTGGACTCCGCGATCCTCATCCGC
<i>spz19_F</i>	CACCAGGAAGGTGCGGATGTCTGTG	<i>spz19_R</i>	CTGGAATCCCCGCATCCACCCG
<i>spz20_F</i>	CAGGTACTTGTCTCCGACGCCCGAC	<i>spz20_R</i>	CTCGACTACGAGCTGCCGATGCTG
<i>spz21_F</i>	CTGGTCTGAACGGGTTTCATGAACCTCG	<i>spz21_R</i>	GCTGACCCGGCATCTCTGTTCAC
<i>spz122_F</i>	CTGGTCAACACGGAGAAGGCCGTC	<i>spz22_R</i>	GATTCCAGATCGAAGCCGATCGGATC
<i>spz23_F</i>	CTAGTCGCCGGAGAGGAGCTGCGC	<i>spz23_R</i>	GTATCGCGTCTCTGCTCCTCAGC
<i>spz24_F</i>	CAACTCCTCGTAGGCGTGCGACAC	<i>spz24_R</i>	CTACCGTCTCTCCTGCGGCTGGATC
<i>spz25_F</i>	GGTGGTGTAGGTGAAGTCCACCCAC	<i>spz25_R</i>	GAAGTCGTACGACCAGAGCGGTGTG
<i>spz26_F</i>	CGAACATCAGGATGGAGGCCGTCAG	<i>spz26_R</i>	GCTGGAGTACTTCTGGTGGGGCTCG
<i>spz27_F</i>	CGTCGAGCCGGAGGTCTGAAGAAG	<i>spz27_R</i>	CAAGGACATCTGGCCGAGTGCTG
<i>spz28_F</i>	CGATCAGCTGGTCTCTCTGGGACCG	<i>spz28_R</i>	GACTAGGGCCAGGCCAAGGAGCG