Metathramycin, a new bioactive aureolic acid discovered by heterologous expression of a metagenome derived biosynthetic pathway


Bacterial natural products have been a rich source of bioactive compounds for drug development, and advances in DNA sequencing, informatics and molecular biology have opened new avenues for their discovery. Here, we describe the isolation of an aureolic acid biosynthetic gene cluster from a metagenome library derived from a New Zealand soil sample. Heterologous expression of this pathway in Streptomyces albus resulted in the production and isolation of two new aureolic acid compounds, one of which (metathramycin) possesses potent bioactivity against a human colon carcinoma cell line (HCT-116, IC_{50} = 14.6 nM). As metathramycin was a minor constituent of the fermentation extract, its discovery relied on a combination of approaches including bioactivity guided fractionation, MS/MS characterisation and pathway engineering. This study not only demonstrates the presence of previously uncharacterised aureolic acids in the environment, but also the value of an integrated natural product discovery approach which may be generally applicable to low abundance bioactive metabolites.

Introduction

Aureolic acids are a family of glycosylated aromatic polyketides that have potent bioactivities against human cancer cells and Gram-positive bacteria.\(^1,2\) All naturally occurring aureolic acids have been discovered from soil or marine actinomycete bacteria, and the biosynthetic gene clusters (BGCs) for two of these have been described: mithramycin (Fig. 1, 1) from Streptomyces argillaceus,\(^1,3\) and chromomycin A\(_3\) (Fig. 1, 2) produced by Streptomyces griseus subsp. griseus.\(^4\) Although aureolic acids have been approved for clinical use (e.g. mithramycin for testicular carcinoma and myeloid leukemia), their clinical applications remain limited due to significant off target toxicity.\(^5,6\)

The potent anticancer bioactivity of mithramycin and chromomycin A\(_3\) has fuelled a search for new congeners. The search for these mithramycin analogues ("mithralogs") has predominantly taken the form of gene knockout or combinatorial biosynthesis based on the known biosynthetic genes and pathways for mithramycin and chromomycin A\(_3\).\(^6\) There are two main regions of the compound structure that have been targeted for diversity generation. The first is the significant sugar decoration of the polyketide core (premithramycinone, Fig. 1, 3), which can be modified by deletion or complementation with different sugar biosynthesis pathways and/or glycosyltransferase enzymes Mtm/\(^6\) CmmGI-GIV.\(^7,8\) The second is the aliphatic tail which is generated by oxidative cleavage of the fourth ring in the tetracyclic precursor (premithramycin B, Fig. 1, 4) by Mtm/CmmOIV, and subsequent sidechain reduction by Mtm/CmmW. Gene knockout of the sidechain reductase has been demonstrated to produce a set of analogue compounds with distinct toxicity profiles.\(^2,4\)

Microbial natural products discovered via strain isolation and bioactivity guided fractionation have been a phenomenally rich source of medically relevant compounds.\(^9\) Another method for uncovering novel chemistry from microbes is the discovery of new biosynthetic pathways from environmental microbes. Gene-first natural product discovery by metagenomic analysis has produced numerous novel bioactive secondary metabolites in recent years,\(^10-15\) providing access to the biosynthetic capacity of microbes independent of our ability to culture these organisms.\(^16-19\) This approach has proven particularly fruitful when applied to aromatic polyketides, leading to the discovery of improved congeners of clinically relevant compounds.\(^20\) In this study, we constructed and screened a soil metagenome cosmid library for ketosynthase alpha (KS\(_\alpha\)) bacterial type II polyketide genes. Sequencing of the recovered library cosmid

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clones resulted in the partial sequence of an aureolic acid biosynthetic gene cluster with low sequence identity to previously reported clusters. Recovery and reconstitution of the full gene cluster from two overlapping cosmid clones permitted heterologous expression in *Streptomyces albus*, ultimately resulting in the discovery of two new aureolic acid compounds that we have named premetathramycin (Fig. 1, 5) and metathramycin (Fig. 1, 6).

Results and discussion

Discovery and recovery of novel gene cluster from a soil metagenomic library

Soil collected in Half Moon Bay, Kaikōura, New Zealand, was used as the source of microbial diversity to construct a metagenome library. High molecular weight environmental-DNA (eDNA) was extracted, purified, and cloned into the cosmid
vector pWEB::tnc using in-house prepared lambda phage packaging extract. The resulting library of >10 million cosmids clones, each containing 30–40 kb of metagenome DNA, was arrayed over four 96 well plates. These arrayed wells were then screened by PCR with degenerate primers targeting bacterial KSz genes, using previously established serial dilution plating protocols.

The resulting cosmids clones were pooled and sequenced as a single library (Illumina HiSeq, PE100 bp chemistry). Following assembly with SPAdes, contigs were matched to individual cosmid insert end sequences by Sanger sequencing, and the assembled metagenome insert sequences analysed with antiSMASH v4. One of the recovered cosmids had 68% gene cluster similarity by clusterBLAST to the aureolic acid pathway producing mithramycin and 58% to chromomycin A₃. Notably, however, the individual amino acid sequences encoded by the genes in the partial gene cluster had a relatively low homology to the known aureolic acid biosynthesis pathways, and the gene arrangements were also unique. The recovered cosmid contained only a partial gene cluster sequence, as it was apparent that the mtrx gene homologue involved in self resistance and DNA repair was truncated at the cosmid insert boundary. Other key resistance genes involved in compound efflux in mithramycin and chromomycin A₃ pathways were not present in the recovered cosmid. Reconstitution of the biosynthetic pathway was achieved by re-screening the soil metagenome library using primers targeting each end of the initially recovered cosmid. Cosmids extending the recovered sequence in both directions were recovered, however only the sequence that complemented the truncated mtrx homologue appeared to contain additional biosynthetic genes. Sequence data for the complete metathramycin BGC has been deposited in GenBank with the accession number MW512268.

The two cosmids containing putative biosynthetic genes were then used to construct a complete and contiguous biosynthetic pathway as a single BAC clone. This was achieved using a modified version of previously published protocols for transformation associated recombination (TAR) in yeast and in which the insert sequences from the two cosmids were liberated from the pWEB::tnc vector using in vitro Cas9 digestion with sgRNA targeted to the vector edges. Double-stranded DNA breaks near assembly sites has been demonstrated to increase recombination frequency in TAR, and the use of targeted Cas9 digestion resulted in free DNA ends within 200 bp of recombination sites. TAR was performed in a previously described S. cerevisiae strain deficient in non-homologous end joining to reduce non-recombinant background colonies, and in a single transformation all emergent colonies (12) analysed tested positive for correct assembly with multiple PCR tests along the BGC length. The reconstituted BGC was then verified by DNA sequencing, and was annotated using results from antiSMASH analysis, as well as BLAST search comparisons of the encoded genes to mithramycin and chromomycin BGCs (Fig. 2A and Table 1). This gene cluster has been assigned the name metathramycin (MMY), as a reference to both the discovery of this gene cluster in a metagenome library, and the gene cluster level similarity to the mithramycin BGC (Table 2).

The metathramycin gene cluster has a length of 42,302 bp, similar to mithramycin (42,374 bp) and chromomycin A₃ (42,074 bp). Of the genes that were assigned to homologous functions, the mean percentage amino acid identity of the products was 60.7% to mithramycin and 52.7% to chromomycin A₃. In general, the highest amino acid identities in the MIBiG database for most genes in the metathramycin pathway were with genes from the two aureolic acid gene clusters, however this was notably not the case for the first six enzyme functions (core polyketide biosynthesis by P, K, S, and cyclisation/aromatisation by X, Y, and Q). Only Q had its nearest homologue in chromomycin A₃, while all other genes aligned to other, non aureolic acid type II PKS biosyntheses.

Fig. 2 Discovery and heterologous expression of an aureolic acid biosynthetic gene cluster from a soil metagenome library. (A) Comparison of our metagenome derived aureolic acid cluster (MMY cluster) to the known clusters for mithramycin and chromomycin A₃. Genes are coloured based on broad function classifications. Amino acid identities for alignments of homologous genes are provided in Table 1. (B) HPLC traces for extracts from S. albus harbouring the MMY biosynthetic gene cluster (S. albus::MMY, red) and an empty vector control (S. albus, blue). Metabolite peaks with the characteristic aureolic acid UV absorbance spectrum (inset, green) are indicated with an asterisk (*). Compound 5 is the major metabolite peak, with a retention time of 13 min. Select minor metabolite peaks were identified by EIC of the LCMS trace; compound 6 is a minor metabolite with retention time 9.4 min, 4-demethylpremithramycinone 7.3 min and premithramycinone (3) peak at 9.8 min. The retention time on the reversed-phase C₁₈ column is shown on the X-axis, and absorbance at 420 nm on the Y-axis.
Comparison of the annotated genes within the metathramycin BGC to those of mithramycin (MMY) and chromomycin A₃ (CMM) biosynthetic gene clusters indicated that all of the genes necessary for biosynthesis of a tetracyclic aureolic acid aglycone premithramycinone (3) (mmyP, K, S, MI, OII, TI, TII) were present. Also present were genes for the subsequent methylation and glycosylation to a tetracyclic aureolic acid aglycone premithramycinone (3).

### Table 1
Gene cluster comparison of aureolic acid biosynthesis genes: homologous genes between the metathramycin (MMY), mithramycin (MTM) and chromomycin A₃ (CMM) biosynthetic gene clusters are indicated, with percentage amino acid identities of Mmy proteins to Mtm and Cmy proteins, respectively.

<table>
<thead>
<tr>
<th>Metathramycin BGC</th>
<th>Mithramycin BGC</th>
<th>Chromomycin A₃ BGC</th>
<th>Proposed function</th>
<th>% aa identity (Mmy:Mtm/Cmm)</th>
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<tr>
<td>CmmA</td>
<td>MtmA</td>
<td></td>
<td>O-acyltransferase</td>
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</tr>
<tr>
<td>CmmC</td>
<td>MtmC</td>
<td>CmmC</td>
<td>AdoMet synthetase</td>
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<td>CmmD</td>
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<td>CmmF</td>
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<td>Adenosylhomocysteinate</td>
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<td>Kethe synthase β</td>
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<td>O-methyltransferase</td>
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<td>Kethsynthe ζ</td>
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<td>CmmQ</td>
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<td>CmmX</td>
<td>Cyclase</td>
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<tr>
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<td>Cyclase</td>
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<td>ATP-binding protein</td>
<td>62.4/48.8</td>
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<td>MmraX</td>
<td></td>
<td></td>
<td>UV-repair system</td>
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### Table 2
Bioactivity assay results of antimicrobial and cytotoxicity assays.

<table>
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<tr>
<th>E. coli ΔtolC</th>
<th>Tetracycline</th>
<th>Nystatin</th>
<th>Mithramycin A (1)</th>
<th>Metathramycin (6)</th>
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<tbody>
<tr>
<td>&gt; 128</td>
<td>0.125</td>
<td>ND</td>
<td>&gt; 32</td>
<td>ND</td>
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<td>B. subtilis E168</td>
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<td>1</td>
<td>ND</td>
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<tr>
<td>S. cerevisiae ΔPDR</td>
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<td>ND</td>
<td>0.5</td>
<td>32</td>
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<tr>
<td>HCT-116</td>
<td>1.91</td>
<td>ND</td>
<td>0.013</td>
<td>0.015</td>
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Antimicrobial minimum inhibitory concentration (MIC) assays were conducted for 5 with positive controls of tetracycline (antibacterial), nystatin (antifungal) and mithramycin A (Gram-positive antibacterial). Tumour cell cytotoxicity was performed with mithramycin A as a positive control agent. The data in the table were determined across at least three independent experiments. MIC values (µg mL⁻¹) are given for E. coli, B. subtilis and S. cerevisiae. IC₅₀ values (µM) are given for HCT-116. ND, not determined.
premithramycin B (4) analogue (mmyMII, GI-GIV) and the final ring opening and sidechain reduction characteristic of aureolic acid biosynthesis (mmyOIV, W). There were four glycosyltransferase genes present, indicating at least four sugars are likely to be present in the final product of the biosynthetic pathway. In previously described aureolic acids, the identity, configuration and connectivity of these sugars differed significantly. All known aureolic acids contain four to six sugars attached as two oligosaccharide chains composed of combinations of D-olivose, D-oliose, D-mycarose and L-chromose B sugars. The metathramycin gene cluster lacked the required genes for chromomycin A₃ style sugars - the epimerase (CmmF) and subsequent reductase (CmmUIII) enzymes required for the production of l-chromose B, and methyl- and acetyltransferase enzymes for further decoration (CmmIII, A). The lack of a sugar epimerase gene indicates that all glycosylated metabolites of the metagenome pathway expression should only contain D-sugars. The metagenome cluster contained homologues indicative of a full sugar biosynthetic pathway for D-olivose, D-oliose and D-mycarose. The metagenome cluster contains a few notable genes that differ from either of the known aureolic acid biosynthesis pathways: The third oxygenase in the pathway (O3) does not have significant sequence homology to MtmmOIII (chromomycin lacks a third oxygenase). The O3 gene has some homology to the DutO3 monooxygenase in dutomycin biosynthesis (51.6% amino acid identity), as the nearest homologue in the MIBIG database. The metagenome gene cluster contains a HAD family hydrolase gene (Hy), which shares low-level identity with an empty vector control in 50 mL R5a medium containing 1 g Dialion HP-20 resin. Following 7 days of cultivation, the resin was washed, eluted with methanol and the resulting crude extracts were analysed by HPLC and LC-MS. Peaks potentially arising from aureolic acids were detected by monitoring absorbance at 420 nm, and these were subsequently confirmed by examining a full absorbance spectrum and comparing this to the characteristic aureolic acid UV signature of a mithramycin standard. The NMR and UV/vis spectral data was consistent with the 13C chemical shifts are consistent with other chromocyclin containing congeners premithramycin B and prechromomycin B. The MS/MS spectrum at low collision energy (20.0 eV) further suggested this tetracyclic structure as opposed to the tricyclic, ring opened structure with a C-3 aliphatic sidechain that is present in many other aureolic acids. This aliphatic sidechain is the most readily fragmentable site of the metabolite and its loss is clearly observable in the MS/MS at low collision energies for both mithramycin (Fig. S1, ESIF) and metathramycin (Fig. S3, ESIF), however
compound 5 shows no such fragmentation (Fig. S2, ESI†). The relative configuration of C-4 and C-5 was determined through interpretation of the ROESY data. Mutual NOE correlations between OCH3-21, H-5 and H-6a suggested they were on the same side of the molecule, corroborated by the relatively small coupling constant between H-5 and H-6a (J = 4.2 Hz) which indicated a syn relationship. For H-6b, the large coupling constant to H-5 (J = 11.0 Hz) and lack of NOE correlations to OCH3-21 further supported this assignment. The relative configuration of C-18 could not be determined from this spectroscopic evidence alone.

The carbohydrate moieties of compound 5 were assigned and characterised using a combination of COSY, HMBC, ROESY and HSQC-TOCSY NMR experiments. The anomeric resonance of sugar A, H-A1 (δH 5.57, dd, J = 2.2, 9.6), showed an HMBC correlation to C-11 (δC 160.4) thus connecting it to the aglycone core directly, while scalar 1H–1H coupling constants and ROESY correlations supported a mycarose sugar configuration (Fig. 3). The original reported structure of chromocyclomycin also contained a C-11 mycarose glycoside, configuration (Fig. 3). The original reported structure of compound 5 did possess moderate/strong Gram-positive antibiotic activity (B. subtilis, MIC 2 µg mL⁻¹), however a mithramycin A standard had an MIC of 0.004 µg mL⁻¹, indicating 500-fold greater Gram-positive antibacterial potency relative to 5. Compound 5 also possessed moderate cytotoxic activity against the HCT-116 cancer cell line, with an IC50 of 1.91 µM. However, mithramycin A was again far more potent, with an IC50 of 0.017 µM.

Chromocyclomycin has only been described in three research publications, where the biological activity is either not defined or it is stated to be “biologically inactive”. Another analogous compound, the tetracyclic aureolic acid precursor premithramycin B, has also been demonstrated to possess no detectable antibiotic activity (no inhibition of Micrococcus luteus growth up to 50 µg in an agar spot bioassay). However, 5 has here been demonstrated to possess Gram-positive antibiotic activity comparable to that of tetracycline. Likewise, 5 possessed moderate cytotoxicity in the HCT-116 cell line. These results indicate that 5 possesses not only a novel chemical structure, but also a novel bioactivity profile.

Bioactivity guided fractionation and discovery of metathramycin

The potent bioactivities of aureolic acid family compounds are exclusively found in those with a tricyclic core and aliphatic tail motif (1, 2), which are generated from tetracyclic precursors by characterised ring opening pathway of two enzymes MtmOV/W/CmmOV,W. The metathramycin BGC contains gene homologues for these enzymes, leading us to speculate that we might have been missing the final product of the pathway. As known mature aureolic acid compounds are highly active Gram-positive antibiotics, we conducted bioactivity-guided fractionation of our fermentation extract to find this putative missing metabolite.

A sample of the crude fermentation extract was fractionated over an HPLC gradient, with fractions testing positive by agar zone of inhibition assays to the Bacillus subtilis 168 test strain being pooled and re-fractionated at a shallower gradient. Three rounds of this process culminated in a sample containing an ion with m/z 967.4186 [M – H]⁻ that possessed potent bioactivity (6, corresponding to C47H68O21 calculated 967.4180, Δ = 0.6 ppm). Notably, the mass difference between 5 and 6 (976.3928–968.4256 = 7.9672 Da) is the same mass difference between premithramycin B and mithramycin (1092.4414–1084.4727 = 7.9687 Da), indicating a similar biochemical transformation.
The production of 6 in \textit{S. albus}:MMY culture was extremely low, and the amount isolated from 7 L of culture was insufficient to allow use of standard quantification methods. The purified sample was instead quantified by comparison to an injection standard of mithramycin by HPLC and measurements of UV absorption. As mithramycin and the predicted structure of 6 carry identical chromophores, the absorbance of the mithramycin standard was used to quantify the sample of 6. After normalisation for the mass difference and assuming the extinction coefficient difference is negligible, the calculated quantity of compound 6 was 7.2 \(\mu\)g.

This low yield (~1 \(\mu\)g L\(^{-1}\)) precluded structure elucidation by NMR, however MS/MS fragmentation analysis was conducted of 6 and the major biosynthesis product compound 1 against an authentic sample of mithramycin (Fig. 4). Compound 6 fragmented similarly to the mithramycin standard, with some of the fragmented ions shifted by a mass corresponding to the loss of a deoxy sugar and the gain of a methyl group, consistent with the mass difference between mithramycin and 6. The observed fragmentation pattern in both cases initiated with the loss of the aliphatic sidechain (daughter ion \(m/z\) 820 or 935) at 20.0 eV, then successive loss of deoxy sugars up to 60.0 eV, leaving the tricyclic core at \(m/z\) 269. The observed mass and fragmentation patterns of 6 are consistent with a “ring opened” aureolic acid. This fragmentation pathway was consistent with the gene cluster analysis, where the presence of OIV and W homologues indicate a ring opening step should occur in the biosynthesis pathway as for mithramycin and chromomycin A\(_3\). Compound 5 did not follow the same fragmentation pattern as the mithramycin standard (Fig. S2, ESI\(^+\)), potentially due to the lack of an aliphatic tail where the initial fragmentation occurs (at 20.0 eV onwards). Instead, the parent ion of 5 remained intact at 20.0 eV, with only minor fragmentation at 40.0 eV, followed by seemingly disordered breakdown at 60.0 eV, from which three of the conserved core masses observed during fragmentation of both mithramycin and compound 6 could be identified (Fig. S2, ESI\(^+\)).

The metathramycin BGC contained homologues of both \textit{mtmOIV} and \textit{mtmW}. These are believed to encode enzymes that catalyse oxidative cleavage of the fourth ring in the tetracyclic core of premithramycin, and reduction of the resulting sidechain, respectively, to yield mithramycin (Fig. 5A).\(^{40,43}\) To further test the hypothesis that compound 6 was indeed the product of enzymatic opening of the fourth ring of 5, we...
amplified the mmyOIV and mmyW genes from our pathway and cloned these into the integrative vector pIJ10257 as a bi-cistronic unit under control of the constitutive promoter ermE*. The PCR primers were designed to include a unique Shine–Dalgarno sequence as a point of homology between the two genes to complete the assembly and allow for expression from a single promoter sequence. The resulting construct was integrated into the S. albus strain harbouring the complete biosynthetic pathway, and production of 6 quantitated using LC-MS. We found that over-expression of the putative ring opening enzymes resulted in a 1.9-fold increase in 6 production compared to the base expression strain (Fig. 5B), providing further evidence to support the hypothesis that compound 5 is not the (sole) final product of the biosynthesis pathway, and that compound 5 may be converted to compound 6 by the same mechanism as conversion of premithramycin B to mithramycin. Given that 6 appears to be a direct conversion product afforded by ring opening and side chain reduction of 5, it is reasonable to assume that the configuration and connectivity of sugars is conserved between these two compounds.

Pleasingly, the 7.2 μg that was isolated was sufficient to conduct a bioactivity assay. When tested against the human tumour cell line HCT-116, metathramycin demonstrated potent cytotoxicity comparable to that of mithramycin (Fig. 5C: mithramycin 1 IC50 = 13.4 nM, metathramycin 6 IC50 = 14.6 nM over 4 biological replicates). Although there was insufficient compound to perform initial toxicity studies, the promising bioactivity of metathramycin against tumour cells suggests that further analysis of this molecule is warranted.

**Concluding remarks**

While the task of generating bioactive mithralogs in the laboratory continues, it is apparent that there are aureolic acids in nature yet to be discovered. Metagenome mining studies in recent years have uncovered polyketide biosynthetic pathways that phylogenetically map to aureolic acid biosynthesis. This study is the first to characterise the aureolic acid chemical output of a metagenome-derived BGC.

Heterologous expression of this biosynthetic gene cluster resulted in a major fermentation product (5) that was not the final product of the biosynthetic pathway (6). The molecular formula of the major metabolite (5) was determined by HR-MS to be C48H64O21, which is identical to a known aureolic acid, chromocyclomycin. Subsequent isolation of 5 revealed that it was not chromocyclomycin, but instead a derivative structure where the glycosidic linkages were exclusively in the β configuration; in contrast, the two mycarose sugars in chromocyclomycin are α-linked. While literature evidence on the bioactivity of chromocyclomycin is limited, our data demonstrated here indicate the different configurations of the mycarose sugars have resulted in a different bioactivity profile between the two compounds.

Uncovering the mature product of the metathramycin gene cluster required us to integrate bioinformatic analysis, pathway engineering, and knowledge from previous biochemical and heterologous expression studies of aureolic acid BGCs. The structure of the major metabolite arising from heterologous expression was not consistent with the gene content of the BGC, suggesting this was not the final product. Bioactivity guided fractionation allowed us to isolate an extremely minor constituent 6 from a complex mixture of biosynthetic intermediates and shunt products, and analysis of fragmentation data from MS/MS experiments allowed a planar structure to be predicted. Finally, overexpression of key genes in the pathway supported our assertion that this was the product of a defined biosynthetic conversion from a metabolite for which we had attained detailed structural data, allowing us to propose likely stereochemistry for the final product of the pathway. Metathramycin was also demonstrated to have potent cytotoxic activity in HCT-116 cells, in keeping with the strong anticancer activities of previously reported aureolic acids. Such a discovery approach may
be generalisable to other families of natural product where biosynthetic gene clusters describe well characterised biosynthetic transformations, as has been previously applied for elucidation of stereochemistry. Without the prior in depth characterisation of aureolic acid biosynthesis by other research groups, such an approach would not have been possible, and it is unlikely that the final product of this pathway would have been discovered.

Future studies may now take place with the metathramycin gene cluster as a basis for combinatorial biosynthesis or derivatisation, as has been performed previously in the search for mithralogs. Other manipulations of biosynthetic pathway regulation may permit greater compound production of mithralogs. The approach used in this study may be applied to future discovery pipelines to find new mithralogs or other low abundance bioactive metabolites.

**Methods**

**Construction of soil metagenome cosmid library**

A soil sample of ~1 kg was collected from Half Moon Bay, Kaikoura New Zealand. A sample of 250 g was weighed into a 1 L centrifuge bottle, with 270 mL heated (70 °C) lysis buffer (100 mM Tris–HCl, 100 mM Na EDTA, 1.5 M NaCl, 1% (w/v) cetyl trimethyl ammonium bromide, pH 8.0) added. This was mixed to ensure the soil was thoroughly wetted, and returned to the water bath to reach 70 °C. A volume of 30 mL heated (70 °C) 20% SDS (w/v) was added to the centrifuge tube, inverted to mix, and returned to the water bath. The sample was incubated for 2 h, with gentle inversion every 30 min. The sample was then rapidly cooled in an ice/water bucket for 30 min, with inversion once in this time. The soil and chilled precipitate were then separated from the clarified lysate by centrifugation at 4500 rcf for 35 min at 4 °C. The supernatant was recovered, volume measured, and returned to room temperature by brief incubation in the warm water bath. Isopropanol was added to 0.7 × the supernatant volume, and mixed by gentle inversion to precipitate the eDNA. Following a 30 min room temperature incubation, the precipitated eDNA was collected by centrifugation at 4500 rcf for 35 min at 4 °C. The supernatant was discarded, and the eDNA pellet and centrifuge bottle was washed with 100 mL of ice cold 70% (v/v) ethanol. The sample was again centrifuged at 4500 rcf for 10 min at 4 °C. The supernatant was discarded, and the bottle briefly centrifuged again to collect the remaining ethanol, which was removed by pipette. The eDNA pellet was briefly air dried (no more than 15 min). A minimum volume of TE buffer was added to cover the eDNA pellet, and this was left to slowly resuspend overnight at room temperature.

The eDNA substrate for cosmid library construction was then size selected by agarose gel extraction in 0.8% agarose 1× TAE gel run at 80 V for 2 h alongside λ HindIII marker, then overnight in fresh TAE at 18 V. DNA above the 23 kb marker was excised from the gel and electroeluted, then concentrated using a 30000 kDa molecular weight cut-off column centrifugal concentrator. The DNA sample was quantified by nanodrop, then end-repaired using End-It DNA End-Repair Kit (Epigenet). The cosmid vector pWEB::tnc was prepared by digested with SmaI restriction enzyme (NEB), and dephosphorylation with Antarctic phosphatase (NEB) according to the manufacturer's instructions. The prepared cosmid vector (250 ng) and eDNA substrate (125 ng) were then ligated in a final reaction volume of 5 μL using Fast-link DNA ligase (Epigenet) according to the manufacturer's instructions. Phage packaging extract was prepared in house according to Winn and Norris (2010), and used to package the ligated DNA. The resulting diluted phage heads were added in a 1 : 10 ratio to an ice cold, day culture of E. coli EC100 (or EC100 ΔentD) grown to OD_{560} of 1.0 in LB 10 mM MgSO_{4}. The phage head/cell mixture was incubated at room temperature for 20 min, then aliquoted across 96 wells and incubated at 37 °C 200 rpm for 75 min. The recovered, transfected E. coli cells were then diluted in LB Amp Chl for cosmid selection, and titre samples of select wells plates plated on LB Amp Chl agar, for colony counting and library size estimations. The remaining cultures were incubated overnight at 37 °C 200 rpm to replicate the library clones in liquid culture. The following morning, samples were taken from each of the 96 aliquots to make glycerol stocks for long term storage, alongside minipreps of each library “well”.

**Sequencing of clones, bioinformatic interrogation**

Cosmids and BACs to be sequenced were isolated from E. coli cultures by miniprep, and Macrogen Inc. performed both Sanger and Illumina HiSeq PE100 bp sequencing. Sanger sequencing primers for cosmid end sequencing were standard T7pro and M13F-pUC primers.

Illumina HiSeq (100 bp PE) reads were pre-processed to remove pWEB::tnc vector sequence and contaminating E. coli gDNA sequences using bowtie2, followed by read quality trimming and sequencing adapter removal by Trimmomatic. The resulting processed Illumina PE reads were then assembled using SPAdes genome assembler.

**Transformation associated recombination**

The E. coli yeasts* Streptomyces shuttle vector, pTARA, was retrofit with a synthetic DNA block via Gibson assembly to produce the TAR capture vector. The synthetic DNA contains 712 bp homology capture arms to each of the cosmid insert sequences, separated by a PmeI restriction site. To prepare for TAR, the vector was linearised by PmeI digestion. Cosmid inserts were liberated from the pWEB::tnc vector by in vitro Cas9 digestion. The CRISPR guide RNAs target to the pWEB::tnc vector ~190 nt upstream of the cosmid insert site (GGTTATGTCACTTAG), and ~115 nt downstream of the cosmid insertion site (GTATAATGTCAACGCAGTC).

The three linear DNA units were transformed into S. cerevisiae BY4727 ΔAhl4 for recombination and reconstitution of the BGC with the BAC. The Kallifidas and Brady (2012) protocol was followed for the preparation of yeast spheroplasts, and for the transformation of the spheroplasts with pTARAs capture vector and overlapping clones.
Conjugation into *Streptomyces albus* expression host

Constructs to be conjugated into *Streptomyces* were first transformed by electroporation into *E. coli* S17-1. Overnight cultures of the S17-1 “donor” strain were sub inoculated into fresh LB with appropriate antibiotics, and cultured at 37 °C until OD<sub>600</sub> reached 0.5–1.0. S17-1 cultures were then chilled and washed in fresh ice-cold LB broth, and resuspended in a final volume of 1 mL LB. The “recipient” *S. albus* J1074 spore stock was diluted into 0.5 mL TSB or LB, and heat shocked at 50 °C for 10 min.

The donor and recipient stocks were then mixed in a 1:1 v/v ratio, and centrifuged at 4000 ref for 5 min. The cell pellet was resuspended in 50–100 μL of the residual LB, and spread onto ISP4 agar, without antibiotics. Plates were incubated for 12–16 h at 30 °C, and then flooded with 1 mL of the appropriate antibiotic stock mix (for pTARA 0.5 mg mL<sup>−1</sup> nalidixic acid, 1.25 mg mL<sup>−1</sup> apramycin sulfate, and for pIJ10257 0.5 mg mL<sup>−1</sup> nalidixic acid, 2.0 mg mL<sup>−1</sup> hygromycin). Plates were incubated at 30 °C for 24–72 h until colonies emerged.

Spectroscopic analysis

Optical rotations were measured using a Rudolph Autopol II polarimeter. A JEOL JNM-ECZ600R with a nitrogen cooled 5 mm SuperCOOL cryogenic probe was used to record the NMR spectra of all compounds (600 MHz for H nuclei and 150 MHz for C nuclei). The residual solvent peak was used as an internal reference for <sup>1</sup>H (δ<sub>H</sub> = 3.31, CD<sub>3</sub>OD; 7.26, CDCl<sub>3</sub> and <sup>13</sup>C (δ<sub>C</sub> = 49.0, CD<sub>3</sub>OD; 71.6, CDCl<sub>3</sub>) chemical shifts. Standard pulse sequences supplied by JEOL were used for NMR analyses. High-resolution (ESI) mass spectrometric data was obtained with an Agilent 6530 Accurate Mass Q-TOF LCMS equipped with a 1260 Infinity binary pump. IR (film) spectra were recorded with an Agilent 6530 Accurate Mass Q-TOF LCMS equipped with a 1260 Infinity binary pump. IR (film) spectra were recorded using a Bruker Platinum Alpha FTIR spectrometer, while UV/vis spectra were extracted from HPLC chromatograms.

Reversed-phase column chromatography was achieved using Supelco Diaion HP20 (PSDVB) chromatographic resin. Size exclusion chromatography was achieved using Sephadex LH20 resin. HPLC purifications were carried out using a Rainin Dynamax SD-200 solvent delivery system with 25 mL pump heads with a Varian Prostar 335 diode array detector. Octadecyl-derivatised silica (C<sub>18</sub>, 5 μm, 100 Å) HPLC columns (Phenomenex) were either analytical (4.6 mm × 250 mm, 1 mL min<sup>−1</sup>) or semi-preparative (10 mm × 250 mm, 4 mL min<sup>−1</sup>), while a Phenomenex Luna octyl-derivatised silica gel (C<sub>8</sub>) column (analytical; 4.6 mm × 250 mm, 1 mL min<sup>−1</sup>) was also used. All solvents used were of HPLC grade and H<sub>2</sub>O was glass distilled. Solvent mixtures are reported as percent volume/volume.

Production and isolation of premetathramycin and metathramycin

Cultures of *S. albus* J1074::MMY were grown in 7 L total R5a medium, with 1% inoculum from a mature seed culture grown in TSB medium. R5a cultures were prepared with the addition of 20 g L<sup>−1</sup> of HP20 resin to collect metabolites as they were produced during the 7-day culture time. The HP20 was washed with 1 L of H<sub>2</sub>O, and 1500 mL of 20% methanol/H<sub>2</sub>O, before elution in 1500 mL 80% methanol/H<sub>2</sub>O. The 1500 mL elution was dried under reduced pressure, resuspended in 80 mL of methanol, and dissolved in acetone removed via centrifugation. The clarified supernatant was dried, with a mass of 1091 mg. Half of this sample was then resuspended in methanol, syringe filtered, and partitioned on an LH20 column run in methanol, with 5 mL fractions collected and combined based on MS analysis for the presence of the protonated molecular ion of 5. Four early fractions were combined, and were dried under reduced pressure to a dry mass of 262 mg. This sample was purified by semi-preparative C<sub>18</sub> HPLC with a gradient of 10–100% methanol/H<sub>2</sub>O (0.2% formic acid) over 60 min. Two fractions were collected, one containing the ion for compound 6 (8.0 mg) with a retention time of 51.5 min and another with compound 5 (19.7 mg) that had a retention time of 52.3 min. The first sample was purified by analytical C<sub>18</sub> HPLC using an isocratic method of 70% methanol/H<sub>2</sub>O (0.2% formic acid), to give compound 6 (7.2 μg). A 10 mg portion of the second sample was also purified by analytical C<sub>18</sub> HPLC (75% methanol/H<sub>2</sub>O (0.2% formic acid)), to give compound 5 (7.9 mg). This represents a recovery of 4.4 mg L<sup>−1</sup> of culture (when accounting for the unprocessed sample at each stage of purification).

Pemetathramycin (5): yellow solid; [α]<sub>D</sub> <sup>20</sup> = 183 (c 0.3, EtOH); UV (MeOH) <sub>λ</sub> <sub>max</sub> 228, 280, 330, 415 nm; IR (film) <sub>v</sub> <sub>max</sub> 3382, 2928, 1628, 1598, 1530, 1370, 1065 cm<sup>−1</sup>; <sup>1</sup>H NMR (1:1 CD<sub>3</sub>OD:CDCl<sub>3</sub>) Tables S1 and S2 ESI;<sup>†</sup> <sup>13</sup>C NMR (1:1 CD<sub>3</sub>OD:CDCl<sub>3</sub>) Tables S1 and S2 ESI;<sup>†</sup> HRESIMS m/z 975.3848 [M – H]<sup>−</sup> (c) calculated for C<sub>49</sub>H<sub>64</sub>O<sub>21</sub>, 975.3867, Δ = −1.9 ppm.

Antimicrobial testing

Minimum inhibitory concentration (MIC) studies were performed using an established protocol<sup>49</sup> to test isolated compounds against test strains of *E. coli* ΔtolC (7NT)<sup>50</sup> and *B. subtilis* 168. All tests were carried out in MH medium at 30 °C, with 16 h growth time in biological triplicates (n = 3 independent experiments). Positive controls of tetracycline and mithramycin A were included in the analysis. All compounds were assayed from 128 μg mL<sup>−1</sup> in two-fold serial dilution steps across 20 dilution stages, except for mithramycin, where the starting concentration was 32 μg mL<sup>−1</sup>. Sterility (media only) and growth (cells only) controls were included in each plate assayed.

The same protocol was followed for fungal MICS against *S. cerevisiae* lacking the pleiotropic drug resistance gene PDR4, with the growth medium YPD, and the positive control antibiotic nystatin. The end-point measurement was changed from an eyesight measure to optical density, where a growth inhibited well was one where the OD measurement was < 90% of the growth control well (as per EUCAST guidelines for fungal susceptibility testing).

Tumour cell cytotoxicity assay

A standard 48 h MTS cell proliferation assay was used to evaluate cytotoxic activity against the human colon carcinoma cell line HCT-116 (n = 4 independent experiments). Cells were treated with compound at various concentrations, and a dose–response was generated relative to a control of untreated...
HCT-116 cells. A positive control of mithramycin A was used in all cases.

Conflicts of interest

There are no conflicts to declare.

References