



Cite this: *Nat. Prod. Rep.*, 2025, 42, 1622

## Recent advances in discovery and biosynthesis of ribosomally synthesized and post-translationally modified peptides (RiPP)-derived lipopeptides

Shumpei Asamizu \*

Covering: This review summarizes recent advances in the discovery, biosynthesis, and bioactivity of RiPP-derived lipopeptides, covering studies published up to 2024.

Ribosomally synthesized and post-translationally modified peptides (RiPPs) are a diverse superfamily of natural products unified by a common biosynthetic logic: The peptide backbone is genetically encoded, and the translated precursor peptide undergoes a series of post-translational modifications catalyzed by maturase enzymes to produce the final bioactive compound. Despite their structural complexity, RiPPs are encoded by relatively small biosynthesis gene clusters. RiPP maturase enzymes are diverse and often promiscuous, offering significant biotechnological potential. However, their lack of conserved features makes genome-based discovery of novel RiPPs challenging. Recent advances in biosynthetic understanding and genome mining techniques have led to the identification of numerous uncharacterized RiPP biosynthetic gene clusters, often flanked by genes encoding non-RiPP moieties, in microbial genomes. Leveraging this information, a new class of natural products, hybrids of RiPPs and non-RiPP elements, has recently been discovered. Among them, RiPPs bearing fatty acyl groups, referred to as RiPP-derived lipopeptides, represent a newly emerging class of lipopeptide natural products with significant antimicrobial activity.

Received 2nd June 2025

DOI: 10.1039/d5np00042d

rsc.li/npr

1. Introduction
2. Polar functionalized fatty acylated RiPPs
  - 2.1. Discovery of RiPP-derived lipopeptides
  - 2.2. Lipolanthines: microvionin and nocavionin
  - 2.3. Polyketide/RiPP-hybrid lipopeptides: goadvionins
  - 2.4. Lipolanthine: albopeptin B
  - 2.5. Lipolanthines: solabiomycins
  - 2.6. Lipolanthines: deoxysolabiomycins
3. Medium-chain fatty acylated RiPPs
  - 3.1. Selidamides: kamptornamide and phaeornamide
4. Short-to-medium-chain acylation
  - 4.1. AviCys-containing thioamide: prethioviridamide
  - 4.2. AviMeCys-containing lipopeptides: lipoavitides
  - 4.3. Polytheonamide
  - 4.4. N-Acetylated RiPPs
5. Diversification of fatty acyl moieties
6. Bioactivity – potential molecular targets of RiPP-type lipopeptides
7. Concluding remarks
8. Data availability

9. Conflicts of interest
10. Acknowledgments
11. References

### 1. Introduction

Ribosomally synthesized and post-translationally modified peptides (RiPPs) are a diverse superfamily of natural products unified by a common biosynthetic logic.<sup>1–3</sup> Their biosynthesis typically begins with a ribosomally synthesized precursor peptide, which often comprises a leader and a core region, although alternative architectures also exist. The core peptide undergoes various post-translational modifications catalyzed by maturase enzymes and is subsequently proteolytically cleaved to yield the final bioactive mature product.

Advances in high-throughput gene sequencing technologies have ushered in a new era of natural product discovery, with RiPPs<sup>3</sup> emerging as one of the fastest-growing fields, supported by the development of specialized bioinformatic tools.<sup>4,5</sup> Despite the structural complexity and large molecular size of RiPPs, their biosynthesis gene clusters (BGCs) are relatively small. The maturase enzymes involved in RiPP biosynthesis are remarkably diverse and often promiscuous in substrate

Engineering Biology Research Center, Kobe University, 1-1 Rokkodai, Nada, Kobe 657-8501, Japan. E-mail: shumpei.asamizu@port.kobe-u.ac.jp



recognition, making them highly attractive for biotechnological applications. However, the discovery of novel RiPPs is hindered by their limited structural predictability in nature and the frequent absence of characteristic features in their maturase enzymes, which restricts their straightforward identification as RiPP-associated enzymes during genome mining.

Growing interest in RiPP-based therapeutics has been fueled by the discovery of compounds exhibiting diverse bioactivities, including antibacterial, antifungal, antiviral, antinociceptive, and antitumor effects.<sup>6</sup> These wide-ranging activities underscore how post-translational modifications drive the diversity of RiPPs, enabling them to engage a broad array of biological targets across different domains of life.

RiPPs exhibit unique modes of action.<sup>6</sup> For example, nisin—a class I lanthipeptide produced by *Lactococcus lactis*—is the most extensively studied RiPP and shows potent antimicrobial activity against many non-proteobacterial species.<sup>6</sup> For over 50 years, nisin has been widely used in the food industry to combat foodborne pathogens. Its mechanism involves binding to the cell wall precursor lipid II and forming stable pores in the bacterial membrane, composed of both nisin and lipid II, ultimately leading to bacterial cell death.<sup>7,8</sup>

Thiostrepton, a thiopeptide antibiotic originally isolated in the 1950s from *Streptomyces azureus*, is used in veterinary medicine and features a characteristic 26-membered macrocyclic core.<sup>6</sup> It exhibits potent growth-inhibitory activity against Firmicutes, including hemolytic streptococci. Thiostrepton exerts its antimicrobial effect by binding to the 50S ribosomal subunit, thereby inhibiting protein synthesis through two mechanisms: it blocks mRNA-tRNA translocation mediated by the GTPase elongation factor G (EF-G), and it interferes with EF-Tu-catalyzed delivery of aminoacyl-tRNA.<sup>9–11</sup>

A more recent finding is darobactin, isolated from *Photorhabdus kharii* HGB1456, which selectively kills Gram-negative pathogens by targeting BamA, the central component of the essential BAM complex that is responsible for folding and inserting outer membrane proteins,<sup>12</sup> mimicking the recognition signal of native substrates and displacing a lipid molecule.<sup>13</sup>

Alongside the discovery of RiPP natural products, novel modification patterns have also been revealed,<sup>1,2,14</sup> including fatty acylation of the RiPP backbone. Lipopeptide natural products (acylated peptides) include clinically valuable antibiotics, such as the antibacterial agents colistin/polymyxin B<sup>15</sup> and daptomycin (DAP),<sup>16</sup> as well as the antifungal agent echinocandin,<sup>17</sup> all of which are produced by nonribosomal peptide synthetase (NRPS) pathways.<sup>18,19</sup> Advances in genome mining have revealed numerous uncharacterized RiPP BGCs, often flanked by genes encoding non-RiPP elements, in microbial genomes. This has led to the discovery of hybrid natural products, including a newly emerging class of RiPP-derived lipopeptides bearing fatty acyl groups, which exhibit notable antimicrobial activity. In addition to the N-acylated RiPPs modified by short-chain acyl groups, this review summarizes recent advances in the discovery, biosynthesis, and bioactivity of RiPP-derived lipopeptides bearing medium- and long-chain acyl modifications, covering reports up to 2024 in this rapidly

developing field. Although RiPPs with short-chain acyl modifications are not typically classified as lipopeptides, the term “lipopeptide” is used in a broader sense in this review to encompass such modifications for the sake of comprehensive coverage.

## 2. Polar functionalized fatty acylated RiPPs

### 2.1. Discovery of RiPP-derived lipopeptides

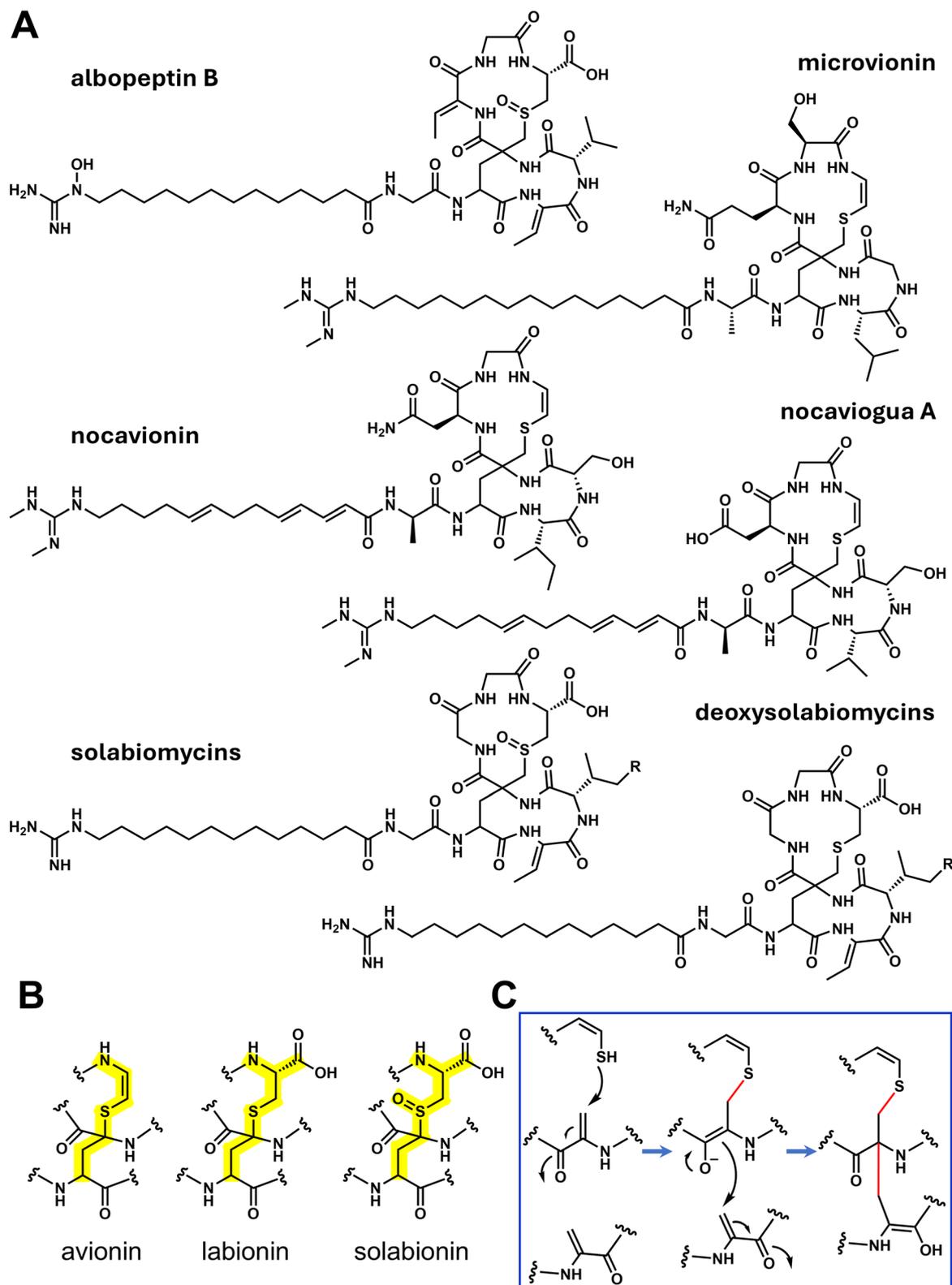
RiPP-derived lipopeptides with basic functional groups at the fatty acid terminus are designated “polar functionalized fatty acylated RiPPs” (PFARs). The first report of the isolation of RiPP-derived lipopeptides was the discovery of albopeptins A and B from *Streptomyces albofaciens* JC-82-120 by Isono *et al.* in 1986 (ref. 20) (Fig. 1A). The structure of albopeptin B was later fully characterized by Oikawa *et al.* in 2022,<sup>21</sup> informed by insights from the biosynthesis genes identified in *S. albofaciens* JC-82-120 (details described in Section 2.4).

### 2.2. Lipolanthines: microvionin and nocavionin

In 2018, with the advancement of genome mining techniques for discovering RiPP natural products, the Süßmuth group reported the isolation and full structural characterization (except for the stereochemistry of a quaternary carbon) of a new class of antibiotics, named lipolanthines. These are hybrids of an *N,N'*-dimethylguanidyl fatty acid and an avionin-type RiPP (see below). The identified lipolanthines include microvionin, produced by *Microbacterium arborescens* 5913, and nocavionin, from *Nocardia terpenica* DSM44935 (ref. 22) (Fig. 1A). A putative BGC was identified for microvionin (the *mic* BGC) and nocavionin (the *noc* BGC); these BGCs were classified as type I lipolanthine BGCs. They contrast with type II lipolanthine BGCs, which additionally contain type I polyketide synthase (PKS)-encoding gene(s). Microvionin exhibited potent growth-inhibitory activity toward both methicillin-susceptible *Staphylococcus aureus* (MSSA) and methicillin-resistant *S. aureus*, with minimum inhibitory concentration (MIC) values of 0.1 and 0.46  $\mu\text{g mL}^{-1}$ , respectively (Table 1).

The RiPP moiety of lipolanthines contains a bicyclic octapeptide featuring a central quaternary carbon atom, referred to as avionin (Fig. 1B), which is generated through the cooperative action of a class III lanthipeptide synthetase (LanKC)<sup>23</sup> and a cysteine decarboxylase (LanD).<sup>24</sup> LanD catalyzes the decarboxylation of a C-terminal cysteine residue to generate 2-aminovinyl-cysteine (AviCys). Meanwhile, the kinase domain of LanKC phosphorylates the hydroxyl groups of two serine residues of the core peptide in an ATP-dependent manner and subsequently eliminates the phosphate to generate dehydroalanine residues. The formation of the avionin structure proceeds *via* a thiol attack on the exomethylene of one dehydroalanine, followed by a nucleophilic attack by the resulting enol on the second dehydroalanine. This sequence of 1,4-additions forms the characteristic bicyclic structure of the avionin-containing peptide (Fig. 1C).





**Fig. 1** Chemical structures of type I lipolanthines from Actinobacteria. (A) Representative compounds include albopeptin B from *Streptomyces albofaciens* and *S. nigrescens* HEK616;<sup>21,31</sup> microvionin from *Microbacterium arborescens*;<sup>22</sup> nocavionin from *Nocardia terpenica*;<sup>22</sup> nocaviogua from *Nocardia* sp. XZ19\_369;<sup>25</sup> and solabiomycons A (R = CH<sub>3</sub>) and B (R = H) from *S. lydicus*.<sup>31</sup> Deoxysolabiomycons A (R = CH<sub>3</sub>) and B (R = H) were obtained from a *S. lydicus*  $\Delta$ solS mutant.<sup>34</sup> (B) Structural features of avionin, labionin, and solabionin.<sup>31</sup> (C) Reaction scheme for avionin formation catalyzed by the enzyme LanKC.<sup>24</sup>



Table 1 Producing strains and bioactivities of RiPP-derived lipopeptides

Isolated product	Group	Original strain	Heterologous expression host	Bioactivity	Reference
Microvionin	Lipolanthine/avionin	<i>Microbacterium arborescens</i> 5913		Methicillin-resistant <i>S. aureus</i> (MRSA) Methicillin-susceptible <i>S. aureus</i> (MSSA)	22
Nocavionin	Lipolanthine/avionin	<i>Nocardia terpenica</i> DSM44935		Weak cytotoxic	22
Nocaviogua A	Lipolanthine/avionin	<i>Nocardia</i> sp. XZ19_369			25
Nocaviogua B	Polyketide/RiPP hybrid	<i>Streptomyces</i> sp. TP-A0584	<i>Streptomyces lividans</i> TK23		26
Goadvionin A1	lipopeptide/avionin	<i>Streptomyces albobacillus</i> JC-82-120		Antifungal	20 and 21
Goadvionin B2	Lipolanthine/solabionin	<i>Streptomyces nigrescens</i> HEK616		Gram-positive bacteria	31
Albopeptin B	Lipolanthine/solabionin	<i>Streptomyces lydicus</i> NBRC 13058		<i>M. tuberculosis</i> H37Rv	31
Solabiomycin A	Lipolanthine/solabionin			<i>M. smegmatis</i>	
Solabiomycin B	Lipolanthine/	<i>Streptomyces lydicus</i> NBRC 13058 ΔsolS		<i>M. smegmatis</i>	34
Deoxysolabiomycin A	labionin			<i>M. smegmatis</i>	
Deoxysolabiomycin B	Selidamide/AviCys	<i>Kamptomena</i> sp. PCC 6506	<i>Escherichia coli</i>	MIC: 25.6 μM	35
Kamptornamide	Selidamide/AviCys	<i>Pseudophaeobacter arcticus</i> DSM 23566	<i>Escherichia coli</i>	MIC: 25.6 μM	35
Phaeornamide	Thioamide/AviCys	<i>Streptomyces olhoviridis</i> NA05001		MIC: 51.2 μM	38
Prethioviridamide	Lipoavitide/AviMeCys	<i>Streptomyces</i> sp. NRRL S-1521	<i>Streptomyces albus</i> J1074	Mitochondrial respiratory chain complex V inhibitor Hemolytic activity	44
Lipoavitide	LAP	<i>Candidatus Entothoonella</i> spp. <i>Streptomyces</i> sp. TP-A0584		P388 murine leukemia cells Anti- <i>Streptomyces</i>	46 55 and 56



The fatty acyl unit of lipolanthines features an N-terminal dimethyl guanidino group. Interestingly, a comparison of BGCs for putative lipolanthines suggests that the terminal guanidine group in microvionin and nocavionin is installed by the combined action of an aldehyde dehydrogenase (MicY or NocY) and a thiamine diphosphate (ThDP)-dependent aceto-lactone synthase (MicJ or NocJ).<sup>22</sup> In contrast, in a type II lipolanthine BGC, which contains a type I PKS, the guanidine group appears to be introduced by an arginine:glycine amidinotransferase, which putatively catalyzes the transfer of a guanidino group from arginine to glycine.<sup>22</sup> However, the product of this type II lipolanthine BGC remains uncharacterized.

In addition to the discovery of nocavionin, two analogs, nocaviogua A and B, with slight differences in the amino acid sequence, have also been isolated from *Nocardia* sp. XZ19\_369 (ref. 25) (Fig. 1A). Nocaviogua A exhibited weak cytotoxic effects against tested cell lines [e.g., non-small cell lung cancer (NCI-H2170) and breast cancer (MDA-MB-231) cell lines], while nocavioguas A and B showed no significant antibacterial or antifungal activity (Table 1). Comparing microvionin and nocavionin analogs, how differences in chemical modifications—such as the presence of *trans* olefins in the fatty acid moiety or variations in the amino acid residues in the RiPP core—affect bioactivity remains unknown.

### 2.3. Polyketide/RiPP-hybrid lipopeptides: goadvionins

Hiroyasu Onaka and colleagues performed genome mining for RiPP BGCs, which led to the identification of the *gdv* BGC for goadvionin in *Streptomyces* sp. TP-A0584. Heterologous expression of the *gdv* BGC in *S. lividans* TK23 resulted in the production of eight congeners and successful isolation of polyketide/RiPP-hybrid lipopeptides, goadvionins A1 and B2 (ref. 26) (Fig. 2A). The RiPP moiety of the goadvionins features a bicyclic octapeptide with an avionin structure, similar to those found in microvionin and nocavionin, but differing in the amino acid residues.

The fatty acyl moiety of goadvionins is highly distinctive, featuring an N-terminal trimethylammonium-containing a linear C31 very-long-chain fatty acid (VLCFA) with multiple hydroxyl groups. Conventional collision-induced dissociation MS/MS was insufficient to elucidate its structure because of limited C–C bond cleavage and low resolution. To address this, radical-induced dissociation techniques—specifically hydrogen attachment/abstraction dissociation (HAD)–MS/MS—were employed.<sup>27</sup> This method uses interactions between precursor ions and neutral hydrogen radicals (H<sup>•</sup>), enabling efficient C–C bond cleavage in mild conditions with high mass resolution.<sup>28</sup> Application of HAD–MS/MS allowed precise localization of all hydroxyl groups within the VLCFA moiety of goadvionins.<sup>26</sup>

In the biosynthesis of the fatty acyl group of goadvionin, it is proposed that, first, L-lysine is converted to trimethyllysine by a methyltransferase (GdvMT), and then the  $\alpha$ -amine is oxidized by an NADH-dependent dehydrogenase (GdvY). A ThDP-dependent enzyme, GdvJ, is assumed to catalyze the decarboxylation of the substrate, forming an aldehyde product. After carboxylation of the aldehyde by unknown enzyme, acyl CoA

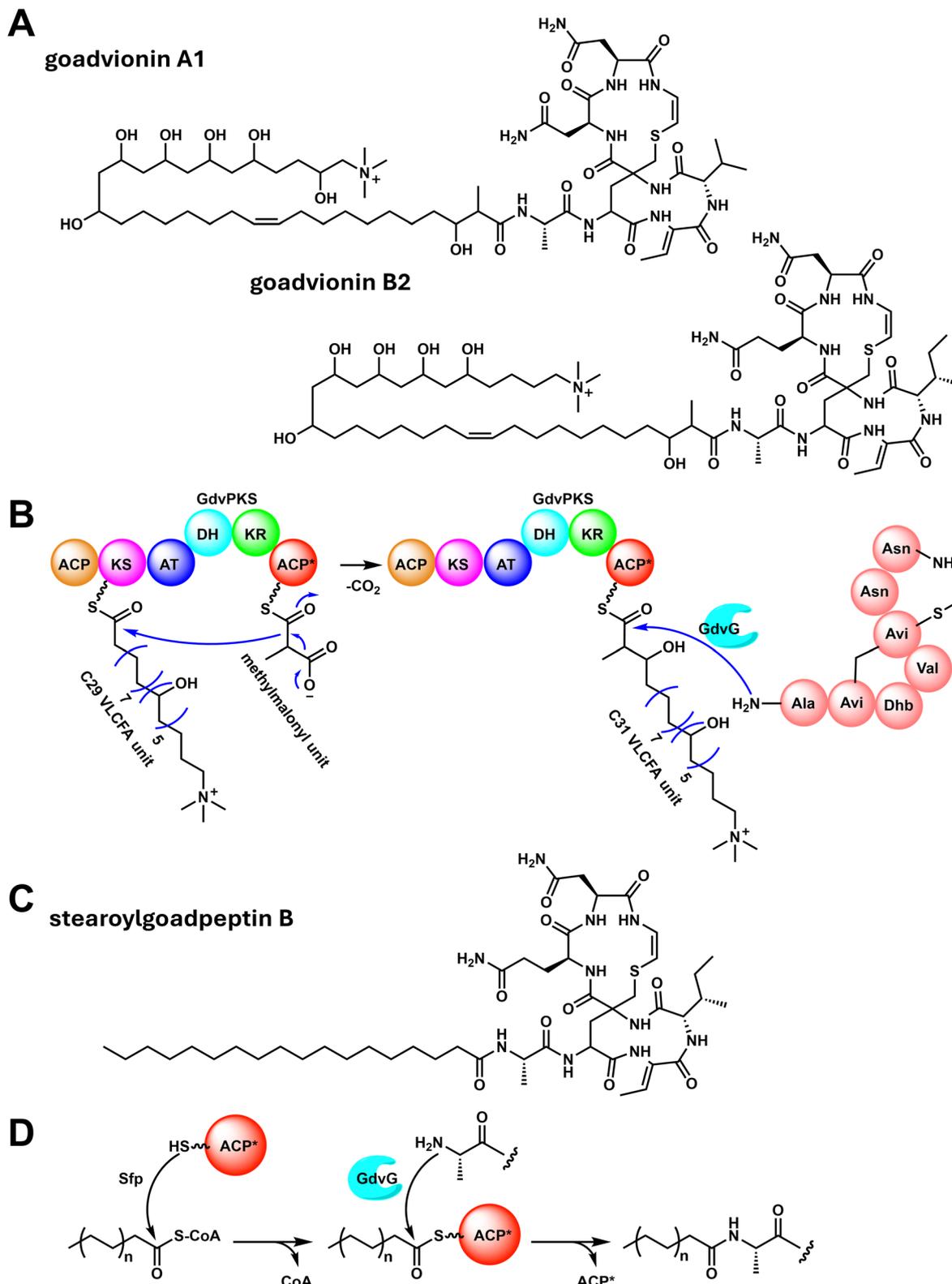
ligase (GdvI) generates the CoA form, which can be loaded onto the acyl carrier protein (ACP) for chain elongation (Fig. 3). This pathway was suggested by genetic work and incorporation studies of <sup>13</sup>C<sub>6</sub>-lysine, which showed a 5 Da mass shift compared with incorporation of <sup>12</sup>C<sub>6</sub>-lysine; were the full <sup>13</sup>C<sub>6</sub>-lysine molecule to be incorporated, there would be a mass shift of 6 Da. The observation of a 5 Da shift indicates that one carbon is lost by decarboxylation.<sup>26</sup> This mechanism is likely conserved in the synthesis of NRP-derived lipopeptides, e.g., fusaricidin A.<sup>29</sup> Interestingly, the incorporation pattern of L-arginine is different in biosynthesis of type I polyketides, e.g., azalomycins from *S. violaceusniger* DSM4137. Azalomycins consist of a 34-membered macrolactone ring and a guanidine-containing side-chain. Arginine monooxygenase and 4-guanidinobutyramide hydrolase are predicted to synthesize 4-guanidinylbutyryl-CoA as the starter unit for the polyketide synthesis.<sup>30</sup>

The VLCFA structure of goadvionin is predicted to be assembled through extension with 12 malonyl-CoA units, beginning with the loading of a trimethyllysine-derived C5 starter unit on the GdvACP, a dedicated ACP in goadvionin biosynthesis, via enzymes associated with the fatty acid synthesis (FAS) system (GdvFB–FG–FZ–FI). In the final step of the fatty acyl synthesis, following the transfer of a fatty acyl group onto an N-terminal ACP domain of GdvPKS, a dedicated type I PKS in goadvionin biosynthesis, a C3 unit derived from methylmalonyl-CoA, is incorporated (Fig. 3). This dedicated PKS system for polyketide/RiPP-hybrid lipopeptides is distinct from that for type I lipolanthines,<sup>22</sup> which presumably uses fatty acyl chains generated solely by the FAS system. The dedicated GDV FAS system likely catalyzes only a ketoreduction step during the first five elongation cycles, resulting in the formation of five consecutive hydroxyl groups along the fatty acyl chain. Structural diversification is introduced during the tailoring steps: a desaturase, GdvC, installs a *cis*-olefin, and an oxygenase (GdvH) hydroxylates the C30' position of the terminal trimethylamine moiety observed in goadvionin A1 (Fig. 2). These variations in the acyl moiety along with the two RiPP cores derived from different precursor peptide genes (*gdvA* and *gdvB*) give rise to eight congeners. The enzymatic condensation of the fatty acyl group to the N-terminus of the RiPP (goadvionins) will be described later (please see Section 5). Goadvionin B2 exhibited growth-inhibitory activity toward Gram-positive bacteria, including MSSA, with an MIC value of 6.4  $\mu\text{g mL}^{-1}$  (Table 1).

### 2.4. Lipolanthine: albopeptin B

Before the isolation of microvionin, nocavionin, and goadvionin, Isono *et al.* reported the isolation of albopeptins A and B from *S. albobifaciens* JC-82-120.<sup>20</sup> Structural elucidation based on spectral data proved challenging, presumably because of the poor solubility and the conformational flexibility of the compounds, which resulted in broadened and low-intensity NMR spectra. The discovery of lipolanthines prompted a revision of the structure of albopeptin B (Fig. 1A), which was later carefully recharacterized by Oikawa *et al.*<sup>21</sup> The genome of *S.*





**Fig. 2** Biosynthesis and chemical structures of goadvionin and synthetic analogs. (A) Chemical structures of goadvionins A1 and B2, isolated from *Streptomyces* sp. TP-A0584.<sup>26</sup> (B) Proposed reaction scheme of the *in vivo* acyl carrier protein (ACP)-dependent acyl transferase reaction catalyzed by GdvG during goadvionin biosynthesis.<sup>26</sup> The C29 fatty acyl group synthesized through the sequential actions of GdvFB (KS), FG (KR), FZ (DH), and FI (ER) is predicted to be loaded onto the N-terminal ACP domain of GdvPKS by GdvL (see details in Fig. 3). A final extension with a building block supplied from methylmalonyl-CoA subsequently generates the C31 very long-chain fatty acid (VLCFA) unit on the C-terminal ACP\* domain of GdvPKS. Although a DH domain is present in GdvPKS, it is likely non-functional, as the final product retains a hydroxyl group.



*albofaciens* JC-82-120 was sequenced, leading to the identification of a putative *alb* BGC. The *alb* cluster contains a LanKC homologue responsible for labionin formation (Fig. 1B), but lacks a LanD homologue, which is typically required for AviCys formation and, consequently, for the formation of avionin. As indicated by the genes in the *alb* BGC, the RiPP moiety of albopeptins indeed lacked the AviCys structure. Notably, the sulfur atom in the labionin is oxygenated to form a sulfoxide, giving rise to a structure named “solabionin”<sup>31</sup> (Fig. 1B).

The presence of a free carboxyl group on the RiPP and an unmodified guanidino terminus on the fatty acyl chain results in a zwitterionic molecule. RiPP-derived lipopeptides often feature positively charged functional groups [e.g., a (dimethyl) guanidino or trimethylamine group] at the terminus of the fatty acyl moiety. The importance of such functional groups is demonstrated, for example, by the NRP-derived lipopeptide fusaricidin A; removal of the guanidino group from the fatty acyl chain of fusaricidin A significantly decreases its activity toward *S. aureus*.<sup>32</sup> Albopeptins exhibited growth-inhibitory activity toward several fungi and Gram-positive bacteria, with MIC values of around 12.5  $\mu\text{g mL}^{-1}$ <sup>20</sup> (Table 1).

## 2.5. Lipolanthines: solabiomycins

Inspired by the discovery of type I lipolanthines and goadvionin, genome mining and stable isotope-guided comparative metabolomics were employed to discover further PFAR natural products, solabiomycins.

Goadvionins were shown to incorporate L-lysine, as demonstrated by the incorporation of <sup>13</sup>C<sub>6</sub>-lysine.<sup>26</sup> In contrast, microvionin and nocavionin were predicted to incorporate L-arginine as the starter unit for their fatty acyl moieties.<sup>22</sup> Genome mining was guided by the presence of acetolactate synthase homologues (MicJ, NocJ, and GdvJ), and putative BGCs for PFARs were identified. Strains harboring such putative BGCs were cultured with stable isotope-labeled amino acids such as <sup>13</sup>C<sub>6</sub>, <sup>15</sup>N<sub>4</sub>-L-arginine or <sup>13</sup>C<sub>6</sub>, <sup>15</sup>N<sub>2</sub>-L-lysine, and incorporation was monitored through comparative secondary metabolomic analysis. This stable isotope labeling method resulted in the discovery of solabiomycins A and B from *S. lydicus* NBRC 13058, along with albopeptin B produced by *S. nigrescens* HEK616 (Fig. 1A).<sup>31</sup> However, when <sup>13</sup>C<sub>6</sub>, <sup>15</sup>N<sub>4</sub>-arginine was incorporated, an unexpected mass shift of 9 Da (rather than 8 Da) was observed for both solabiomycins and albopeptins, suggesting that no decarboxylation occurred during biosynthesis of the guanidino fatty acid chain.<sup>31</sup> This result made ambiguous the function of acetolactate synthase homologs (GdvJ, MicJ, NocJ, SolJ, and NapJ) in the biosynthesis.

The involvement of the identified BGCs in the production of solabiomycins and albopeptins was confirmed through gene knockout experiments: disruption of *solG*, encoding a putative Gcn5-related N-acetyltransferase (GNAT) family acyltransferase

in the *sol* BGC, abolished solabiomycin production in *S. lydicus*,<sup>31</sup> while deletion of *napKC*, encoding a putative type III lanthionine synthase in the *nap* BGC, resulted in the loss of albopeptin biosynthesis in *S. nigrescens*.<sup>31</sup>

The antituberculosis activity of solabiomycins was evaluated by assessing the viability of *Mycobacterium tuberculosis* H37Rv through ATP quantification using the BacTiter assay system. Both solabiomycins exhibited a 95% inhibitory concentration (MIC<sub>95</sub>) of 3.125  $\mu\text{g mL}^{-1}$ <sup>31</sup> (Table 1), indicating their potential as antituberculosis agents. However, the mode of action of RiPP-type lipopeptides, as well as the mechanisms by which producer strains resist their own compounds (“self-resistance”), remains largely unknown (see also Section 6). RNA-Seq analysis was performed on the *nap* BGC responsible for albopeptin B biosynthesis in *S. nigrescens* HEK616.<sup>31</sup> Transcription of the genes encoding a putative ABC transporter permease, NapT, and two hypothetical proteins—NapV, which contains a conserved domain associated with cell wall-active antibiotic response proteins,<sup>33</sup> and NapW—was observed alongside the expression of other biosynthesis enzymes. Although further investigation is needed, characterization of these gene products may shed light on the self-resistance mechanism and reveal potential target molecules of RiPP-derived lipopeptides.

## 2.6. Lipolanthines: deoxysolabiomycins

Similar to albopeptins, solabiomycins feature a characteristic sulfoxide group within the labionin moiety, a structural motif referred to as solabionin in the RiPP portion of the molecule (Fig. 1B). A comparative analysis of the chemical structures of albopeptins and solabiomycins, and of the respective *nap* and *sol* BGCs, revealed that the *sol* BGC contains a single putative oxidase gene, *sols*, whereas the *nap* BGC harbors two oxidase-like genes, *napS* and *napN*. Given that albopeptins also possess a unique N-hydroxyl group, the conserved oxidase genes *sols* and *napS* were predicted to be involved in the sulfoxide formation. Gene knockout experiments showed that the *sols* gene, encoding a putative flavin adenine dinucleotide (FAD)-nicotinamide adenine dinucleotide phosphate (NAD(P))-binding protein, is involved in the sulfoxidation of an alkyl sulfide in solabiomycin.<sup>31</sup> Deoxysolabiomycins A and B were isolated from the *S. lydicus*  $\Delta$ *sols* mutant.<sup>34</sup> The catalytic activity of Sols was demonstrated *in vitro* using recombinant protein expressed in *Escherichia coli*. When incubated with the putative substrates (*i.e.*, deoxysolabiomycins) and the cofactors FAD and NADPH, Sols catalyzed sulfoxidation, converting deoxysolabiomycins into solabiomycins.<sup>34</sup>

The bioactivity of deoxysolabiomycins against Gram-positive bacteria, including *Mycolicibacterium smegmatis*, was evaluated. The results indicated that the sulfoxide group is critical for antibacterial activity, because deoxysolabiomycins exhibited significantly decreased growth inhibition compared with the

Abbreviations: KS, ketosynthase; AT, acyltransferase; DH, dehydratase; KR, ketoreductase; ER, enoylreductase. (C) Chemical structure of stearoylgoadpeptin B, generated by *in vitro* synthesis.<sup>26</sup> (D) Reaction scheme of the *in vitro* ACP\*-dependent non-natural acyl transferase reaction catalyzed by GdvG for the synthesis of stearoylgoadpeptin B.<sup>26</sup> ACP\* refers to a truncated version of the C-terminal ACP domain from GdvPKS.



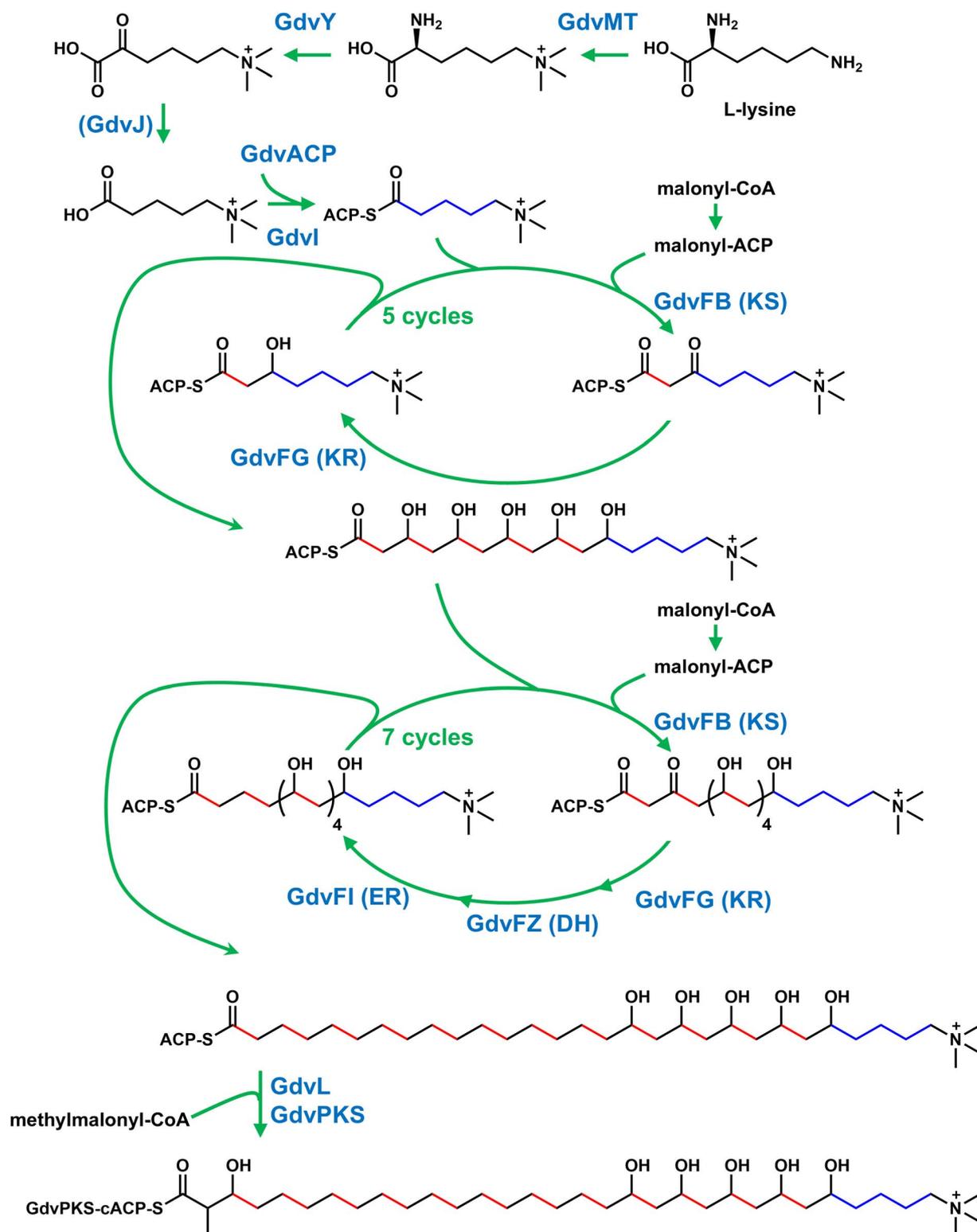


Fig. 3 Proposed biosynthesis pathway of the C31 very-long-chain fatty acid (VLCFA) in goadionin biosynthesis. L-lysine undergoes modification to serve as the trimethylamine-containing C5 starter unit for fatty acid synthesis.<sup>26</sup> In the biosynthesis of the intermediate C29 VLCFA moiety in goadionins, dehydration and enoyl reduction steps are skipped during the first five elongation cycles. The final chain extension to generate a C31 VLCFA is catalyzed by Gdv polyketide synthase (PKS) using methylmalonyl-CoA as the substrate.



equivalent solabimycins. Specifically, the MIC of deoxy-solabimycin A increased from 6.4  $\mu\text{M}$  to 25.6  $\mu\text{M}$  upon removal of the sulfoxide moiety<sup>34</sup> (Table 1).

### 3. Medium-chain fatty acylated RiPPs

#### 3.1. Selidamides: kamptornamide and phaeornamide

In addition to N-terminal acylation of RiPPs by polar functionalized fatty acyl chains, as described above, a RiPP-derived lipopeptide featuring N-acylation on an amino acid side-chain was discovered following heterologous expression of cyanobacterial genes. Jörn Piel and colleagues undertook genome mining and discovered a unique class of RiPP-derived lipopeptides *via* heterologous gene expression in *E. coli*. The identified BGCs from cyanobacteria contained LanM-type class II lanthipeptide synthases<sup>23</sup> along with GNAT family acyltransferases.<sup>35</sup> Putative gene clusters from *Kamptonema* sp. PCC 6506 (*kspA-kspRMN*) and *Pseudophaeobacter arcticus* DSM 23566 (*phaA-phaNRIM*) were heterologously expressed in *E. coli* BL21 (DE3). N-terminal His-tagged precursor peptides (KspA or PhaA) were co-expressed with the associated modification enzymes, and the resulting products—named kamptornamide and phaeornamide respectively—were obtained after cleavage of the leader peptide by the endoproteinase LahT150, which specifically cleaves peptide bonds between consecutive glycine residues<sup>36</sup> (Fig. 4).

These compounds were not detected in the original strains; it thus remains unclear whether they are true natural products or artifacts of heterologous expression. Nonetheless, the products exhibited unique acylation features. Both compounds were N-acylated at the side-chain of an ornithine residue derived from arginine, presumably generated by arginase activity (of KspR or PhaR). Kamptornamide was N-acylated with dodecanoic acid (C12), and phaeornamide was modified with 3-(*S*)-hydroxydecanoic acid (C10) at the side-chain of 4-(*S*)-hydroxyornithine. Because the BGCs did not encode enzymes for fatty acid biosynthesis, it is likely that the fatty acids were supplied by the primary metabolism of *E. coli*. However, *E. coli* does not typically accumulate medium-chain fatty acids such as C10 and C12 forms; so, how, specifically, the fatty acid modifications take place is interesting. A possible explanation is that the culture temperature (isopropyl  $\beta$ -D-1-thiogalactopyranoside induction at 18  $^{\circ}\text{C}$ ) may have affected the composition of the *E. coli* fatty acids.<sup>37</sup> Additionally, because the BGCs do not encode a dedicated ACP, this observation raises the possibility that the acylation steps catalyzed by KspN and PhaN, which have a conserved CoA-binding pocket predicted in their primary structures, are likely to use acyl-CoA as substrates rather than acyl-ACP (which is observed in the biosynthesis of goadivonin). However, this hypothesis requires experimental validation. Because of their structural features, in which N-acylation occurs on the amine group of an ornithine side-chain, this compound

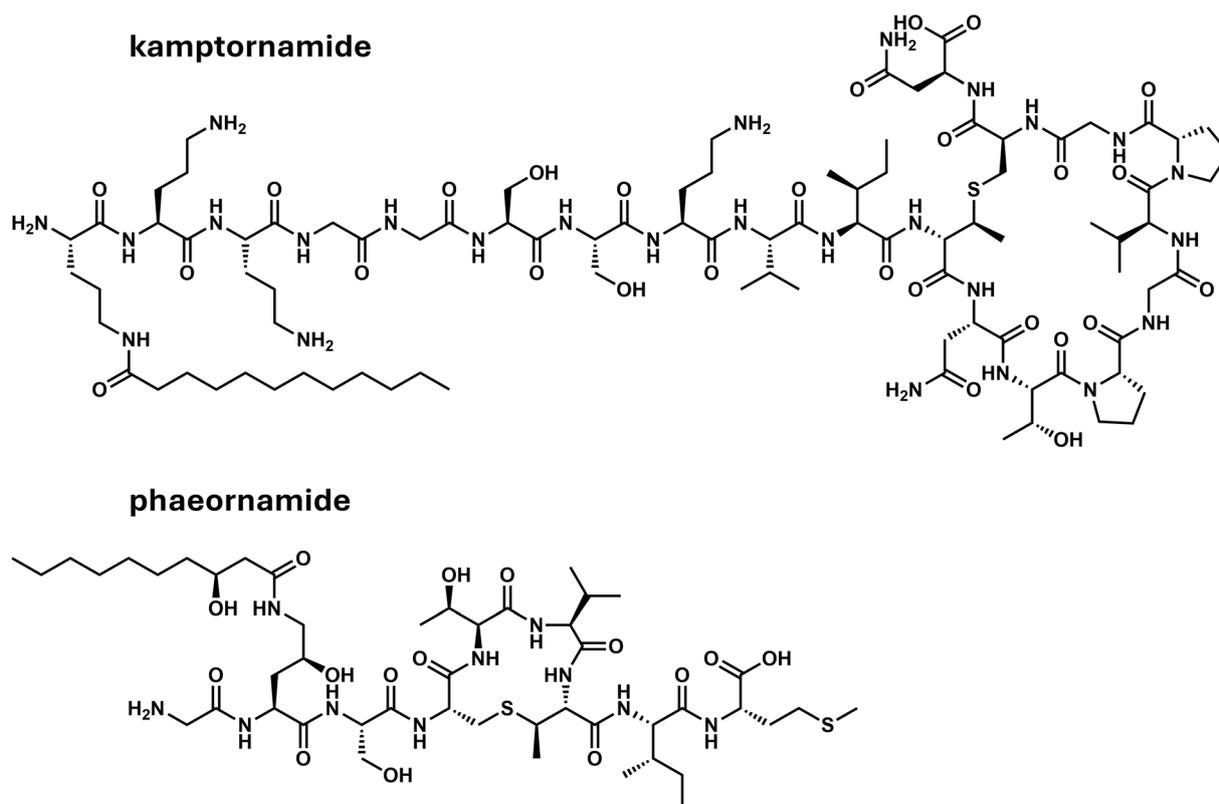


Fig. 4 Chemical structures of selidamides of cyanobacterial origin. Kamptornamide was identified following the heterologous expression of *ksp* genes from *Kamptonema* sp. in *Escherichia coli*.<sup>35</sup> Phaeornamide was obtained following the heterologous expression of *pha* genes from *Pseudophaeobacter arcticus* in *E. coli*.<sup>35</sup>



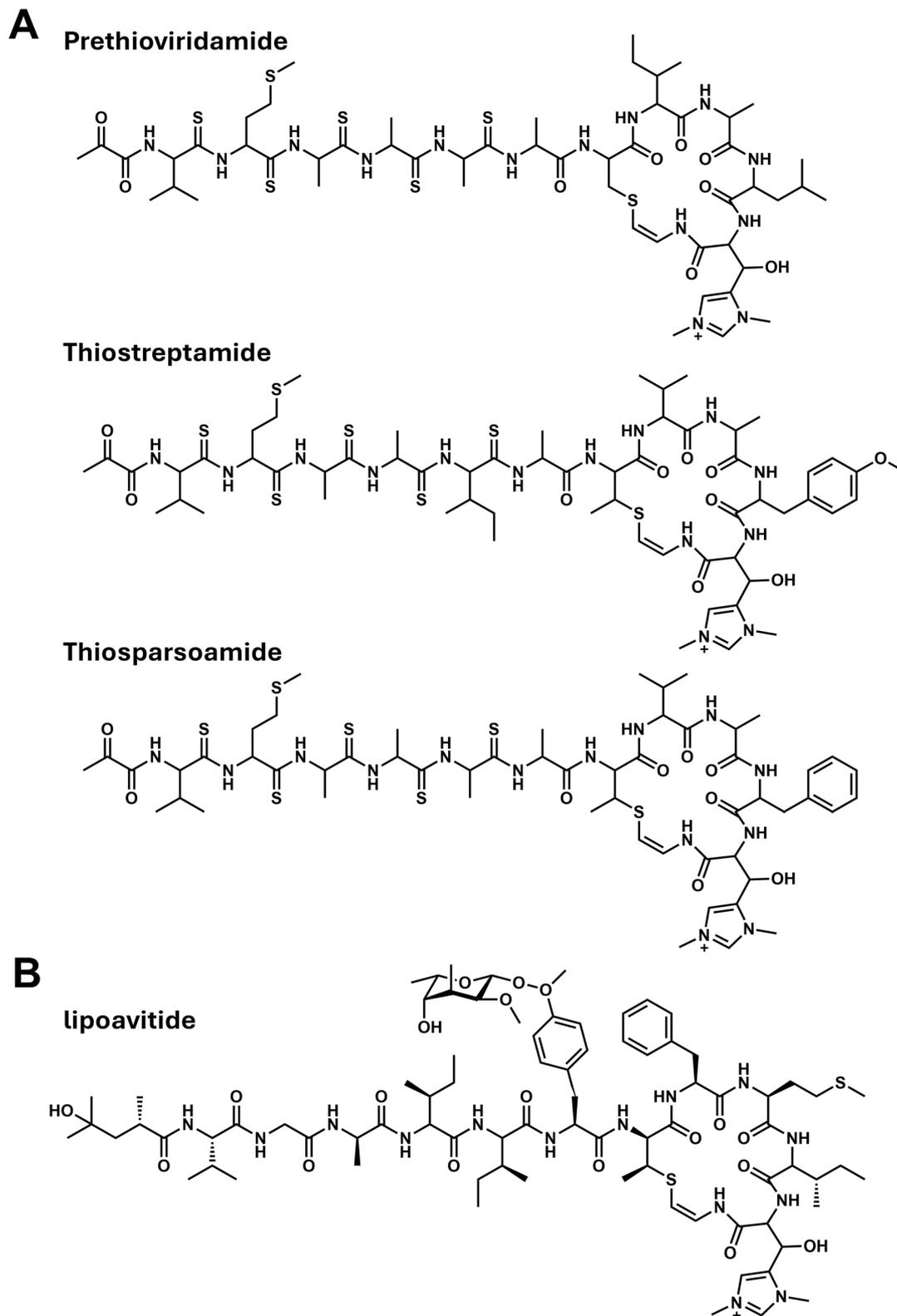


Fig. 5 Chemical structures of 2-aminovinyl-cysteine (AviCys)-containing peptides from Actinobacteria. (A) Prethioviridamide from *S. olivoviridis* NA05001,<sup>38</sup> thiostreptamide from *Streptomyces* sp. NRRL S-4,<sup>42</sup> and thiosparsoamide from *S. sparsogenes* ATCC 25498.<sup>43</sup> (B) Lipoavitide from *Streptomyces* sp. NRRL S-1521.<sup>44</sup>



class was named selidamides (“*selida*” means “side” in ancient Greek). The bioactivities of these compounds have not been elucidated.

## 4. Short-to-medium-chain acylation

### 4.1. AviCys-containing thioamide: prethioviridamide

Relatively short-chain acyl groups have been found in several RiPP natural products. Thioviridamide is an AviCys-containing cyclic peptide, with, uniquely, five thioamides in its side-chain. It was found to contain a 2-hydroxy-2-methyl-4-oxopentanoyl (HMOP) group at its N-terminus. Thioviridamide was isolated from *S. olivoviridis* NA05001.<sup>38</sup> It was later revealed that the true metabolite is prethioviridamide,<sup>39</sup> which contains a pyruvyl group at the N-terminus of the RiPP core (Fig. 5A). Investigation of the *tva* BGC in *S. olivoviridis* NA05001 showed that it lacks an acyltransferase homolog. It was therefore proposed that the

pyruvyl moiety is formed from the N-terminal serine residue of the peptide *via* dehydration, enamine–imine tautomerization, and hydrolysis (Fig. 6A).<sup>40</sup>

Prethioviridamide induces selective apoptosis of various cancer cells—particularly those expressing the adenoviral oncogene E1A—by binding to and inhibiting mitochondrial respiratory chain complex V (F<sub>1</sub>F<sub>0</sub>-ATP synthase)<sup>41</sup> (Table 1).

The formation of AviCys-containing macrocycles is a unique feature of these thioamide-containing RiPP natural products. In the biosynthesis of thiostreptamide S4 from *Streptomyces* sp. NRRL S-4 (Fig. 5A), it is predicted that a phosphotransferase (TsaC) and a HopA1-like protein (TsaD), a class V lanthipeptide synthetases, catalyze the formation of dehydroalanine (or dehydrobutyryne), while a LanD-like enzyme (TsaF) generates the AviCys residue. Notably, a canonical lanthionine synthetase (such as LanKC or LanM), which are typically involved in the macrocyclization steps

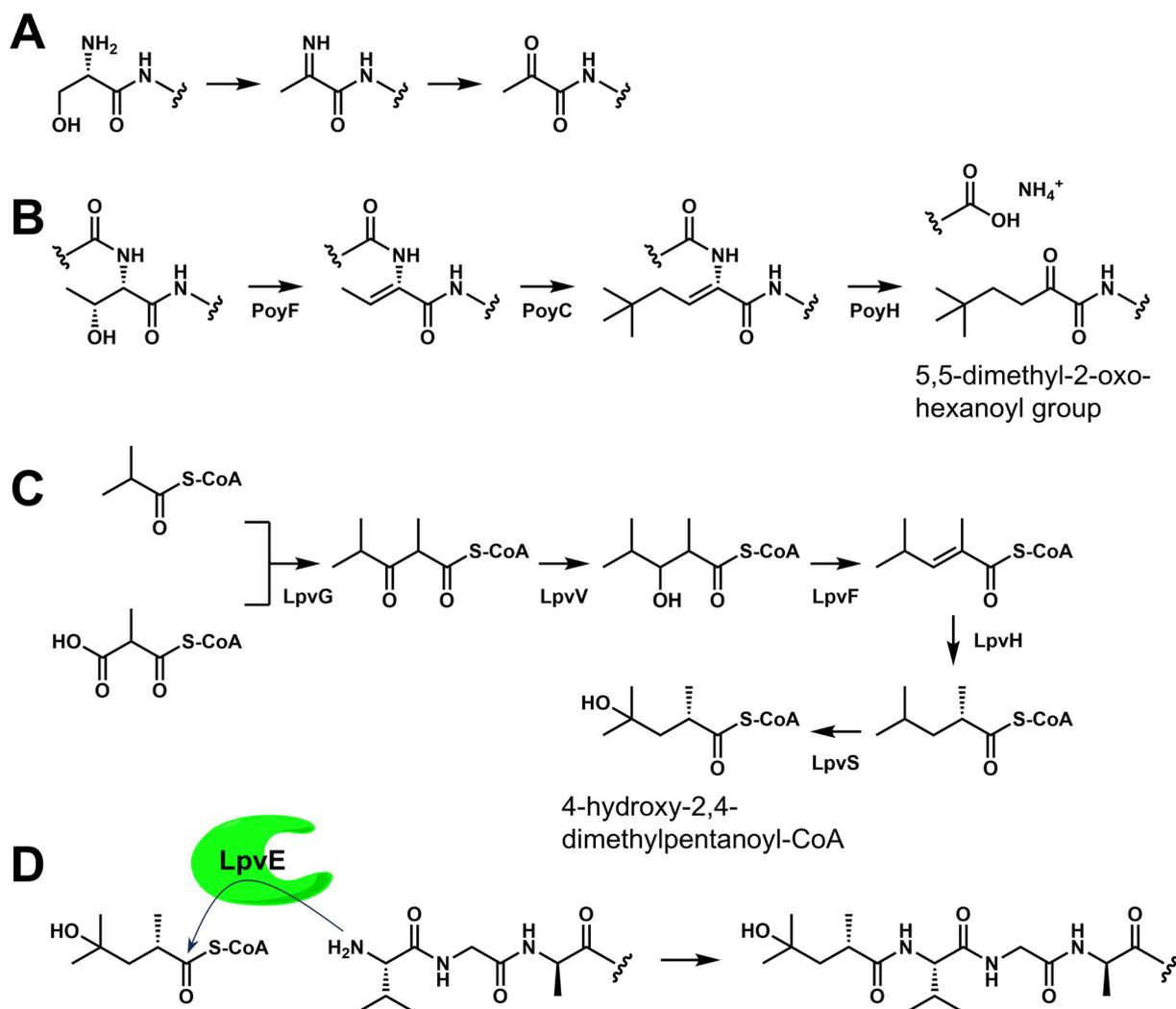


Fig. 6 N-Terminal acyl modifications in ribosomally synthesized and post-translationally modified peptide (RiPP) biosynthesis. (A) Formation of the N-terminal pyruvyl moiety from a serine residue in prethioviridamide biosynthesis.<sup>40</sup> (B) Formation of the N-terminal 5,5-dimethyl-2-oxo-hexanoyl group from a threonine residue in polytheonamide biosynthesis.<sup>50</sup> (C) Proposed biosynthesis pathway of 4-hydroxy-2,4-dimethylpentanoyl-CoA (HMP-CoA) from isobutyryl-CoA and methylmalonyl-CoA in lipoavotide biosynthesis.<sup>44</sup> (D) CoA-dependent N-acyltransferase reaction catalyzed by LpvE in lipoavotide biosynthesis.<sup>44</sup>



in the formation of lipolanthines, goadvionins, and selidamides, is absent from the BGCs for biosynthesis of RiPPs with thioamide groups (*i.e.*, *tva* and *tsa* for prethioviridamide and thiostreptamide, respectively).<sup>42</sup> Interestingly, in the biosynthesis of thio-sparsoamide in *S. sparsogenes* ATCC 25498, a LanKC homolog (SpaKC) located outside the *spa* cluster was found to catalyze the dehydration of a serine residue and was involved in the formation of the AviCys-containing macrocycle with LanD homologue SpaF.<sup>43</sup> Further experimental validation is required to reveal the functions of the phosphotransferase and HopA1-like protein conserved in the thioamide BGC.

#### 4.2. AviMeCys-containing lipopeptides: lipoavitides

In thioamide biosynthesis, the N-terminal pyruvyl moiety is predicted to be derived from a serine residue (Fig. 6A). A newly identified class of fatty acylated RiPPs features an AviCys-containing core with structural similarity to thioamides (Fig. 5B). Huimin Zhao and colleagues performed genome mining for RiPP BGCs encoding conserved AviCys (or AviMeCys) biosynthesis genes, such as *tsaC*, *tsaD*, and *tsaF*, known from thiostreptamide S4 biosynthesis. Using the CAPTURE method (Cas12a-assisted precise targeted cloning with *in vivo* Cre-lox recombination), they cloned the *lpv* BGC from *Streptomyces* sp. NRRL S-1521. Heterologous expression in *S. albus* J1074 led to the discovery of lipoavitides, a new class of AviMeCys-containing lipopeptides modified with a 4-hydroxy-2,4-dimethylpentanoyl (HMP) group (Fig. 5B).<sup>44</sup> While lipoavitides lack thioamide groups,<sup>45</sup> they feature unique modifications, including *O*-glycosylation (2-*O*-methyl- $\beta$ -6-deoxygulose) of a tyrosine side-chain and incorporation of a *D*-alanine residue derived from serine.

The biosynthesis pathway of 4-hydroxy-2,4-dimethylpentanoate was systematically investigated.<sup>44</sup> It was found that LpvG, a 3-ketoacyl-ACP synthase III, catalyzes the initial condensation of isobutyryl-CoA and methylmalonyl-CoA to produce 3-oxo-2,4-dimethylpentanoyl-CoA (Fig. 6C). This intermediate undergoes sequential transformations: ketoreduction by LpvV, dehydration by LpvF, and enoyl reduction also by LpvH, yielding 2,4-dimethylpentanoyl-CoA. The P450 enzyme LpvS subsequently hydroxylates this intermediate to generate HMP-CoA. The enzymatic condensation of the fatty acyl group to the N-terminus of the RiPP is described later (please see Section 5). Notably, the acyltransferase LpvE displays substrate promiscuity, accepting various acyl-CoA derivatives to modify the RiPP scaffold.

The lipoavitides exhibit hemolytic activity, with the N-terminal HMP moiety playing a critical role in this bioactivity; removal of the HMP group, leaving only the RiPP core, led to a marked reduction in the hemolytic activity.<sup>44</sup> While the fatty acyl moieties of other RiPP-derived lipopeptides, such as lipolanthins and goadvionins, have been identified, their precise contributions to antibacterial activity remain unclear. Elucidating the role of these lipid modifications will be an important direction for future research.

#### 4.3. Polytheonamide

Polytheonamide, one of the most extensively modified RiPPs, is composed of 48 amino acid residues and features a remarkable

array of modifications, including eight *tert*-leucines, three  $\beta$ -hydroxyvalines, six  $\gamma$ -*N*-methylasparagines, two  $\gamma$ -*N*-methyl- $\beta$ -hydroxyasparagines, and a  $\beta$ , $\beta$ -dimethylmethionine sulfoxide. Notably, 18 of its residues are *D*-amino acids. It was originally isolated from the marine sponge *Theonella swinhoei* by the Fusetani and Matsunaga groups.<sup>46</sup> Uncultivated filamentous bacteria "*Candidatus Entotheonella*" spp. have been proposed as symbionts of marine sponge and a source of marine natural products.<sup>47</sup> It was later found that polytheonamide is produced by symbiotic *Entotheonella* spp., members of the uncultivated candidate phylum "Tectomicrobia," with genome sizes exceeding 9 megabases.<sup>48</sup>

Because of its structural complexity and the presence of numerous nonproteinogenic and *D*-amino acids, polytheonamide was initially thought to be a nonribosomal peptide product. However, Piel and colleagues identified the ribosomal origin of polytheonamide through metagenomic analysis of *T. swinhoei*.<sup>49</sup> The *poy* BGC responsible for polytheonamide biosynthesis was discovered in the assembled metagenomic data, revealing that only eight gene products are required to construct this highly complex molecule.<sup>50</sup> The N-terminus of polytheonamide features a unique 5,5-dimethyl-2-oxohexanoyl group (Fig. 6B). Structural comparison with the core peptide suggests that a threonine residue is transformed into this unusual acyl moiety through a remarkable series of biosynthetic steps, including dehydration catalyzed by PoyF, formal *tert*-butylation at a nonactivated carbon by PoyC, and formation of the 2-oxo group *via* hydrolytic cleavage of the enamide intermediate by PoyH (Fig. 6B). These insights led to the successful biosynthesis of the polytheonamide-related compound, aeronamide A, in the culturable host *Microvirgula aerodenitrificans*, and further enabled the production of polygeonamides, whose biosynthetic gene cluster (BGC) originates from the uncultivated Rhodospirillaceae bacterium BRH-c57.<sup>51</sup>

Polytheonamide B exhibits potent cytotoxicity against P388 murine leukemia cells, with an IC<sub>50</sub> value of 68 pg mL<sup>-1</sup>.<sup>46</sup> Its structure is a right-handed  $\beta$ -helix approximately 45 Å long, which forms a hydrophilic pore with inner diameter ~4 Å. This unique structure enables the formation of transmembrane channels that selectively permeate monovalent cations. The high cytotoxicity of polytheonamide B is attributed to its ability to form single-molecule ion channels in biological membranes.<sup>52</sup> The Inoue group, in addition to accomplishing the total synthesis of polytheonamide,<sup>53</sup> has extensively investigated its structure–activity relationship. Notably, they demonstrated that N-terminal modifications with hydrophobic groups tend to enhance cytotoxic activity.<sup>54</sup> In particular, the palmitamide derivative bearing an N-terminal C16 palmitoyl group exhibited even greater potency than the natural polytheonamide. These findings underscore the critical role of N-terminal acylation in modulating the bioactivity of this peptide.

#### 4.4. N-Acetylated RiPPs

Goadsporin was first isolated from *Streptomyces* sp. TP-A0584 through a screen for secondary metabolites that induce pigment production and morphological differentiation in *S.*



Table 2 Condensation mechanism of RiPP and fatty acyl moiety<sup>a</sup>

Isolated product	Fatty acyl moiety (putative donor substrate)	Acylation position in RiPPs	Putative acyl transferase homologs	Condensation mechanism	Experimental evidence	Reference
Microvionin	Dimethyl guanidine terminal C15 fatty acyl group	RiPP N-terminal	MicB	Fatty acyl-ACP dependent (putative)	N/A	22
Nocavionin	Dimethyl guanidine terminal C13 (with 3 trans double bonds) fatty acyl group	RiPP N-terminal	NocB1/NocB2	Fatty acyl-ACP dependent (putative)	N/A	22
Goadvionins	Trimethylamine terminal C31 fatty acyl group	RiPP N-terminal	GdvG	Fatty acyl-ACP dependent	<i>In vivo</i> (gene disruption): Detection of intermediates <i>In vitro</i> reaction	26
Solabionmycins	Stearoyl group	RiPP N-terminal	GdvG	Fatty acyl-ACP dependent	<i>In vivo</i> (gene disruption): Abolishment of production	31
	Guanidine terminal C13 fatty acyl group	RiPP N-terminal	SolG	Fatty acyl-ACP dependent (putative)	N/A	21 and 31
Albopeptins	Guanidine terminal C13 fatty acyl group	RiPP N-terminal	AlbG/NapG	Fatty acyl-ACP dependent (putative)	N/A	21 and 31
Kamptornamide	C12 fatty acyl group	Side chain amine of N-terminal ornithine residue	KspN	Fatty acyl-CoA dependent (putative)	<i>In vivo</i> (heterologous biosynthesis)	35
Phaeornamide	3-(S)-Hydroxydecanoyl (C10) group	Side chain amine of N-terminal ornithine residue	PhaN	Fatty acyl-CoA dependent (putative)	<i>In vivo</i> (heterologous biosynthesis)	35
Lipoavitide	4-Hydroxy-2,4-dimethylpentanoyl group	RiPP N-terminal	LpvE	Fatty acyl-CoA dependent	<i>In vitro</i> reaction	44

<sup>a</sup> N/A: not available.

*lividans*.<sup>55,56</sup> The *god* BGC was identified *via* heterologous expression, revealing that goadsporin is a ribosomally synthesized linear azole-containing peptide featuring dehydroamino acids and N-terminal acetylation.<sup>57</sup> The GNAT family acetyltransferase GodH was shown to catalyze N-terminal acetylation using acetyl-CoA as a substrate.<sup>58</sup> Whether GodH exhibits substrate promiscuity and can transfer alternative acyl groups—similar to LpvE in lipoavotide biosynthesis—remains to be investigated. Goadsporin likely targets the bacterial signal recognition particle (SRP), because overexpression of paralogous Ffh homologs (the product of the *godI* gene), a component of SRP, confers self-resistance (Table 1).<sup>57</sup>

In addition to goadsporin, N-acetylated RiPPs have been identified in a variety of bacterial species. For example, members of the microviridin family—tricyclic depsipeptides produced by *Microcystis* spp. (cyanobacteria)—feature N-terminal acetylation.<sup>59</sup> Albusnodin, a lasso peptide from *Streptomyces albus*, is also post-translationally modified with an acetyl group.<sup>60</sup> Paenibacillin, a lantibiotic produced by the Firmicute *Paenibacillus polymyxa*, likewise exhibits N-terminal acetylation.<sup>61</sup>

## 5. Diversification of fatty acyl moieties

*In vitro* experiments revealed that, in goadvionin biosynthesis, the GNAT family acyltransferase GdvG recognizes the C-terminal ACP domain in GdvPKS and catalyzes the condensation reaction between a non-natural stearyl (C18) unit loaded onto the ACP and the N-terminus of the RiPP moiety (Fig. 2B–D and Table 2).<sup>26</sup> The transferase reaction was successfully performed using a stearyl substrate, which is markedly different from the native substrate (Fig. 2B), demonstrating the substrate promiscuity of GdvG. This suggests the potential of the enzyme for generating analogs with non-natural acyl groups. In contrast, in lipoavotide biosynthesis, LpvE was shown to recognize acyl-CoA as a substrate,<sup>44</sup> bypassing the need for a specific ACP (Fig. 6D and Table 2). LpvE appears to preferentially recognize shorter acyl chains (less than C6), but also accepts acyl-CoA with a branched chain, an aromatic group, and even an alkynyl group, likely determined by the size of the cavity for the acyl-CoA substrate.<sup>44</sup>

Based on comparison of BGCs with those for goadvionin,<sup>26</sup> it appears that synthesis of microvionin from *M. arborescens*,<sup>22</sup> nocavionin from *N. terpenica*,<sup>22</sup> solabiomycins from *S. lydicus*,<sup>31</sup> and albopeptins from *S. albobacillans* or *S. nigrescens*<sup>21,31</sup> may use ACP-dependent transfer of acyl moieties by a GNAT family acyltransferase, because specific ACP homologues are present in their BGCs (Table 2). In contrast, similar to the BGCs for lipoavotide,<sup>44</sup> the synthesis of kamptornamide from *Kamptornema* sp.<sup>35</sup> and phaornamide from *P. arcticus*<sup>35</sup> may not require an ACP, instead directly using acyl-CoA as a substrate (Table 2).

The range of substrate tolerance of the acyltransferase enzymes enables the incorporation of diverse acyl groups in RiPP bioengineering. However, expanding structural diversity further will require overcoming the challenge of substrate

tolerance toward the RiPP core. Detailed three dimensional structural characterization of relevant enzymes will be essential for more precise and rational protein engineering.

Fatty acyl moieties play a critical role in bioactivity, as demonstrated in NRPS-derived antibiotics such as DAP, where the acyl chain modulates both antibacterial efficacy<sup>62</sup> and hemolytic toxicity.<sup>32</sup> DAP, a semisynthetic derivative of the A21978C complex from *S. roseosporus*, is produced industrially by supplementing toxic decanoic acid.<sup>63</sup> Recently, Kang and colleagues developed a synthetic biology strategy for selective biosynthesis of C10 decanoic acid, streamlining DAP production.<sup>64</sup> Such control over fatty acid chain length could enable synthetic biology approaches to generate novel RiPP-derived lipopeptides.

## 6. Bioactivity – potential molecular targets of RiPP-type lipopeptides

RiPP-derived lipopeptides exhibit diverse bioactivities, including antibacterial, antifungal, and cytotoxic hemolytic effects (Table 1). However, their mechanisms of action remain largely uncharacterized, and further investigation will be essential to fully realize their potential as therapeutic agents. Nevertheless, insights into their modes of action may be drawn from studies of compounds with similar structural features to those found in RiPP-derived lipopeptides:

Peptide antibiotics containing AviCys are known to interact with the bacterial cell wall precursor lipid II, forming pores in the membrane.<sup>65</sup> This dual mechanism—lipid II sequestration combined with pore formation—leads to bacterial cell lysis and death, and has been described for several lanthipeptides, including mutacin-1140 and epidermin. Other lanthipeptides, such as mersacidin, are currently believed to function solely through lipid II sequestration without pore formation.

The NRPS-derived lipopeptide DAP acts by forming calcium-dependent micelles, enabling its insertion into phosphatidylglycerol-containing membranes. Once inserted, it oligomerizes and either forms cation-selective pores or disrupts membrane integrity by lipid extraction, leading to ion leakage and cell death.<sup>66</sup> Another NRPS-derived lipopeptides, polymyxins (*e.g.*, colistin) act by binding to lipid A in lipopolysaccharide (LPS), displacing divalent cations and destabilizing the outer membrane. The lipid tail inserts into the membrane, promoting self-mediated uptake into the periplasm, where colistin targets LPS precursors in the inner membrane. This disrupts membrane integrity, causes cytoplasmic leakage, and leads to bacterial cell death.<sup>67</sup> These NRPS-derived lipopeptides may offer valuable insights into the mechanisms of action of RiPP-derived lipopeptides.

## 7. Concluding remarks

Recent genome mining efforts have uncovered a novel class of RiPP-based natural products that undergo fatty acylation during their biosynthesis, generating hybrid molecules that combine RiPP cores with fatty acid moieties. These RiPP-derived



lipopeptides exhibit antibacterial and antifungal activities, and hold significant promise as therapeutic agents. The RiPP precursors are genetically encoded, but structural diversification can be achieved through *in vitro* post-translational enzymatic modifications.<sup>68–70</sup> Efficient selection of functional variants can be facilitated using target-guided mRNA display screening,<sup>71</sup> which may aid the development of this class of molecules for therapeutic uses.

The molecular diversity revealed through genome mining, combined with rational design for the biological production of RiPP-derived lipopeptides, may advance drug discovery efforts in response to the growing challenge of antimicrobial resistance.<sup>72</sup> Moreover, integrating *in vitro* engineering strategies with *in vivo* screening and production platforms for RiPPs<sup>73,74</sup> holds great promise for the development of novel RiPP-derived lipopeptide therapeutics in the future.

## 8. Data availability

No primary research results, software or code have been included and no new data were generated or analysed as part of this review.

## 9. Conflicts of interest

The author declares no conflict of interest.

## 10. Acknowledgments

This project was supported by a research grant from the Novozymes Japan Research Foundation. We thank Edanz (<https://www.jp.edanz.com/ac>) for editing a draft of this manuscript. This article was made open access with funding support from Kobe University.

## 11. References

- 1 D. Richter and J. Piel, *Curr. Opin. Chem. Biol.*, 2024, **80**, 102463.
- 2 Q. Guo and B. I. Morinaka, *Curr. Opin. Chem. Biol.*, 2024, **81**, 102483.
- 3 P. G. Arnison, M. J. Bibb, G. Bierbaum, A. A. Bowers, T. S. Bugni, G. Bulaj, J. A. Camarero, D. J. Campopiano, G. L. Challis, J. Clardy, P. D. Cotter, D. J. Craik, M. Dawson, E. Dittmann, S. Donadio, P. C. Dorrestein, K. D. Entian, M. A. Fischbach, J. S. Garavelli, U. Goransson, C. W. Gruber, D. H. Haft, T. K. Hemscheidt, C. Hertweck, C. Hill, A. R. Horswill, M. Jaspars, W. L. Kelly, J. P. Klinman, O. P. Kuipers, A. J. Link, W. Liu, M. A. Marahiel, D. A. Mitchell, G. N. Moll, B. S. Moore, R. Muller, S. K. Nair, I. F. Nes, G. E. Norris, B. M. Olivera, H. Onaka, M. L. Patchett, J. Piel, M. J. Reaney, S. Rebuffat, R. P. Ross, H. G. Sahl, E. W. Schmidt, M. E. Selsted, K. Severinov, B. Shen, K. Sivonen, L. Smith, T. Stein, R. D. Sussmuth, J. R. Tagg, G. L. Tang, A. W. Truman, J. C. Vederas, C. T. Walsh, J. D. Walton, S. C. Wenzel, J. M. Willey and W. A. van der Donk, *Nat. Prod. Rep.*, 2013, **30**, 108–160.
- 4 M. M. Zdouc, J. J. van der Hooft and M. H. Medema, *Trends Pharmacol. Sci.*, 2023, **44**, 532–541.
- 5 A. M. Kloosterman, M. H. Medema and G. P. van Wezel, *Curr. Opin. Biotechnol.*, 2021, **69**, 60–67.
- 6 C. Ongpipattanakul, E. K. Desormeaux, A. DiCaprio, W. A. van der Donk, D. A. Mitchell and S. K. Nair, *Chem. Rev.*, 2022, **122**, 14722–14814.
- 7 E. Breukink, I. Wiedemann, C. van Kraaij, O. P. Kuipers, H. G. Sahl and B. de Kruijff, *Science*, 1999, **286**, 2361–2364.
- 8 S. T. Hsu, E. Breukink, E. Tischenko, M. A. Lutters, B. de Kruijff, R. Kaptein, A. M. Bonvin and N. A. van Nuland, *Nat. Struct. Mol. Biol.*, 2004, **11**, 963–967.
- 9 G. L. Conn, D. E. Draper, E. E. Lattman and A. G. Gittis, *Science*, 1999, **284**, 1171–1174.
- 10 B. T. Wimberly, R. Guymon, J. P. McCutcheon, S. W. White and V. Ramakrishnan, *Cell*, 1999, **97**, 491–502.
- 11 M. V. Rodnina, A. Savelsbergh, N. B. Matassova, V. I. Katunin, Y. P. Semenov and W. Wintermeyer, *Proc. Natl. Acad. Sci. U. S. A.*, 1999, **96**, 9586–9590.
- 12 Y. Imai, K. J. Meyer, A. Iinishi, Q. Favre-Godal, R. Green, S. Manuse, M. Caboni, M. Mori, S. Niles, M. Ghiglieri, C. Honrao, X. Ma, J. J. Guo, A. Makriyannis, L. Linares-Otoya, N. Bohringer, Z. G. Wuisan, H. Kaur, R. Wu, A. Mateus, A. Typas, M. M. Savitski, J. L. Espinoza, A. O'Rourke, K. E. Nelson, S. Hiller, N. Noinaj, T. F. Schaberle, A. D'Onofrio and K. Lewis, *Nature*, 2019, **576**, 459–464.
- 13 H. Kaur, R. P. Jakob, J. K. Marzinek, R. Green, Y. Imai, J. R. Bolla, E. Agustoni, C. V. Robinson, P. J. Bond, K. Lewis, T. Maier and S. Hiller, *Nature*, 2021, **593**, 125–129.
- 14 Y. Zheng, Y. Cong, E. W. Schmidt and S. K. Nair, *Acc. Chem. Res.*, 2022, **55**, 1313–1323.
- 15 K. D. Roberts, M. A. Azad, J. Wang, A. S. Horne, P. E. Thompson, R. L. Nation, T. Velkov and J. Li, *ACS Infect. Dis.*, 2015, **1**, 568–575.
- 16 L. Robbel and M. A. Marahiel, *J. Biol. Chem.*, 2010, **285**, 27501–27508.
- 17 Q. Yue, L. Chen, X. Zhang, K. Li, J. Sun, X. Liu, Z. An and G. F. Bills, *Eukaryotic Cell*, 2015, **14**, 698–718.
- 18 R. D. Sussmuth and A. Mainz, *Angew Chem. Int. Ed. Engl.*, 2017, **56**, 3770–3821.
- 19 M. Dell, K. L. Dunbar and C. Hertweck, *Nat. Prod. Rep.*, 2022, **39**, 453–459.
- 20 K. Isono, K. Kobinata, H. Okawa, H. Kusakabe, M. Uramoto, K. Ko, T. Misato, S.-W. Tai, C.-T. Ni and Y.-C. Shen, *Agric. Biol. Chem.*, 1986, **50**, 2163–2165.
- 21 H. Oikawa, Y. Mizunoue, T. Nakamura, E. Fukushi, J. Yulu, T. Ozaki and A. Minami, *Biosci., Biotechnol., Biochem.*, 2022, **86**, 717–723.
- 22 V. Wiebach, A. Mainz, M. J. Siegert, N. A. Jungmann, G. Lesquame, S. Tirat, A. Dreux-Zigha, J. Aszodi, D. Le Beller and R. D. Sussmuth, *Nat. Chem. Biol.*, 2018, **14**, 652–654.
- 23 M. Montalban-Lopez, T. A. Scott, S. Ramesh, I. R. Rahman, A. J. van Heel, J. H. Viel, V. Bandarian, E. Dittmann,



- O. Genilloud, Y. Goto, M. J. Grande Burgos, C. Hill, S. Kim, J. Koehnke, J. A. Latham, A. J. Link, B. Martinez, S. K. Nair, Y. Nicolet, S. Rebuffat, H. G. Sahl, D. Sareen, E. W. Schmidt, L. Schmitt, K. Severinov, R. D. Sussmuth, A. W. Truman, H. Wang, J. K. Weng, G. P. van Wezel, Q. Zhang, J. Zhong, J. Piel, D. A. Mitchell, O. P. Kuipers and W. A. van der Donk, *Nat. Prod. Rep.*, 2021, **38**, 130–239.
- 24 V. Wiebach, A. Mainz, R. Schnegotzki, M. J. Siegert, M. Hugelland, N. Pliszka and R. D. Sussmuth, *Angew Chem. Int. Ed. Engl.*, 2020, **59**, 16777–16785.
- 25 S. Chang, Y. Luo, N. He, X. Huang, M. Chen, L. Yuan and Y. Xie, *Front. Chem.*, 2023, **11**, 1233938.
- 26 R. Kozakai, T. Ono, S. Hoshino, H. Takahashi, Y. Katsuyama, Y. Sugai, T. Ozaki, K. Teramoto, K. Teramoto, K. Tanaka, I. Abe, S. Asamizu and H. Onaka, *Nat. Chem.*, 2020, **12**, 869–877.
- 27 H. Takahashi, S. Sekiya, T. Nishikaze, K. Kodera, S. Iwamoto, M. Wada and K. Tanaka, *Anal. Chem.*, 2016, **88**, 3810–3816.
- 28 H. Takahashi, Y. Shimabukuro, D. Asakawa, S. Yamauchi, S. Sekiya, S. Iwamoto, M. Wada and K. Tanaka, *Anal. Chem.*, 2018, **90**, 7230–7238.
- 29 J. Li and S. E. Jensen, *Chem. Biol.*, 2008, **15**, 118–127.
- 30 H. Hong, T. Fill and P. F. Leadlay, *Angew Chem. Int. Ed. Engl.*, 2013, **52**, 13096–13099.
- 31 S. Asamizu, S. Ijichi, S. Hoshino, H. Jo, H. Takahashi, Y. Itoh, S. Matsumoto and H. Onaka, *ACS Chem. Biol.*, 2022, **17**, 2936–2944.
- 32 N. Bionda, J. P. Pitteloud and P. Cudic, *Future Med. Chem.*, 2013, **5**, 1311–1330.
- 33 S. Boyle-Vavra, S. Yin, D. S. Jo, C. P. Montgomery and R. S. Daum, *Antimicrob. Agents Chemother.*, 2013, **57**, 83–95.
- 34 S. Ijichi, S. Hoshino, S. Asamizu and H. Onaka, *Bioorg. Med. Chem. Lett.*, 2023, **89**, 129323.
- 35 F. Hubrich, N. M. Bosch, C. Chepkirui, B. I. Morinaka, M. Rust, M. Gugger, S. L. Robinson, A. L. Vagstad and J. Piel, *Proc. Natl. Acad. Sci. U. S. A.*, 2022, **119**, e2113120119.
- 36 S. C. Bobeica, S. H. Dong, L. Huo, N. Mazo, M. I. McLaughlin, G. Jimenez-Oses, S. K. Nair and W. A. van der Donk, *Elife*, 2019, **8**, e42305.
- 37 L. Hoogerland, S. P. H. van den Berg, Y. Suo, Y. W. Moriuchi, A. Zoumaro-Djayoon, E. Geurken, F. Yang, F. Bruggeman, M. D. Burkart and G. Bokinsky, *Nat. Commun.*, 2024, **15**, 9386.
- 38 Y. Hayakawa, K. Sasaki, H. Adachi, K. Furihata, K. Nagai and K. Shin-ya, *J. Antibiot.*, 2006, **59**, 1–5.
- 39 M. Izawa, S. Nagamine, H. Aoki and Y. Hayakawa, *J. Gen. Appl. Microbiol.*, 2018, **64**, 50–53.
- 40 M. Izawa, T. Kawasaki and Y. Hayakawa, *Appl. Environ. Microbiol.*, 2013, **79**, 7110–7113.
- 41 S. Takase, R. Kurokawa, Y. Kondoh, K. Honda, T. Suzuki, T. Kawahara, H. Ikeda, N. Dohmae, H. Osada, K. Shin-Ya, T. Kushiro, M. Yoshida and K. Matsumoto, *ACS Chem. Biol.*, 2019, **14**, 1819–1828.
- 42 T. H. Eyles, N. M. Vior, R. Lacroix and A. W. Truman, *Chem. Sci.*, 2021, **12**, 7138–7150.
- 43 J. Lu, Y. Wu, Y. Li and H. Wang, *Angew Chem. Int. Ed. Engl.*, 2021, **60**, 1951–1958.
- 44 H. Ren, C. Huang, Y. Pan, S. R. Dommaraju, H. Cui, M. Li, M. G. Gadgil, D. A. Mitchell and H. Zhao, *Nat. Chem.*, 2024, **16**, 1320–1329.
- 45 A. Sikandar, M. Lopatniuk, A. Luzhetskyy, R. Muller and J. Koehnke, *J. Am. Chem. Soc.*, 2022, **144**, 5136–5144.
- 46 T. Hamada, S. Matsunaga, G. Yano and N. Fusetani, *J. Am. Chem. Soc.*, 2005, **127**, 110–118.
- 47 E. W. Schmidt, A. Y. Obraztsova, S. K. Davidson, D. J. Faulkner and M. G. Haygood, *Mar. Biol.*, 2000, **136**, 969–977.
- 48 M. C. Wilson, T. Mori, C. Ruckert, A. R. Uria, M. J. Helf, K. Takada, C. Gernert, U. A. Steffens, N. Heycke, S. Schmitt, C. Rinke, E. J. Helfrich, A. O. Brachmann, C. Gurgui, T. Wakimoto, M. Kracht, M. Crusemann, U. Hentschel, I. Abe, S. Matsunaga, J. Kalinowski, H. Takeyama and J. Piel, *Nature*, 2014, **506**, 58–62.
- 49 M. F. Freeman, C. Gurgui, M. J. Helf, B. I. Morinaka, A. R. Uria, N. J. Oldham, H. G. Sahl, S. Matsunaga and J. Piel, *Science*, 2012, **338**, 387–390.
- 50 M. F. Freeman, M. J. Helf, A. Bhushan, B. I. Morinaka and J. Piel, *Nat. Chem.*, 2017, **9**, 387–395.
- 51 A. Bhushan, P. J. Egli, E. E. Peters, M. F. Freeman and J. Piel, *Nat. Chem.*, 2019, **11**, 931–939.
- 52 T. Hamada, S. Matsunaga, M. Fujiwara, K. Fujita, H. Hirota, R. Schmucki, P. Güntert and N. Fusetani, *J. Am. Chem. Soc.*, 2010, **132**, 12941–12945.
- 53 M. Inoue, N. Shinohara, S. Tanabe, T. Takahashi, K. Okura, H. Itoh, Y. Mizoguchi, M. Iida, N. Lee and S. Matsuoka, *Nat. Chem.*, 2010, **2**, 280–285.
- 54 N. Shinohara, H. Itoh, S. Matsuoka and M. Inoue, *ChemMedChem*, 2012, **7**, 1770–1773.
- 55 Y. Igarashi, Y. Kan, K. Fujii, T. Fujita, K. Harada, H. Naoki, H. Tabata, H. Onaka and T. Furumai, *J. Antibiot.*, 2001, **54**, 1045–1053.
- 56 H. Onaka, H. Tabata, Y. Igarashi, Y. Sato and T. Furumai, *J. Antibiot.*, 2001, **54**, 1036–1044.
- 57 H. Onaka, M. Nakaho, K. Hayashi, Y. Igarashi and T. Furumai, *Microbiology*, 2005, **151**, 3923–3933.
- 58 T. Ozaki, K. Yamashita, Y. Goto, M. Shimomura, S. Hayashi, S. Asamizu, Y. Sugai, H. Ikeda, H. Suga and H. Onaka, *Nat. Commun.*, 2017, **8**, 14207.
- 59 N. Ziemert, K. Ishida, A. Liaimer, C. Hertweck and E. Dittmann, *Angew Chem. Int. Ed. Engl.*, 2008, **47**, 7756–7759.
- 60 C. Zong, W. L. Cheung-Lee, H. E. Elashal, M. Raj and A. J. Link, *Chem. Commun.*, 2018, **54**, 1339–1342.
- 61 E. Huang and A. E. Yousef, *Microbiol. Res.*, 2015, **181**, 15–21.
- 62 J. H. Lakey and E. J. A. Lea, *Biochim. Biophys. Acta*, 1986, **859**, 219–226.
- 63 F. M. Huber, R. L. Pieper and A. J. Tietz, *J. Biotechnol.*, 1988, **7**, 283–292.
- 64 C. H. Ji, S. Park, K. Lee, H. W. Je and H. S. Kang, *J. Am. Chem. Soc.*, 2024, **146**, 30434–30442.
- 65 E. S. Grant-Mackie, E. T. Williams, P. W. R. Harris and M. A. Brimble, *JACS Au*, 2021, **1**, 1527–1540.
- 66 W. R. Miller, A. S. Bayer and C. A. Arias, *Cold Spring Harbor Perspect. Med.*, 2016, **6**, a026997.



- 67 E. V. K. Ledger, A. Sabnis and A. M. Edwards, *Microbiology*, 2022, **168**, 001136.
- 68 J. S. Chang, A. A. Vinogradov, Y. Zhang, Y. Goto and H. Suga, *ACS Cent. Sci.*, 2023, **9**, 2150–2160.
- 69 A. A. Vinogradov, M. Shimomura, Y. Goto, T. Ozaki, S. Asamizu, Y. Sugai, H. Suga and H. Onaka, *Nat. Commun.*, 2020, **11**, 2272.
- 70 Y. C. Zhang, K. Hamada, D. T. Nguyen, S. Inoue, M. Satake, S. Kobayashi, C. Okada, K. Ogata, M. Okada, T. Sengoku, Y. Goto and H. Suga, *Nat. Catal.*, 2022, **5**, 682.
- 71 S. E. Iskandar and A. A. Bowers, *ACS Med. Chem. Lett.*, 2022, **13**, 1379–1383.
- 72 C. E. Seyfert, C. Porten, B. Yuan, S. Deckarm, F. Panter, C. D. Bader, J. Coetzee, F. Deschner, K. Tehrani, P. G. Higgins, H. Seifert, T. C. Marlovits, J. Herrmann and R. Muller, *Angew Chem. Int. Ed. Engl.*, 2023, **62**, e202214094.
- 73 T. Si, Q. Tian, Y. Min, L. Zhang, J. V. Sweedler, W. A. van der Donk and H. Zhao, *J. Am. Chem. Soc.*, 2018, **140**, 11884–11888.
- 74 R. S. Ayikpoe, C. Shi, A. J. Battiste, S. M. Eslami, S. Ramesh, M. A. Simon, I. R. Bothwell, H. Lee, A. J. Rice, H. Ren, Q. Tian, L. A. Harris, R. Sarksian, L. Zhu, A. M. Frerk, T. W. Precord, W. A. van der Donk, D. A. Mitchell and H. Zhao, *Nat. Commun.*, 2022, **13**, 6135.

