

# Biosynthesis of thiomarinol A and related metabolites of *Pseudoalteromonas* sp. SANK 73390†

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The biosynthesis of the mixed PKS-FAS-NRPS hybrid antibiotic thiomarinol A was investigated using feeding studies to both wild type and mutant strains of the marine bacterium *Pseudoalteromonas*. Particularly interesting features of the pathway include assembly of the 8-hydroxyoctanoic acid side-chain *via* chain extension of a C<sub>4</sub>-precursor (4-hydroxybutyrate), and construction of the pyrrothine unit from cysteine *via* (HoA-D, F-H) prior to intact incorporation into thiomarinol (catalysed by TmU). A series of thiomarinol-related and other minor metabolites have been isolated from wild-type and mutant strains. The results of these investigations are rationalised in terms of the overall biosynthetic pathway.

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## Introduction

Thiomarinol A **1** (Fig. 1), is the major metabolite produced by the marine bacterium *Pseudoalteromonas* sp. SANK 73390, along with minor analogues B to G (**2** to **7**).<sup>1–3</sup> They display antibiotic activity against both Gram-negative and Gram-positive bacteria, and have particularly potent activity against methicillin-resistant *Staphylococcus aureus* (MRSA). They are unusual in being hybrid antibiotics, effectively two separate antibiotic classes in one molecule, and biosynthetically can be regarded as comprising three main elements: a highly functionalised polyketide-derived acid, which is esterified by a hydroxy fatty acid, which itself forms an amide with the bicyclic amino acid-derived pyrrothine. The main carbon skeleton of the thiomarinols is very closely related to that of mupirocin, an important antibiotic used clinically against MRSA.<sup>4</sup> Mupirocin consists of a mixture of pseudomonic acids (PA A–C, **8** to **10**),<sup>5–7</sup> with PA-A comprising *ca.* 95% of the mixture. The major differences in thiomarinol A are the replacement of the 9-hydroxynonanoic acid moiety in the PAs with 8-hydroxyoctanoate in the

thiomarinols, the presence of an extra 4-hydroxyl group, the lack of the 10,11-epoxide and the pyrrothine. Many pyrrothine natural products themselves display antibiotic activity, *e.g.* holomycin **11**, *N*-propionylholothin **12**, thiolutin **13**, and aureothricin **14**.<sup>8–10</sup> Stable isotope labelling studies were used to confirm the origins of all of the atoms in pseudomonic acid A and were consistent with a polyketide/fatty acid origin.<sup>11,12</sup> This was subsequently confirmed by isolation of the biosynthetic gene cluster<sup>13</sup> which revealed that mupirocin was one of the first members of what is now known to comprise an increasingly large number of biologically active compounds. These are most often from diverse and unusual bacterial sources and are produced *via trans*-AT modular PKSs that have evolved independently from other modular systems.<sup>14</sup> The main identifying feature of these systems is the lack of an integral acyl transferase (AT) domain in each chain extension module, this function being supplied “*in trans*” to each module by separately encoded mono- (or di-) functional ATs. In addition they show many other unusual features including extensive involvement of other *trans* acting enzymes during polyketide assembly. In particular  $\beta$ -branches are introduced onto  $\beta$ -keto-thiol ester intermediates by a cassette of enzymes that include an HMG CoA synthase (HMGS) analogue using chemistry similar to that involved in mevalonate formation in terpene biosynthesis.<sup>15</sup> The biosynthesis of the pyrrothine class of natural products has not been studied in detail, although cysteine has been implicated as a precursor *via* radioisotope labelling studies.<sup>16</sup>

We have recently identified the thiomarinol (*tml*) biosynthetic gene cluster *via* full genome sequencing of SANK 73390. It is contained on a 97 kb plasmid consisting almost entirely of the thiomarinol biosynthetic genes.<sup>17</sup> These consist of *trans*-AT

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pathways. A regular acetate incorporation pattern is observed in the C1–C14 fragment of thiomarinol. A high level of  $^{13}\text{C}$  incorporation (5–6%) is observed at all seven expected positions in the polyketide-derived moiety with high levels of  $^{18}\text{O}$  retention (*ca.* 80%) at carbons 1, 5, 7 and 13 (Table 1). The 4- and 6-hydroxyl groups are not labelled from  $^{18}\text{O}$ -acetate and are at positions consistent with their introduction being *via* tailoring enzymes using molecular oxygen. The 16-methylene (55% enriched from [methyl- $^{13}\text{C}$ ]methionine) and 17-methyl carbons (40% enriched) are methionine derived consistent with the presence of C-MeT domains in the first and third extension modules, and the 15-methyl is derived from the methyl of a cleaved acetate unit consistent with it being derived *via* the HMGS  $\beta$ -branch pathway as in pseudomonic acid A.

### Biosynthetic origin of the side-chain

Interestingly the labelling pattern in the octanoate side chain did not follow a standard polyketide labelling pattern. Whilst expected head-to-tail incorporation of intact acetate units from

**Table 1** Incorporations of isotopically labelled acetates into thiomarinol A

Carbon	$\delta_{\text{C}}^{\text{c}}$	[1,2- $^{13}\text{C}_2$ ]-acetate <sup>a</sup> % ( $J_{\text{C-C}}$ Hz)	[2- $^{13}\text{C}$ ]-acetate <sup>a</sup> %	[1- $^{13}\text{C}$ , $^{18}\text{O}_2$ ]-acetate <sup>a</sup> % $\Delta\delta_{\text{C}}^{\text{b}}$ (ppm)
1	166.1	2 (76)		5 (0.03)
2	114.3	2 (76)	7	5
3	160.8	2 (43)		6 (0.02)
4	72.4	2 (43)	7	
5	76.2	2 (43)		6 (0.02)
6	63.8	2 (43)	7	
7	69.6	2 (35)		6 (0.02)
8	42.2	2 (35)	7	
9	31.8	2 (44)		6
10	127.7	2 (44)	7	
11	134.1	2 (44)		6
12	43.1	2 (44)	7	
13	69.2	2 (39)		6 (0.02)
14	20.0	2 (39)	7	
15	15.7	2	7	
16	64.2			
17	15.7			
1'	171.8	2 (50)		6 (0.03)
2'	34.6	2 (50)	7	
3'	24.9	2 (35)		6
4'	28.4	2 (35)	7	
5'	28.3	<b>1</b> (34)		<b>3</b>
6'	25.3	<b>1</b> (34)	<b>4</b>	
7'	28.2	<b>0.3</b> (38)	<b>2</b>	
8'	63.0	<b>0.3</b> (38)	<b>2</b>	<b>1</b>
1''	167.9			
2''	115.3			
3''	133.9			
4''	133.6			
5''	110.3			

<sup>a</sup> Percentage enrichment (>1 to nearest integer); significant differences from average values indicated in bold. <sup>b</sup> Average ratio of enriched ( $^{13}\text{C}$ ) to enriched + isotopically shifted ( $^{13}\text{C}^{18}\text{O}$ ) signals is 1 : 4 except C-1' which is *ca.* 1 : 1, all values to nearest integer. <sup>c</sup> Spectra determined in  $\text{d}_6$ -DMSO, and  $\text{d}_4$ -MeOH in which the 15- and 17-methyls appear respectively at 14.9 and 15.2 ppm.

[1,2- $^{13}\text{C}_2$ ]acetate was observed for C1' to C4' (2% incorporation), a lower level of incorporation was observed in C5' and C6' (1%) and much lower into C7' and C8' (0.3%). [2- $^{13}\text{C}$ ]-Acetate also gave a similar pattern with much lower incorporation of  $^{13}\text{C}$  into both C7' and C8' (~2% compared to 7% in other acetate-derived positions), and an intermediate level of enrichment (4%) at C6' (Table 1). [1- $^{13}\text{C}$ ,  $^{18}\text{O}_2$ ]-Acetate gave incorporation levels of  $^{13}\text{C}$  (~6%) at carbons 1' and 3' as was observed elsewhere, but incorporation into C5' was half this level and at C7' was negligible. A low incorporation, *ca.* 1%, into C8' was observed. These results suggest that a  $\text{C}_4$  precursor is likely to be the starter unit for two rounds of FAS catalysed extension to give the octanoate side-chain. The observed scrambling of acetate incorporation at C7' and C8' suggested the possible involvement of a citric acid (Krebs) cycle intermediate.

Feeding [2,3- $^{13}\text{C}_2$ ]succinate to cultures of wild-type *Pseudoalteromonas* SANK 73390 revealed an intact incorporation confirmed by observation of a  $^{13}\text{C}$ - $^{13}\text{C}$  coupling of 34 Hz between C6' and C7' (*ca.* 0.5% enrichment, Scheme 2, see insert expansion in Fig. S10†) in accord with the proposed  $\text{C}_4$  precursor. A lower level of incorporation (0.15%) was also observed elsewhere within the molecule resulting from degradation of [2,3- $^{13}\text{C}_2$ ]succinate to [1,2- $^{13}\text{C}_2$ ]acetate.

It is known that 4-hydroxybutyrate can be formed *via* reduction of succinyl CoA.<sup>24</sup> The overall results of acetate incorporation are also consistent with this: acetate is incorporated intact into only carbons 3 and 4 of succinyl CoA (Scheme S2, ESI†) which explains the significant but lower incorporation into C5'/C6' of thiomarinol, presumably due to dilution by a pool of endogenously derived Krebs Cycle intermediate. Further cycling of labelled succinate would give single labels at carbons 1 and 2 of succinyl CoA which explains the lower levels of labelling from [2- $^{13}\text{C}$ ]acetate into C7' and C8'. To explain the low level of intact acetate labelling of C7'/C8', the conversion of succinyl CoA to symmetrical succinate must be partially reversible so that *ca.* 30% of the succinate is incorporated *via* this route. The low level of labelling at C8' from [1- $^{13}\text{C}$ ]acetate is completely consistent with this hypothesis.

To further investigate the proposed  $\text{C}_4$  side-chain precursor, [2,3- $^{13}\text{C}_2$ ]-4-hydroxybutyrate was prepared from [2,3- $^{13}\text{C}_2$ ]succinate *via* cyclisation to succinic anhydride, reduction to the lactone with  $\text{LiAlH}_4$  followed by hydrolytic ring opening with aqueous NaOH (Scheme S1†). Feeding [2,3- $^{13}\text{C}_2$ ]-4-hydroxybutyrate to cultures of the wild-type *Pseudoalteromonas* SANK 73390 showed a site-specific labelling (0.2%) of thiomarinol A at C6', C7' confirming intact incorporation (Scheme 2, see insert expansion in Fig. S12†).



**Scheme 2** Incorporation of [ $^{13}\text{C}$ ]-labelled succinate, 4-hydroxybutyrate and cystine into thiomarinol A.



Thus the biosynthetic origin of C5' to C8' is somewhat similar to the situation presumed for 9-hydroxynonanoic in mupirocin biosynthesis, where labelling studies are consistent with a 3-hydroxypropionate starter unit, which then undergoes three reductive chain elongations. 3-Hydroxypropionate can be formed in bacteria by reduction of the thioester of malonyl CoA<sup>25</sup> and both the *mup* and *tml* clusters show the presence of genes encoding a malonyl/succinyl CoA dehydrogenase clustered with an acyl CoA synthetase and an ACP.

Interestingly, the biosynthetic gene cluster for diffidin,<sup>26</sup> another *trans*-AT metabolite of *Bacillus amyloliquefaciens*, also contains an oxo-acyl CoA reductase encoded by *difE* which as in mupirocin/thiomarinol is grouped with an acyl CoA synthetase and an ACP (*difD* and *difC*). In addition there is an adjacent kinase (*difB*) which would be suitable for catalysing dehydration of 3-hydroxypropionate to produce the acryl thiol ester required to prime diffidin chain elongation.

The retention of <sup>18</sup>O label at C1' of thiomarinol A is slightly less than half that at the other acetate-oxygen bearing positions. This is consistent with C1' existing as the free carboxylate, resulting in randomisation of the label between both carboxylate oxygens, before ligation *via* an amide bond to the pyrrothine. Analysis of the structure of the pyrrothine moiety of thiomarinol suggest that it is formed *via* two molecules of cysteine. To gain evidence for this proposal, [2-<sup>13</sup>C]cystine was prepared from ethyl [2-<sup>13</sup>C]pyruvate, (Scheme S2†) and fed to WT *Pseudoalteromonas*. A high level of incorporation of <sup>13</sup>C into the C2'' and C4'' positions (16 and 20% respectively) confirmed that the pyrrothine is indeed cysteine, and therefore NRPS, derived (Scheme 2).

The biosynthetic gene cluster for holomycin biosynthesis in *Streptomyces clavuligerus* has been isolated and a pathway from cysteine proposed. This pathway involves initial dipeptide formation followed by a series of oxidative cyclisations, the final step being acetylation of the free pyrrothine.<sup>18</sup> To prove that the pyrrothine moiety in thiomarinol is assembled and then incorporated as an intact unit rather than being assembled on marinolic acid, pyrrothine was synthesised as previously described<sup>27</sup> and fed to the  $\Delta$ NRPS mutant strain of SANK 73390. The mutant was prepared by a deletion in the *holA* gene encoding the amino acid activation domain which, from sequence analysis of the active site, is predicted to selectively activate cysteine. As indicated above, the  $\Delta$ NRPS mutant normally produces mainly marinolic acid. When cultures were supplemented with pyrrothine, the normal WT phenotype was restored indicating that *Pseudoalteromonas* takes the intact pyrrothine as substrate (Scheme 3). Interestingly, the concentration of the feeds was critical, with maximum restoration of thiomarinol A production being observed at a pyrrothine concentration of 20 mg ml<sup>-1</sup>. At concentrations of >80 mg ml<sup>-1</sup>, no thiomarinol A was observed and production of marinolic acid was also completely inhibited (Fig. S3, ESI†).

### Novel acyl pyrrothines *via* mutasynthesis

Previous studies have shown that when TmlU, a putative amide ligase is knocked out, the mutant strain produces no



Scheme 3 Feeding pyrrothine to the NRPS mutant gives thiomarinol A.

thiomarinol but does produce marinolic acid and the full set of xenorhabdins.<sup>20</sup> The HolE analogue, ORF3483, from the holomycin cluster in *S. clavuligerus* has been expressed and purified and it has been shown<sup>18</sup> to act as an acyl transferase capable of transferring acetyl and longer acyl groups to pyrrothine. Thus it appears that the thiomarinol cluster has two ligases, TmlU which forms the amide in thiomarinol, and HolE which links pyrrothine to a wide range of acyl CoAs available in *Pseudoalteromonas* to produce the xenorhabdins.

To test if HolE can also transfer a range of non-endogenous substrates, various short, medium and long chain fatty acids and others with  $\omega$ -functionalized side chains, *e.g.* hydroxyl, amino, carboxyl and phenyl were fed to the  $\Delta$ PKS mutant (Scheme 4). Some of these were incorporated with varying efficiencies, presumably after conversion to their respective CoA esters, sometimes after catabolic chain shortening to give **19** and the novel acyl-pyrrothines **26–30**.

### The biosynthesis of thiomarinol A and related metabolites

The isolation of the minor metabolites **15–18** from both WT and mutant strains and the incorporation of isotopically labelled substrates can be rationalised as shown in Scheme 5. Marinolic acid **15** is produced by the  $\Delta$ NRPS and  $\Delta$ TmlU mutants, and is present in trace amounts in the WT which suggests that it is both a genuine biosynthetic product and intermediate in its own right and is not produced as an artefact, *e.g.* as a degradation product of thiomarinol A. The intermediacy of marinolic acid **15** was confirmed by feeding it to the  $\Delta$ PKS mutant whereupon, thiomarinol production was restored (Fig. S19†). The amide **16** on the other hand was only seen in the WT cultures which suggests that it is formed by degradation of thiomarinol A, and is not formed by, *e.g.* amidation of marinolic acid. This was further evidenced by feeding thiomarinol A to the  $\Delta$ PKS mutant when the presence of both marinolic acid and marinolic amide in the crude extract could be observed by LCMS. This suggests that the amide is indeed a degradation product of thiomarinol.

The isolation of truncated metabolites marinolic acids A<sub>4</sub> and A<sub>6</sub>, is consistent with the isotope labelling studies which suggest that a C<sub>4</sub> intermediate is added to the product of the PKS, *e.g.* by *trans*-esterification. This would then be elongated to



Scheme 4 Novel acyl-pyrrothines generated using the  $\Delta$ NRPS mutant.





Scheme 5 Overall biosynthesis of thiomarinol and related metabolites in *Pseudoalteromonas*.

give an enzyme bound hexanoate analogue which would be further elongated to give the completed octanoate side chain. In the absence of the pyrroline in the  $\Delta$ NRPS mutant they would be released either by enzymatic or spontaneous hydrolysis as marinolic acids. We have previously shown that release of assembly intermediates occurs when flux of intermediates along the normal mupirocin pathway is impaired by a range of targeted mutations of *trans*-acting and other tailoring genes.<sup>28</sup> Similar mechanisms would allow leakage of marinolic acids A<sub>4</sub> and A<sub>6</sub> into the medium. It is significant that this appears to be more pronounced in the mutant than the WT. Truncated forms of mupirocin and related metabolites with C<sub>7</sub> and C<sub>5</sub> hydroxy acids have also been observed in both mutant strains and when the quorum sensing regulation mechanism has been manipulated to increase mupirocin production.<sup>29</sup>

## Conclusions

Using a combination of genetic and isotopic labelling studies, evidence has been presented for a mixed PKS-NRPS biosynthetic pathway to the antibiotic thiomarinol A in *Pseudoalteromonas* sp. SANK 73390. Particularly interesting features of the pathway include assembly of the 8-hydroxyoctanoic acid side-chain, which is generated *via* chain extension of a C<sub>4</sub> precursor (4-hydroxybutyrate), and both marinolic acids A<sub>4</sub> 18 and A<sub>6</sub> 17 have been isolated as minor metabolites from both wild-type and the  $\Delta$ NRPS mutant. There was a good incorporation of [<sup>13</sup>C]cysteine into thiomarinol A and feeding studies with synthetic pyrroline to the

$\Delta$ NRPS mutant are in accord with heterocycle assembly (HoloA-D, F-H) prior to intact incorporation into thiomarinol A (TmlU). A further minor metabolite, marinolic amide 16, present in wild-type but not the  $\Delta$ NRPS mutant, is a degradation product of thiomarinol. Isotopic labelling studies indicate that the biosynthesis of the C1–C17 fragment is analogous to that of pseudomonic acid, a *trans*-AT modular PKS pathway with many fascinating and, as yet, unexplained features and these studies lay the foundation for a further understanding of the pathways to these compounds.

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## Notes and references

- 1 H. Shiozawa, I. T. Kagasak, T. Kinoshita, H. Haruyama, H. Domon, Y. Utsui, K. Kodama and S. Takahashi, *J. Antibiot.*, 1993, **46**, 1834–1842.
- 2 H. Shiozawa, T. Kagasaki, A. Torikata, N. Tanaka, K. Fujimoto, T. Hata, Y. Furukawa and S. Takahashi, *J. Antibiot.*, 1995, **48**, 907–909.



- 3 H. Shiozawa, A. Shimada and S. Takahashi, *J. Antibiot.*, 1997, **50**, 449–452.
- 4 C. M. Thomas, J. Hotherhall, C. L. Willis and T. J. Simpson, *Nat. Rev. Microbiol.*, 2010, **8**, 281–289.
- 5 A. T. Fuller, G. Mellows, M. Woolford, G. T. Banks, K. D. Barrow and E. B. Chain, *Nature*, 1971, **234**, 416–417.
- 6 E. B. Chain and G. Mellows, *J. Chem. Soc., Perkin Trans. 1*, 1977, 294–309.
- 7 E. B. Chain and G. Mellows, *J. Chem. Soc., Perkin Trans. 1*, 1977, 318–322.
- 8 L. Ettlinger, E. Gäumann, R. Hütter, W. Keller-Schierlein, F. Kradolfer, L. Neipp, V. Prelog and H. Zähner, *Helv. Chim. Acta*, 1959, **42**, 563–569.
- 9 W. D. Celmer and I. A. Solomons, *J. Am. Chem. Soc.*, 1955, **77**, 2861–2865.
- 10 K. Okamura, K. Soga, Y. Shimauchi, T. Ihikura and J. Lein, *J. Antibiot.*, 1977, **30**, 334–336.
- 11 T. C. Feline, R. B. Jones, G. Mellows and L. Phillips, *J. Chem. Soc., Perkin Trans. 1*, 1977, 309–318.
- 12 F. M. Martin and T. J. Simpson, *J. Chem. Soc., Perkin Trans. 1*, 1989, 207–209.
- 13 A. K. El-Sayed, J. Hotherhall, S. M. Cooper, E. Stephens, T. J. Simpson and C. M. Thomas, *Chem. Biol.*, 2003, **10**, 419–430.
- 14 J. Piel, *Nat. Prod. Rep.*, 2010, **27**, 996–1047.
- 15 C. T. Calderone, *Nat. Prod. Rep.*, 2008, **25**, 845–853.
- 16 M. Okanishi and H. Umezawa, *Genetics of the Actinomycetales*, Gustav Fischer, Stuttgart, New York, 1978.
- 17 D. Fukuda, A. S. Haines, Z. Song, A. Murphy, J. Hotherhall, E. R. Stephens, R. Gurney, R. J. Cox, J. Crosby, C. L. Willis, T. J. Simpson and C. M. Thomas, *PLoS One*, 2011, **6**, e18031.
- 18 B. Li and C. T. Walsh, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 12629–12633.
- 19 S. Huang, Y. Zhao, Z. Qin, X. Wang, M. Onega, L. Chen, J. He, Y. Yu and H. Deng, *Process Biochem.*, 2011, **46**, 811–816.
- 20 A. C. Murphy, D. Fukuda, Z. Song, J. Hotherhall, R. J. Cox, C. L. Willis, C. M. Thomas and T. J. Simpson, *Angew. Chem., Int. Ed.*, 2011, **123**, 3271–3274.
- 21 B. V. McInerney, R. P. Gregson, M. J. Lacey, R. J. Akhurst, G. R. Lyons, S. H. Rhodes, D. R. J. Smith, L. M. Engelhardt and A. H. White, *J. Nat. Prod.*, 1991, **54**, 774–784.
- 22 J. Li, G. Chen, J. M. Webster and E. Czyzewska, *J. Nat. Prod.*, 1995, **58**, 1081–1086.
- 23 S. Paik, Y. H. Park, S. I. Suh, H. S. Kim, I. S. Lee, M. K. Park, C. S. Lee and S. H. Park, *Bull. Korean Chem. Soc.*, 2001, **22**, 372–374.
- 24 I. A. Berg, D. Kockelkorn, W. Buckel and G. Fuchs, *Science*, 2007, **318**, 1782–1786.
- 25 M. Hügler, C. Menendez, H. Schägger and G. Fuchs, *J. Bacteriol.*, 2002, **184**, 2404–2410.
- 26 X.-H. Chen, J. Vater, J. Piel, P. Franke, R. Scholz, K. Schneider, A. Koumoutsis, G. Hitzeroth, N. Grammel, A. W. Strittmatter, G. Gottschalk, R. D. Suessmuth and R. Borriss, *J. Bacteriol.*, 2006, **188**, 4024–4036.
- 27 T. Hjelmgaard, M. Givskovb and J. Nielsen, *Org. Biomol. Chem.*, 2007, **5**, 344–348.
- 28 J.-e. Wu, J. Hotherhall, C. Mazetti, Y. O'Connell, J. A. Shields, A. S. Rahman, R. J. Cox, J. Crosby, T. J. Simpson, C. M. Thomas and C. L. Willis, *ChemBioChem*, 2008, **9**, 1500–1508.
- 29 J. Hotherhall, A. C. Murphy, Z. Iqbal, G. Campbell, E. R. Stephens, J.-e. Wu, H. Cooper, S. Atkinson, P. Williams, J. Crosby, C. L. Willis, R. J. Cox, T. J. Simpson and C. M. Thomas, *Appl. Microbiol. Biotechnol.*, 2011, **90**, 1017–1026.

