

REVIEW

View Article Online
View Journal | View Issue



Cite this: *Nat. Prod. Rep.*, 2024, **41**, 990

Opportunities and challenges of RiPP-based therapeutics

Isabel P.-M. Pfeiffer,  † Maria-Paula Schröder  † and Silja Mordhorst  *

Covering: up to 2024

Ribosomally synthesised and post-translationally modified peptides (RiPPs) comprise a substantial group of peptide natural products exhibiting noteworthy bioactivities ranging from anti-infective to anticancer and analgesic effects. Furthermore, RiPP biosynthetic pathways represent promising production routes for complex peptide drugs, and the RiPP technology is well-suited for peptide engineering to produce derivatives with specific functions. Thus, RiPP natural products possess features that render them potentially ideal candidates for drug discovery and development. Nonetheless, only a small number of RiPP-derived compounds have successfully reached the market thus far. This review initially outlines the therapeutic opportunities that RiPP-based compounds can offer, whilst subsequently discussing the limitations that require resolution in order to fully exploit the potential of RiPPs towards the development of innovative drugs.

Received 26th October 2023

DOI: 10.1039/d3np00057e

rsc.li/npr

- 1 Introduction
- 2 Opportunities
 - 2.1 Antibiotic RiPPs
 - 2.2 Antifungal RiPPs
 - 2.3 Antiviral RiPPs
 - 2.4 Antiparasitic RiPPs
 - 2.5 Anticancer/antitumour RiPPs
 - 2.6 Analgesic RiPPs
 - 2.7 Further bioactivities
 - 2.8 Imaging and diagnostic agents
 - 2.9 Synthetic, engineered RiPP natural products
- 3 Advanced RiPP (candidate) drugs
 - 3.1 RiPPs in clinical trials
 - 3.2 RiPPs on the market
- 4 Challenges and limitations
 - 4.1 Oral bioavailability/gastrointestinal stability
 - 4.2 Absorption
 - 4.3 Metabolic stability
 - 4.4 Toxicity
- 5 Concluding remarks
- 6 Author contributions
- 7 Conflicts of interest
- 8 Acknowledgements
- 9 Notes and references

1 Introduction

Ribosomally synthesised and post-translationally modified peptides (RiPPs) constitute a large superfamily of peptide natural products and possess an extensive structural diversity.¹ The different structural elements lead to various biological functions, including a range of therapeutic effects. The biological effects of these compounds rely on chemical motifs, which are introduced by post-translational modifications in precursor peptides that are produced by ribosomes. Most precursor peptides comprise an N-terminal leader peptide and a C-terminal core peptide, while some precursors bear an additional follower peptide at the C-terminus. RiPP biosynthetic gene clusters encode the precursor peptide(s), post-translationally acting maturases, transporter proteins, and often proteases to cleave the modified core sequences from the leader peptide releasing the mature natural product.²

The family of RiPP natural products is continuously expanding and its members encompass redox cofactors,^{3–5} chalkophores,⁶ siderophores,^{7–9} and antimicrobials that participate in natural competition within microbial communities. Although most researchers categorise “classical” RiPPs as bacterial, fungal, and plant specialised metabolites, we also classify certain human, amphibian, and mollusk hormones or toxins as RiPPs in this review, since they also fall under the category of ribosomally synthesised and post-translationally modified peptides. In contrast to “classical” RiPPs, these molecules only contain disulphide bridges as post-translational modifications. Peptides facilitate an array of biological

University of Tübingen, Pharmaceutical Institute, Department of Pharmaceutical Biology, Auf der Morgenstelle 8, 72076 Tübingen, Germany. E-mail: silja.mordhorst@uni-tuebingen.de

† These authors have contributed equally.



processes in all domains of life,⁹ and are believed to be the pharmaceuticals of tomorrow. Their large chemical diversity, typically low toxicity/immunogenicity and their notable target specificity make them a prime subject for drug discovery.^{10,11} The molecular weight of peptide drugs falls between small molecules and large biologicals, such as antibodies. Therefore, peptide drugs are deemed to combine the high specificity of biologicals that are capable of inhibiting protein–protein interactions, with the favourable physicochemical properties of classical small molecule drugs, that are essential for bioavailability. While peptides tend to have low metabolic stability, and linear peptides can display lower binding affinity to their target structure owing to their structural flexibility, modifications can be introduced that incorporate structural characteristics to enhance peptide stability and binding affinity.^{12,13}

Numerous review articles have been published on RiPP research, including general overviews of RiPPs,^{1,2} their physiological and ecological roles,⁹ their modes of action,^{14,15} their antiviral activities,¹⁶ their engineering potential,^{17–19} and reviews on certain RiPP families.^{20–25} This review will concentrate on the potential therapeutic applications of RiPPs and will describe the

promising opportunities RiPPs can offer in drug discovery, as well as the limitations currently preventing their application.

2 Opportunities

RiPP natural products exhibit significant bioactivities in laboratory-based *in vitro* and *in vivo* experiments. Numerous research groups have directed their focus on antibiotic RiPPs, due to the prevalent issue of antibiotic resistance, commonly referred to as “the silent pandemic”. Nonetheless, RiPP natural products have demonstrated additional pharmaceutically relevant effects including antifungal, antiviral, antiparasitic, anti-tumour, and analgesic activities. In the upcoming sections we will present sample compounds for each of the bioactivities mentioned, underscoring the immense potential of this natural product class for pharmaceutical purposes.

2.1 Antibiotic RiPPs

Over 80 antibiotically active RiPPs with distinct targets and modes of action are described to date. Most of them have already been reviewed elsewhere;¹⁵ therefore, only a few examples shall be highlighted here. Along with members of popular RiPP classes, peptides with remarkable activity against drug-resistant and Gram-negative bacteria, and RiPPs with unique targets or modes of action have been chosen for this chapter. First, members of the RiPP class of lanthipeptides, including lanthidins and lipolanthins, are described, followed by linear azol(in)e containing peptides (LAPs), glycocins, thiopeptides, and lasso peptides.

Cinnamycin (also known as Ro 09-0198 or lanthiopeptin) and duramycin (*syn.* PA48009 or leucopeptin, Fig. 1) are two closely related class II lanthipeptides, which are produced by *Streptomyces cinnamoneus*.²⁶ They differ solely in one amino acid and display antibiotic activity against Gram-positive bacteria,^{27–29} by binding to the cell membrane component



Isabel P.-M. Pfeiffer

Isabel Pfeiffer studied Biology (BSc) and Microbiology (MSc) at the University of Tübingen, Germany. Since 2022, she has been a PhD student in the lab of Silja Mordhorst at the department of Pharmaceutical Biology, University of Tübingen. Her research focuses on the investigation of novel bacterial RiPP natural product biosynthetic pathways.



Maria-Paula Schröder

Currently Maria-Paula Schröder is pursuing her PhD at the University of Tübingen in the group of Silja Mordhorst. She has a Bachelor of Science in Chemistry and a Master of Science in Biochemistry from TU Dresden (Germany). Her research focus lies in elucidating new RiPP natural products, exploring their biosynthetic pathways, uncovering the final products, and investigating their potential applications.



Silja Mordhorst

Silja Mordhorst studied Pharmaceutical Sciences and earned her doctoral degree at the University of Freiburg, Germany. In 2019, she started her postdoc as a Feodor-Lynen fellow of the Alexander-von-Humboldt foundation in the natural product lab of Professor Jörn Piel at ETH Zurich, Switzerland, where she investigated novel peptide-modifying enzymes from RiPP biosynthetic gene clusters. In 2022, she was appointed assistant

professor for Pharmaceutical Biology at the University of Tübingen, Germany. Her research includes studies on the discovery and biosynthesis of (mainly RiPP) natural products and the biochemical characterisation of enzymes with a special focus on their potential for peptide engineering.



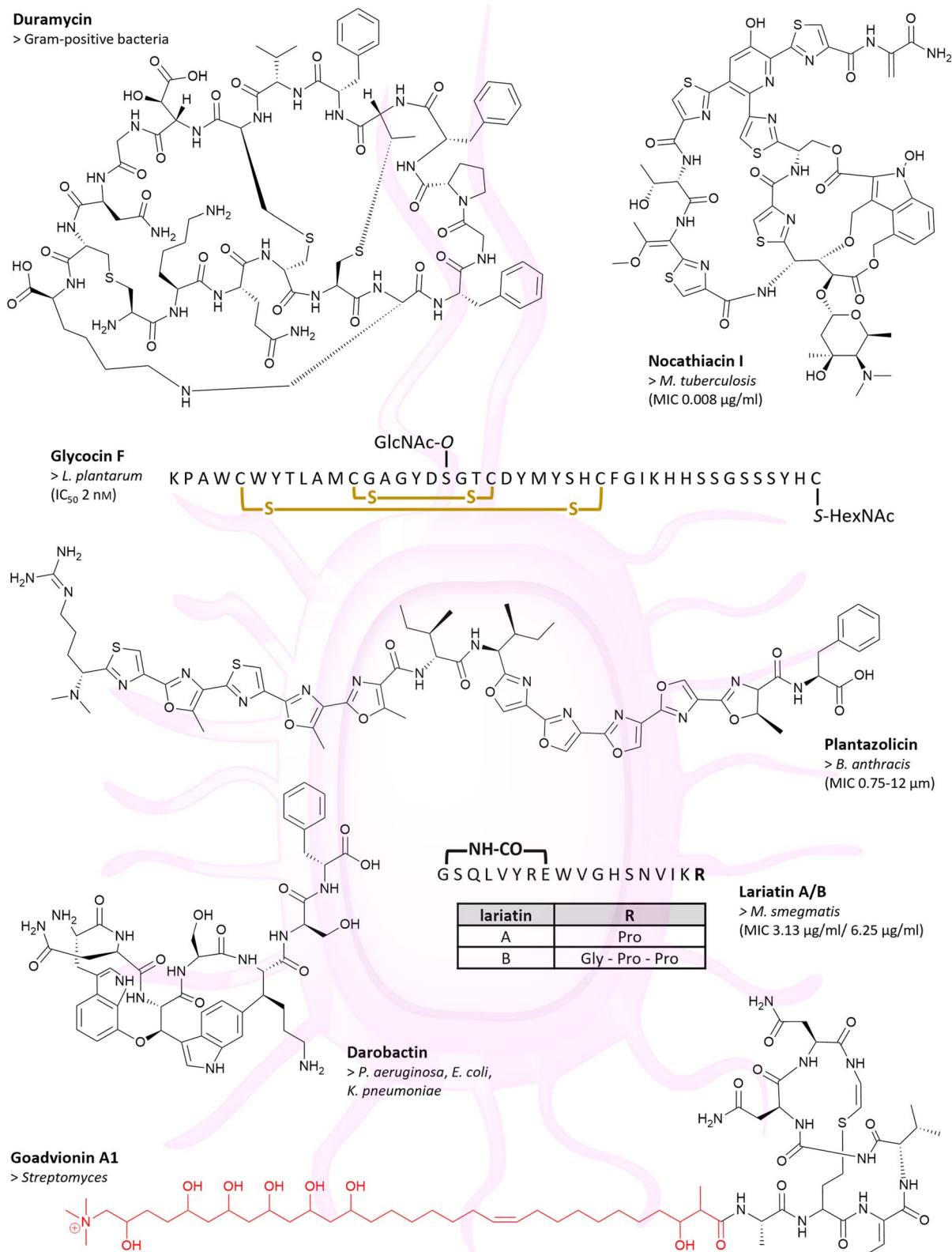


Fig. 1 Many RiPPs show antibiotic activities. Chemical structures of duramycin, nocathiacin I, glycocin F, plantazolicin, darobactin, lariat A/B and goadvionin A1 are depicted. Example(s) of affected bacteria with MIC or IC₅₀ value are listed below the compound name. For glycocin F, amino acids are shown in the single-letter code and yellow indicated crosslinks are disulphide bridges. The acyl group of goadvionin A1 is indicated in red.



phosphatidylethanolamine (PE) in a highly specific manner. While cinnamycin causes membrane disruption, duramycin induces membrane permeabilisation. The precise mechanism of action for both peptides, however, remains unclear.^{15,30}

Lexapeptide and cacaoidin represent the first members of the class V lanthipeptides, which are characterised by an atypical three-component lanthionine synthetase. The two peptides exhibit an *N,N*-dimethylation at their N-termini.³¹ Lexapeptide, derived from *Streptomyces rochei* Sal35, displays antibiotic activity against various Gram-positive bacteria in a low micromolar range, encompassing *Mycobacterium smegmatis* (MIC‡ 0.26 μM , Table 1), § *Enterococcus faecalis* (MIC 0.52 μM), methicillin-resistant *Staphylococcus aureus* (MRSA; MIC 0.52 μM) and *Staphylococcus epidermidis* (MRSE; MIC 1.03 μM). Neither the target nor the mode of action have yet been elucidated.³² Cacaoidin is produced by *Streptomyces cacaoidi* CA-170360 and shows antimicrobial activity against MRSA (MIC 0.22 μM) and *Clostridioides* (formerly *Clostridium*) *difficile* (MIC 1.7 μM). It is believed that cacaoidin targets cell wall biosynthesis, through the inhibition of transglycosylases in the cell wall and through the binding of the cell wall precursor lipid II.^{31,33}

Lipidated lanthipeptides are found in the class of lipolanthines. The first lipolanthine, microvionin, was identified through a bioactivity-guided screening of culture extracts from *Micobacterium arborescens* 5913. This lipolanthine shows potent antibacterial activity against Gram-positive bacteria, notably against MRSA and *Streptococcus pneumoniae* with low minimal inhibitory concentrations of 0.46 $\mu\text{g ml}^{-1}$ and 0.15 $\mu\text{g ml}^{-1}$, respectively.³⁴ Another family of lipolanthines are the goadvionins, which comprise currently eight polyketide/RiPP hybrid lipopeptides, named goadvionin A1–A4 and B1–B4 (Fig. 2). Extracts from a strain that produces goadvionin, namely *Streptomyces* sp. TP-A0584, displayed antibacterial effects against *Streptomyces* and other Gram-positive bacteria, but no effect was observed on Gram-negative bacteria.³⁵

Plantazolicin (Fig. 2), microcin B17, and goadsporin are classified as LAPs. Plantazolicin, a highly selective, narrow-spectrum antibiotic was isolated from *Bacillus amyloliquefaciens* FZB42. It demonstrated noteworthy efficacy against *Bacillus anthracis* (MIC 0.75–12 μM). Following localisation to the cell envelop of *B. anthracis*, depolarisation and lysis of the cellular membrane was observed as a result of plantazolicin treatment. Plantazolicin utilises a distinct mechanism of action in contrast

to other antibiotics targeting the cell envelope. It is suggested that it takes advantage of a transient weakened cell membrane, caused by increased membrane fluidity or changes in the lipid composition, such as an aberrant cardiolipin portion. Cell death ultimately occurs through cell lysis.³⁶ The prototypical example of class B microcins is microcin B17 (MccB17),³⁷ which was isolated from *Escherichia coli* and demonstrated to block DNA replication. In contrast to other DNA gyrase inhibitors, like the quinolones, MccB17 does not bind the A subunit but the B subunit of bacterial DNA gyrase.³⁸ This causes a cascade of reactions. Double-stranded DNA breaks initiate the SOS response, resulting in DNA degradation and, finally, cell death. A single molecule of MccB17 is enough to kill a bacterium.^{39,40} Antibiotic activity of goadsporin, produced by *Streptomyces* sp. TP-A0584, was found against actinomycetes, but not against other bacteria and fungi. Goadsporin was demonstrated to bind the intracellular signal recognition particle (SRP) and interfering with the correct cellular localisation of nascent proteins. But the target of its antibiotic activity remains unknown.^{41,42}

Further compounds with antibiotic activity can be found in the RiPP family of glycosylated bacteriocins, the glycocins. Bacteriocins are synthesised by bacteria and belong to the class of antimicrobial peptides, which demonstrate antibiotic activity only against highly similar strains.⁴³ Glycocin F (or plantaricin KW30, Fig. 1), produced by *Lactobacillus plantarum* KW30, exhibits uncommon structural features. These include an N-acetylglucosamine (GlcNAc), which is β -O-linked to serine 18 of the peptide, and an N-acetylhexosamine (HexNAc), which is S-linked to the C-terminal cysteine 43. Inhibitory activity of glycocin F was observed in *Lactobacillus* strains, particularly *L. plantarum* strains were strongly inhibited (IC₅₀‡ 2 nM). It was demonstrated that GlcNAc is required for bacteriostasis and HexNAc crucial for a maximal efficacy. To our knowledge, the exact mode of action has not yet been reported.^{44,45}

Several thiopeptide antibiotics with diverse structures have been characterised so far. They are unified by the presence of a six-membered heterocycle and a common mode of action, the inhibition of bacterial protein biosynthesis.⁴⁶ Thiostrepton was already isolated in the 1950s from *Streptomyces* sp. and inhibits different strains of *Streptococcus* and *Staphylococcus*.⁴⁷ Potent antibacterial activity against MRSA and vancomycin-resistant *Enterococcus* (VRE) was observed by the thiopeptides thiazomycin (MIC MRSA: 0.032 $\mu\text{g ml}^{-1}$; VRE: 0.064 $\mu\text{g ml}^{-1}$) produced by *Amycolatopsis fastidiosa*,⁴⁸ nocathiacin (Fig. 1, MIC MRSA: 0.007 $\mu\text{g ml}^{-1}$; VRE: 0.03 $\mu\text{g ml}^{-1}$)^{48,49} and nosiheptide (MIC MRSA: 0.03 $\mu\text{g ml}^{-1}$; VRE: 0.125 $\mu\text{g ml}^{-1}$).⁵⁰ Nocathiacins, isolated from *Nocardia* sp. ATCC202099, were additionally effective against *Mycobacterium tuberculosis* (MIC 0.008 $\mu\text{g ml}^{-1}$), *Mycobacterium avium* (MIC 0.06 $\mu\text{g ml}^{-1}$) and like thiazomycin against penicillin-resistant *Streptococcus pneumoniae* (PRSP) with a minimal inhibitory concentration <0.002 $\mu\text{g ml}^{-1}$.^{48,49} Nosiheptide, which is produced by *Streptomyces* sp. CNT-373, was further shown to inhibit a hypervirulent B1 strain of *Clostridioides difficile* (MIC 0.008 $\mu\text{g ml}^{-1}$).⁵⁰

Antimycobacterial activity was detected for the lasso peptides lassomycin and lariatin A and B (Fig. 1). Lassomycin is a unique bactericidal compound produced by *Lentzea*

‡ The following different ratings are employed to characterise the activity/strength of the different compounds. Please find their respective definitions below. EC₅₀ = half maximal effective concentration; GI₅₀ = concentration required for 50% growth inhibition; IC₅₀ = half maximal inhibitory concentration; ID₅₀ = median infectious dose; LC₁₀₀ = lethal concentration (kills 100% of a test sample); LD₅₀ = median lethal dose; LD₉₀ = 90% lethal dose (kills 90% of a test sample); MIC = minimal inhibitory concentration; TD₅₀ = median toxic dose. The lower the value, the more potent (or toxic) the compound is.

§ To help the reader classify the values mentioned, we summarise below typical values of drugs that are in clinical use: MIC values of potent antibiotics and antifungals are in the low μM range. IC₅₀ values of potent antifungals and antivirals are in the submicromolar to low μM range. First line treatments for antiparasitic drugs usually have IC₅₀ values in the nM to low μM range. Chemotherapy drugs that have made it through clinical trial show IC₅₀ values in the nM range for their target.



Table 1 Overview of antibiotic RiPPs^a

Compound name	Bioactivity	Target (if known)	Activity metrics
Achromonodins	Gram-negative bacteria	RNA polymerase	<i>A. pulmonis</i> : MIC 1.3 μM (ref. 55)
Acinetodin	Not observed	RNA polymerase ⁶³	n.d.
Cacaoidin	Gram-positive bacteria	Cell wall	MRSA: MIC 0.22 μM
Capistruin	Gram-negative bacteria	DNA-dependent RNA polymerase	<i>C. difficile</i> : MIC 1.7 μM (ref. 33)
Cinnamycin	Gram-positive bacteria	Cell membrane (PE) ³⁰	<i>P. aeruginosa</i> : MIC 50 μM
Citrocin	Gram-negative bacteria	RNA polymerase	<i>B. caledonica</i> : MIC 12 μM (ref. 57)
Cloacaenodin	Gram-negative bacteria	RNA polymerase	n.d.
Darobactin	Gram-negative bacteria	Outer membrane protein BamA ³⁹	<i>E. coli</i> BW25113: MIC 31 μM (ref. 64)
Duramycin	Gram-positive bacteria	Cell membrane (PE) ³⁰	<i>E. cloacae</i> ATCC 13047: MIC 0.94 μM (ref. 65)
Glycocin F	Gram-positive bacteria	n.d.	n.d.
Goadsporin	Gram-positive bacteria ⁴¹	n.d.	<i>L. plantarum</i> : IC ₅₀ 2 nM (ref. 44)
Goadvionins	Gram-positive bacteria ³⁵	n.d.	n.d.
Klebsidin	Gram-negative bacteria	RNA polymerase	n.d.
Lariatins A & B	Mycobacteria	Mycobacterial cell wall	<i>K. pneumoniae</i> : MIC 256 μM (ref. 63)
Lassomycin	Mycobacteria	ClpC1 ATPase ⁵¹	Lariatins A: <i>M. smegmatis</i> : MIC 3.13 $\mu\text{g ml}^{-1}$
Lexapeptide	Gram-positive bacteria	n.d.	<i>M. tuberculosis</i> : MIC 0.39 $\mu\text{g ml}^{-1}$
			Lariatins B: <i>M. smegmatis</i> : MIC 6.25 $\mu\text{g ml}^{-1}$ (ref. 52)
			n.d.
			<i>M. smegmatis</i> : MIC 0.26 μM
			<i>E. faecalis</i> : MIC 0.52 μM
			MRSA: MIC 0.52 μM
			MRSE: MIC 1.03 μM (ref. 32)
Microcin B17	Gram-negative bacteria ⁶⁶	Bacterial DNA gyrase, subunit B ³⁸	n.d.
Microcin J25	Gram-negative bacteria	DNA-dependent RNA polymerase	<i>S. Newport</i> : MIC 0.01 $\mu\text{g ml}^{-1}$ (ref. 56)
Microcin Y	Gram-positive and Gram-negative bacteria	RNA polymerase	<i>B. subtilis</i> ATCC 6633: MIC 4 μM
Microvionin	Gram-positive bacteria	n.d.	<i>S. infantis</i> : MIC 0.04 μM (ref. 67 and 68)
Nocathiacin	Gram-positive bacteria	Bacterial protein biosynthesis	MRSA: MIC 0.46 $\mu\text{g ml}^{-1}$
			<i>S. pneumoniae</i> : MIC 0.15 $\mu\text{g ml}^{-1}$ (ref. 34)
			MRSA: MIC 0.007 $\mu\text{g ml}^{-1}$
			VRE: MIC 0.03 $\mu\text{g ml}^{-1}$ (ref. 48)
			<i>M. tuberculosis</i> : MIC 0.008 $\mu\text{g ml}^{-1}$ (ref. 49)
			PRSP: MIC <0.002 $\mu\text{g ml}^{-1}$ (ref. 48)
Nosiheptide	Gram-positive bacteria	Bacterial protein biosynthesis	MRSA: MIC 0.03 $\mu\text{g ml}^{-1}$
			VRE: MIC 0.125 $\mu\text{g ml}^{-1}$
			<i>C. difficile</i> : MIC 0.008 $\mu\text{g ml}^{-1}$ (ref. 50)
Plantazolicin	Gram-positive bacteria	Cell membrane	<i>B. anthracis</i> : MIC 0.75–12 μM (ref. 36)
Siamycins	Gram-positive bacteria	Cell wall/lipid II ⁵⁴	n.d.
Thiazomycin	Gram-positive bacteria	Bacterial protein biosynthesis	MRSA: MIC 0.032 $\mu\text{g ml}^{-1}$
			VRE: MIC 0.064 $\mu\text{g ml}^{-1}$ (ref. 48)
Thiostrepton	Gram-positive bacteria	Bacterial protein biosynthesis ⁴⁷	n.d.
Ubonodin	Gram-negative bacteria	RNA polymerase	<i>B. cepacia</i> ATCC 25416: MIC 4 μM (ref. 69)
Marketed (non-RiPP) antibiotics			
Ceftazidime	Gram-positive and Gram-negative bacteria	Cell wall	<i>E. coli</i> : MIC 0.06–0.25 $\mu\text{g ml}^{-1}$ (ref. 70)
Vancomycin	Gram-positive bacteria	Cell wall	<i>Streptococcus pneumoniae</i> : MIC 0.25–0.5 $\mu\text{g ml}^{-1}$ (ref. 71)

^a n.d. = not determined.

kentuckyensis, targeting the ClpC1 ATPase. It triggers the ATPase activity and in parallel abolishes the ClpC1's proteolytic activity. In addition, lassomycin exhibits a high specificity for mycobacteria, but is inactive against bacteria of the human microbiota.⁵¹ Lariatins A is an 18 amino acid peptide, and lariatins B contains two additional amino acids in the tail region, glycine and proline. Both peptides were isolated from

Rhodococcus jostii K01B0171 and specifically inhibited growth of *M. smegmatis* (lariatins A: MIC 3.13 $\mu\text{g ml}^{-1}$; lariatins B: MIC 6.25 $\mu\text{g ml}^{-1}$); *M. tuberculosis* was only inhibited by lariatins A with a MIC of 0.39 $\mu\text{g ml}^{-1}$. They displayed no activity against other bacterial and fungal test organisms. Mycobacteria possess an uncommon cell wall structure, differentiating them from other bacteria. Since the lariatins only inhibited



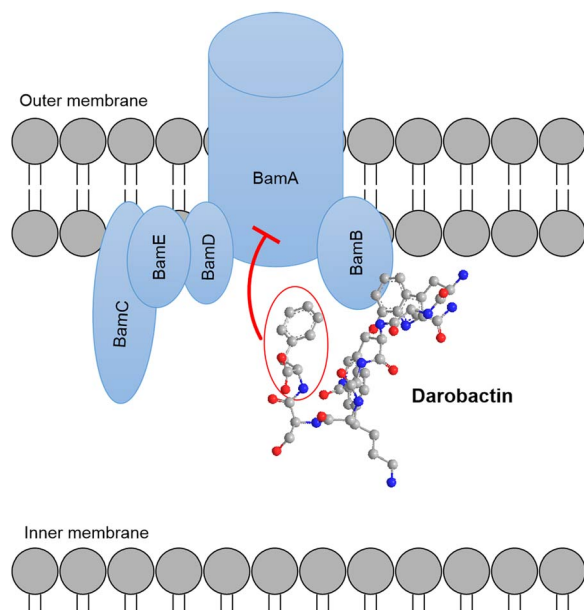


Fig. 2 Darobactin targets the outer membrane protein complex Bam of Gram-negative bacteria. Binding to the subunit BamA leads to inhibition of activity.

mycobacteria, it is assumed they are targeting a step specific for mycobacterial cell wall biosynthesis.⁵² *Streptomyces* sp. AA6532 produces siamycin I (Fig. 5), which also belongs to the class of lasso peptides. Strong antibiotic activity was observed against Gram-positive bacteria, including MRSA and VRE. Siamycin I binds to the pyrophosphate-sugar motif of lipid II at the outside of Gram-positive cell walls. Incorporation of the *N*-acetyl glucosamine-*N*-acetylmuramic acid disaccharide into the cell wall is thought to be hindered by that.^{53,54} The lasso peptides achromonodin-1 and achromonodin-2 identified from *Achromobacter* sp. show narrow-spectrum antibiotic activity against other pathogenic *Achromobacter* sp., which often infect cystic fibrosis patients.⁵⁵

Compounds targeting Gram-negative bacteria are scarce and novel anti-Gram-negative compounds are rarely discovered. The lasso peptides microcin J25 (MccJ25) and capistrin are both highly active against Gram-negative bacteria. MccJ25, isolated from *E. coli*, was found to inhibit the DNA-dependent RNA polymerase (RNAP) and antibiotic activity with a MIC of 0.01 $\mu\text{g ml}^{-1}$ against *Salmonella newport* was demonstrated.⁵⁶ Capistrin, produced by *Burkholderia thailandensis* E264, displays antibiotic activity against *Pseudomonas* and *Burkholderia* strains that are closely related to the producer. For *Pseudomonas aeruginosa* AT27853 a MIC of 50 μM and for *Burkholderia caledonica* a MIC of 12 μM were determined.⁵⁷ Like MccJ25, capistrin was found to bind and inhibit RNAP.⁵⁸ A bactericidal antibiotic with negligible anti-Gram-positive, but strong anti-Gram-negative activity is darobactin (Fig. 1). It is produced by *Photothabdus kharii* HGB1456. Inhibition of clinically relevant drug resistant pathogens, including *P. aeruginosa*, *E. coli* and *Klebsiella pneumoniae* strains, was demonstrated. The effect was explained by darobactin binding the outer membrane protein BamA, thereby preventing

proper assembly of the Gram-negatives outer membrane (Fig. 2).⁵⁹ The binding site of darobactin at BamA is not addressed by other commercially available antibiotics making darobactin a valuable compound to treat infections with multidrug resistant strains. Genetically engineered derivatives of darobactin show an increased antibacterial activity (128-fold increase compared to darobactin) and a promising ADMET (absorption, distribution, metabolism, excretion, and toxicity) profile.^{60–62}

2.2 Antifungal RiPPs

Fungal diseases pose a serious threat to human health. Mortality rates above 50% of invasive fungal infections underline the need for novel antifungal drugs.⁷² Antifungal RiPPs are found in several RiPP families, including lanthipeptides, cyanobactins, thiopeptides, and lasso peptides.

Pinensin A and B (Fig. 3) are two new lanthipeptides obtained from *Chitinophaga pinensis* DSM 28390, representing the first antifungal lantibiotics produced by a Gram-negative bacterium. The pinensins A and B both contain two methyl-lanthionine rings, and differ by an additional alanine at the C-terminus in pinensin A. Both peptides were found to exhibit a weak antibacterial but in combination a broad antifungal activity against filamentous fungi and yeast (MIC 2.1–4.2 $\mu\text{g ml}^{-1}$, Table 2). Their target and the mechanism behind the activity remain elusive.⁷³

Four cyclic heptaepptides of the cyanobactin family, hymenamide A–C (Fig. 3) and hymenamide E, were isolated from the marine sponge *Hymeniacidon* sp. Antifungal activity of these proline-rich peptides was observed against *Cryptococcus neoformans* with a MIC of 133 $\mu\text{g ml}^{-1}$ for congeners A, C, E and 33 $\mu\text{g ml}^{-1}$ for B. Hymenamide A and B showed additionally activity against *Candida albicans* with a MIC of 33 $\mu\text{g ml}^{-1}$ and 66 $\mu\text{g ml}^{-1}$, respectively.^{74,75}

Cyclothiazomycin B1 (Fig. 3) is a cyclic peptide that belongs to the class of thiopeptides. It was isolated from a *Streptomyces* sp. strain and found to be fungistatic. By binding chitin in the fungal cell wall, it inhibits growth of filamentous fungi in a submicromolar range. This leads putatively to cell wall fragility. Activity against bacteria, yeast, or cytotoxic effects on mammalian cells were not observed.⁷⁶

A lasso peptide with high structural similarities to the siamycins, is humidimycin (or MDN-0010, Fig. 5). It was isolated from *Streptomyces humidus* F-100.629. The peptide is not antifungal itself but exhibits a synergistic effect with approved antifungal drugs, like caspofungin (CAS). This makes humidimycin an antifungal enhancer. The antifungal effect of CAS in combination with humidimycin was shown to be enhanced by 4.5-fold, resulting in a remarkable low IC_{50} of 0.007 $\mu\text{g ml}^{-1}$. Its target and the mechanism behind this effect are not yet fully understood. The high osmolarity glycerol (HOG) pathway is assumed to be involved in the response to humidimycin. When treated only with CAS, the HOG pathway may be employed to bypass the CAS-blocked pathway. Thus, tackling a different target than CAS could explain the observed synergistic effect. This could present a possible starting point to enhance CAS activity.⁷⁷



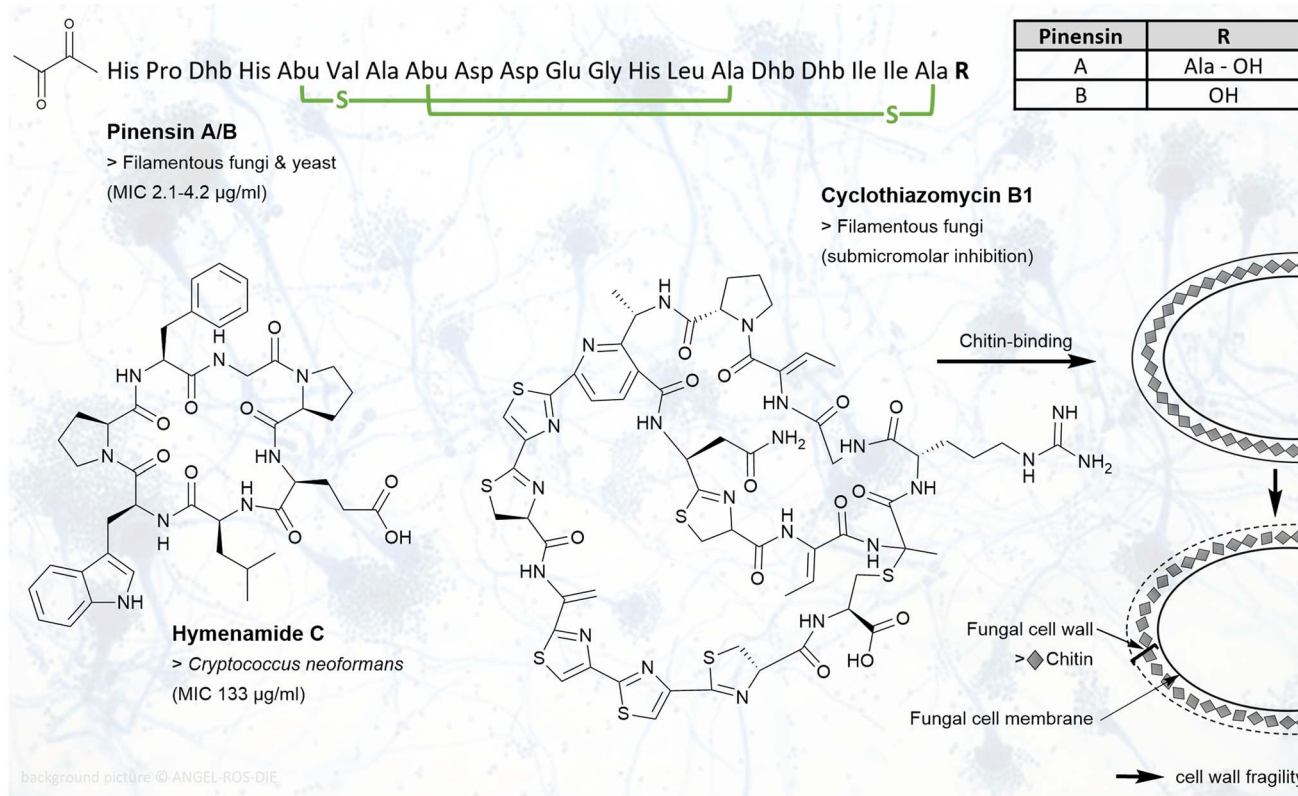


Fig. 3 RiPPs with antifungal activities: chemical structures of pinensin A/B, hymenamide C, and cyclothiazomycin B1 are shown. Example(s) of affected fungi with MIC values are listed below the compound name. For pinensin A/B, the three-letter code is used. Dhb, dehydrobutyrine; Abu, 2-aminobutyric acid. Green indicated crosslinks are lanthionine bridges. For cyclothiazomycin B1, the target chitin in the fungal cell wall is shown. Binding of cyclothiazomycin B1 to chitin leads to cell wall fragility.

2.3 Antiviral RiPPs

Epidemics and pandemics of viral diseases recurred frequently throughout human history. The recent emergence of the severe acute respiratory syndrome coronavirus (SARS-CoV2) and the coronavirus disease 2019 (COVID-19) represents just one example of the keen interest in new antiviral therapeutics.⁸¹ More than 40 peptides from different RiPP families are known to have antiviral activity, including members of proteusins, lasso peptides, and lanthipeptides.¹⁶

Landornamide A (Fig. 3) is a member of the proteusin family and produced by *Kamptonema* sp. PCC 6506. It exhibits rare antiarenaviral activity, inhibiting the infection with lymphocytic choriomeningitis virus (LCMV) with an IC₅₀ of 1.4–2.9 µM (Table 3).⁸² Investigations aiming to determine the target and mode of action of landornamide A are ongoing.

Several RiPPs with anti-HIV activity are known to date. The lasso peptides siamycin I and II differ in position 4: the valine residue is replaced by isoleucine in siamycin II. An ID₅₀† 7 µg ml^{−1} for HIV and ID₅₀ 48 µg ml^{−1} for HSV (herpes simplex virus) was

Table 2 Overview of antifungal RiPPs^a

Compound name	Bioactivity	Target (if known)	Activity metrics
Cyclothiazomycin B1	Filamentous fungi	Fungal cell wall	Submicromolar inhibition ⁷⁶
Humidimycin	Antifungal enhancer	HOG pathway	+Caspofungin IC ₅₀ 0.007 µg ml ^{−1} (ref. 77)
Hymenamides	Non-filamentous fungi	n.d.	<i>C. neoformans</i> : MIC 133 µg ml ^{−1} (hymenamides A, C, and E), ^{74,75} 33 µg ml ^{−1} (hymenamide B) <i>C. albicans</i> : MIC 33 µg ml ^{−1} (hymenamide A) 66 µg ml ^{−1} (hymenamide B) ⁷⁵
Pinensins	Filamentous fungi and yeast	n.d.	Yeast: MIC 2.1–4.2 µg ml ^{−1} (ref. 73)
Marketed (non-RiPP) antifungal drug			
Caspofungin	Filamentous fungi and yeast	Fungal cell wall ⁷⁸	<i>Candida</i> sp.: MIC 0.015–2 µg ml ^{−1} (ref. 79) <i>Aspergillus</i> sp.: MIC 0.25–16 µg ml ^{−1} (ref. 80)

^a n.d. = not determined.



Table 3 Overview of antiviral RiPPs^a

Compound name	Bioactivity	Target (if known)	Activity metrics
Cinnamycin	HSV-1	Viral proliferation ^{15,88}	n.d.
Divamide A	HIV	Cell entry or exit ⁹¹	n.d.
Duramycin	HSV-1, Ebola, West Nile virus, Dengue virus ⁸⁹	Cell entry ^{15,88}	n.d.
Labyrinthopeptins	HIV & HSV	Cell entry	LabyA1: HIV: EC ₅₀ † 0.79–3.3 μM HSV: EC ₅₀ 0.29–2.7 μM LabyA2: HIV: EC ₅₀ of 26 μM (ref. 92) LCMV: IC ₅₀ 1.4–2.9 μM (ref. 82)
Landornamide A	Arenavirus	n.d.	(I) HIV: ID ₅₀ † 7 μg ml ⁻¹ HSV: ID ₅₀ 48 μg ml ⁻¹ (ref. 93)
Siamycins I-III	HIV & HSV	Viral cell fusion	(II) HIV: ID ₅₀ 9 μg ml ⁻¹ HSV: ID ₅₀ 27 μg ml ⁻¹ (ref. 93) (III) HIV-1 reverse transcriptase: IC ₅₀ 4 μg ml ⁻¹ , HIV aspartyl protease: IC ₅₀ 48 μg ml ⁻¹ (ref. 84)
Marketed (non-RiPP) antiviral drugs			
Docosanol	HSV	Cell entry	EC ₅₀ 2.5 mg ml ⁻¹ (ref. 94)
Maraviroc	HIV	Cell entry	IC ₅₀ 43 nM (ref. 95)
Saquinavir	HIV	HIV protease	IC ₅₀ 1–10 nM (ref. 96)

^a n.d. = not determined.

measured for siamycin I. Siamycin II, produced by *Streptomyces* sp. AA3891, displayed an ID₅₀ of 9 μg ml⁻¹ for HIV and an ID₅₀ of 27 μg ml⁻¹ for HSV. Their anti-HIV and anti-HSV activity is hypothesised to be related to blocking viral cell fusion.⁵³ Structural similarities to gp41 transmembrane protein functional domains of HIV-1 were observed in siamycin III (or RP 71955 or arborycin), implying an analogous mode of action of all siamycins. Siamycin III is produced by *Streptomyces* sp. SP9440 and contains an isoleucine at position 4 like siamycin II, and a valine instead of isoleucine at position 17.⁸³ Furthermore, siamycin III was found to

affect HIV-1 by inhibiting the production of the reverse transcriptase (IC₅₀ 4 μg ml⁻¹) and the aspartyl protease (IC₅₀ 48 μg ml⁻¹) of HIV. Knowledge on the exact mechanism of action of the siamycins is not available yet.^{15,84,85} Based on its structural relatedness to the siamycins, humidimycin (mentioned above) is thought to have anti-HIV properties as well.⁸⁶

Activity against HIV and HSV can be attributed to labyrinthopeptin A1 (LabyA1, Fig. 5) and A2 (LabyA2), which belong to a new class of carbacyclic lantibiotics, isolated from *Actinomadura namibiensis* DSM 6313. These compounds contain labionin, an αC quaternary substituted amino acid. This was the first time such a carbocyclic side chain linkage was described. Two serine and one cysteine residue(s) are linked by LabKC to form this unique structural feature. The cell entry of HIV and HSV is inhibited, in case of HIV by interaction of LabyA1 with the envelope protein gp120. Depending on the cell line tested, an EC₅₀† of 0.79–3.3 μM for HIV and an EC₅₀ of 0.29–2.7 μM for HSV was observed. LabyA1 further acts as anti-HIV enhancer. It showed an additive to synergistic effect together with anti(retro)viral drugs such as aciclovir, tenofovir and saquinavir. In contrast, LabyA2 was found to be about 10-fold less potent against HSV than LabyA1 and displayed only an EC₅₀ of 26 μM against HIV. This notable difference in potency may be explained by the structural differences of the labyrinthopeptins. LabyA1 has two additional amino acids and LabyA1 and LabyA2 vary at five amino acid positions.⁸⁷

In addition to their antimicrobial activity, cinnamycin and duramycin exhibit antiviral activity against HSV-1. By targeting phosphatidylethanolamine (PE), the viral proliferation is inhibited.^{15,88} Duramycin effects furthermore Ebola, West Nile and Dengue virus specifically by preventing virus attachment, thereby stopping cellular entry (Fig. 4).⁸⁹

The divamides are cinnamycin-like anti-HIV peptides that were originally isolated from a marine tunicate. Later, it was

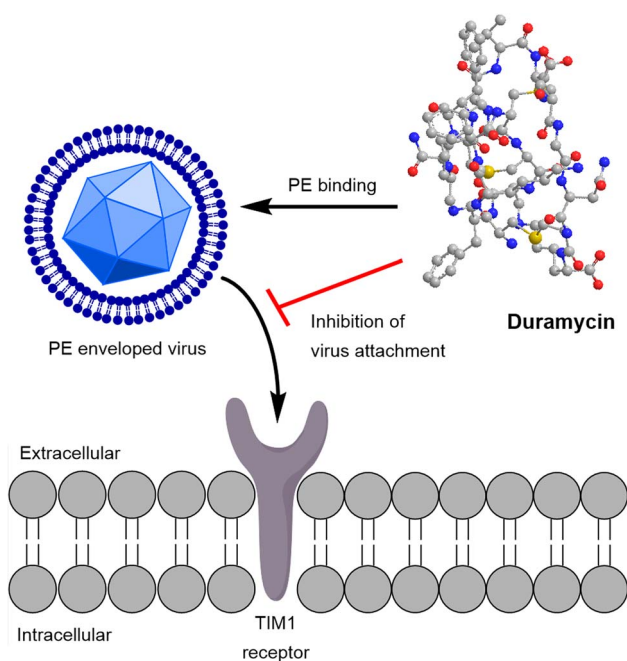


Fig. 4 Duramycin targets enveloped viruses by binding PE (phosphatidylethanolamine). This leads to inhibition of virus attachment to the TIM1 membrane receptor in human cells and blocks cell entry.



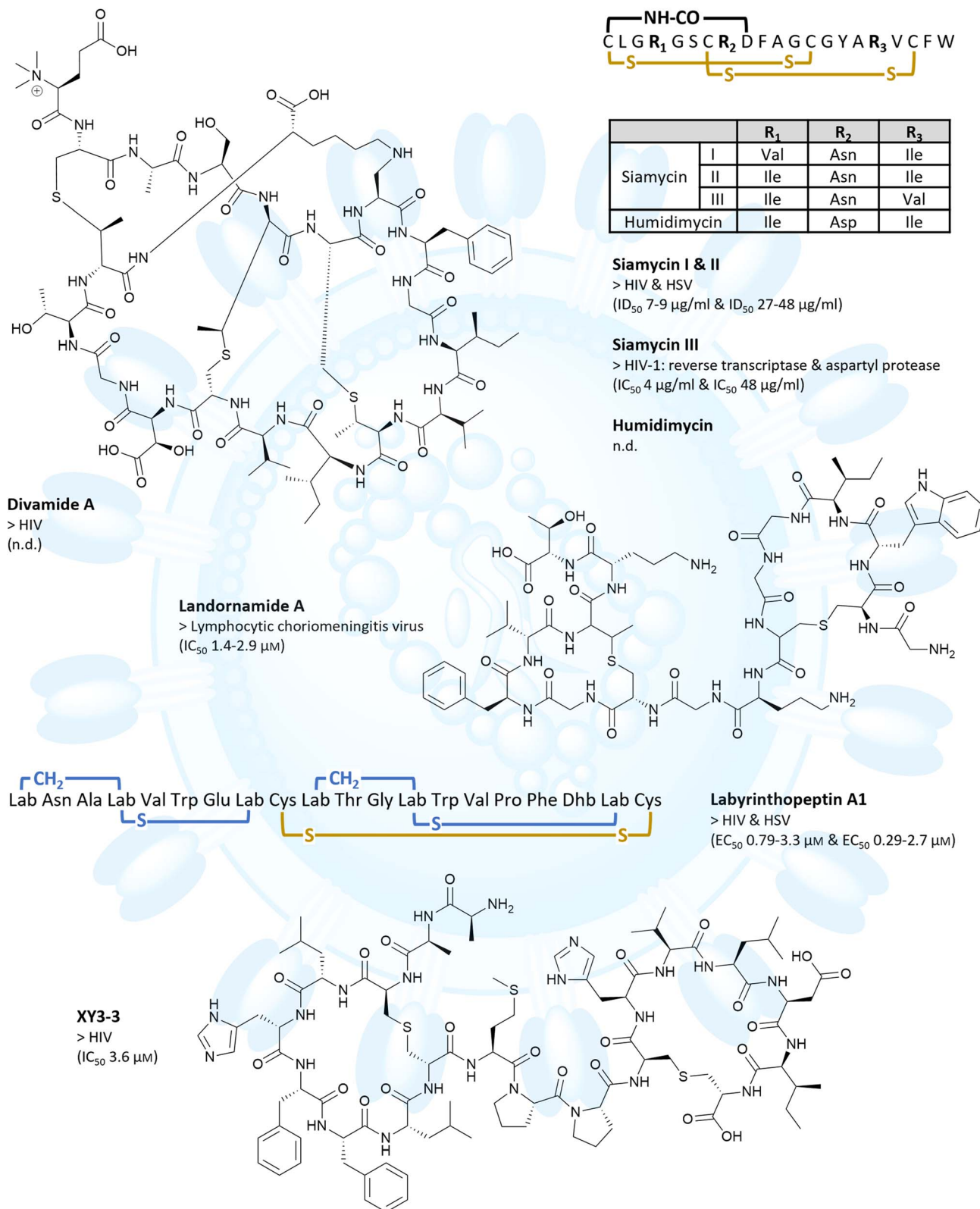


Fig. 5 RiPPs with antiviral activities: chemical structures of divamide A, siamycins and humidimycin, landornamide A, labyrinthopeptin A1, and XY3-3 are depicted. Example(s) of affected viruses with ID₅₀, IC₅₀ or EC₅₀ values are listed below the compound name. For labyrinthopeptin A1, the three-letter code is used. Lab, labionin; Dhb, dehydrobutyrine. Yellow indicated crosslinks are disulphide bridges, blue indicated crosslinks are labionin bridges.



found that divamide A–C were produced by a symbiont of the tunicate, the cyanobacterium *Prochloron didemnid*. Employing bioassay-guided fractionation, the anti-HIV activity could be assigned only to divamide A (Fig. 5). By structure activity relationship studies, the lysinoalanine residue was shown to be crucial for its antiviral activity. The exact underlying mode of action could not be clarified yet. Interaction of divamide A with T-cell membranes was observed, thus it was concluded it may block the viral cell entry or exit. Another bioactivity related discovery was the splitting of divamide A's cytotoxic and antiviral effect by minor changes in the sequence of amino acids.⁹⁰

2.4 Antiparasitic RiPPs

Parasitic diseases affect 3.5 billion people worldwide according to WHO.⁹⁷ Economical and public health effects disproportionately burden the poorest nations and the drugs currently in use to treat these infections often show sub-optimal efficacy or serious toxicity. They were mostly discovered over 50 years ago, and today many drug-resistant strains exist. To combat these infections and mitigate the long lasting effects, it is imperative to find new drug leads.^{97,98} First, we will discuss antimicrobial RiPPs that have additional antiparasitic properties. Subsequently, we will provide examples from the defensin, cyanobactin, and cyclotide families. Defensins are antimicrobial peptides found in eukaryotes and prokaryotes; they are ribosomally synthesised and post-translationally modified, and can therefore be attributed to the natural product class of RiPPs.^{2,99,100}

Besides its antimicrobial activity (see Section 2.1), thiostrepton (Fig. 8) has potential as an antimalarial drug lead. It was shown to target the proteasome of the human malaria parasite *Plasmodium falciparum*. The natural compound itself shows modest antimalarial activity with an IC₅₀ of 8.9 μM

(Table 4), while optimised thiostrepton derivatives achieved IC₅₀ values as low as 1.0 μM .¹⁰¹ Two independent targets were identified, the parasitic 20S proteasome and the large ribosomal subunit of the prokaryotic apicoplast, an essential cell organelle found in parasites belonging to the phylum Apicomplexa. Addressing two targets presents a substantial advantage, as it lessens the risk of resistance development. Additionally thiostrepton-based derivatives show considerable selectivity for the parasite proteasome over the human proteasome and no toxicity to human cell lines, making these compounds attractive antimalarial drug leads.¹⁰¹

The defensins found in the dragonfly *Aeshna cyanea*, known as *Aeshna* defensin, and in the flesh fly *Protophormia terraenovae*, referred to as phormicin, exhibit antiparasitic activity against *P. gallinaceum*.^{102–104} *P. gallinaceum* is a malarial parasite to birds of the genus *Gallus* and it is used as a model system in malaria research.^{105,106} Injecting *Aeshna* defensin or phormicin into the haemolymph of *Aedes aegypti* mosquitos 3–4 days after parasite ingestion significantly reduced the oocyte density, without showing toxicity against the mosquito hosts.¹⁰⁷ A defensin from the sandfly *Phlebotomus duboscqi* active against *Leishmania major* was identified in a *Leishmania* infected sand fly. *Leishmania* spp. are parasites that cause leishmaniasis, a tropical disease with potentially fatal consequences. Recombinant *Phlebotomus* defensin was then tested against *L. major*, i.e. strain MHOM/YE/84, *in vitro* and IC₅₀ values ranged from 68–85 μM , demonstrating antiparasitic activity for the *Phlebotomus* defensin.^{108,109}

Balgacyclamides A–C (Fig. 8) belong to the family of cyanobactins and were isolated from *Microcystis aeruginosa* EAWAG 251. They were tested against a range of parasites, such as *P. falciparum*, *Trypanosoma brucei rhodesiense*, *Trypanosoma cruzi*, and *Leishmania donovani*. *T. b. rhodesiense* is the cause of African trypanosomiasis, which can lead to coma

Table 4 Overview of RiPPs with antiparasitic properties^a

Compound name	Bioactivity	Target (if known)	Activity metrics
<i>Aeshna</i> defensin	Antimalaria	n.d.	Dosed at 1 μg (ref. 107)
Balgacyclamides A–C	Antimalaria	n.d.	IC ₅₀ 8.2–9 μM (ref. 110)
Balgacyclamides A–C	Anti-trypanosomatid	n.d.	IC ₅₀ 51–59 μM (ref. 110)
Cycloviolacin O14	Antihelminthic	n.d.	IC ₅₀ 1.40 μM (ref. 116)
Kalata B1	Antihelminthic	n.d.	IC ₅₀ 3.36 μM (ref. 116)
Mollamide B	Antimalaria	n.d.	IC ₅₀ 2 $\mu\text{g ml}^{-1}$ (ref. 111)
Mollamide B	Leishmanicidal	n.d.	IC ₅₀ 18 $\mu\text{g ml}^{-1}$ (ref. 111)
<i>Phlebotomus</i> defensin	Leishmanicidal	n.d.	IC ₅₀ 68–85 μM (ref. 108)
Phormicin	Antimalaria	n.d.	Dosed at 1 μg (ref. 107)
Thiostrepton	Antimalaria	Proteasome	IC ₅₀ 8.9 μM (ref. 101)
Venturamides	Antimalaria	n.d.	IC ₅₀ 5.2–8.2 μM (ref. 112)
Venturamides	Anti-trypanosomatid	n.d.	IC ₅₀ 14.6–15.8 μM (ref. 112)

Marketed (non-RiPP) antiparasitic drugs

Atovaquone/Proguanil	Antimalaria	Mitochondrial <i>cytb</i> gene	IC ₅₀ 3.4 nM and 36.5 μM (ref. 117)
Nifurtimox	Anti-trypanosomatid	Inhibition trypanozhione reductase, oxidative stress ¹¹⁸	IC ₅₀ 0.9–3.4 μM (ref. 119)
Meglumine antimoniate	Leishmanicidal	n.d.	EC ₅₀ 29.1–60.1 $\mu\text{g ml}^{-1}$ (depending on strain) ¹²⁰

^a n.d. = not determined.



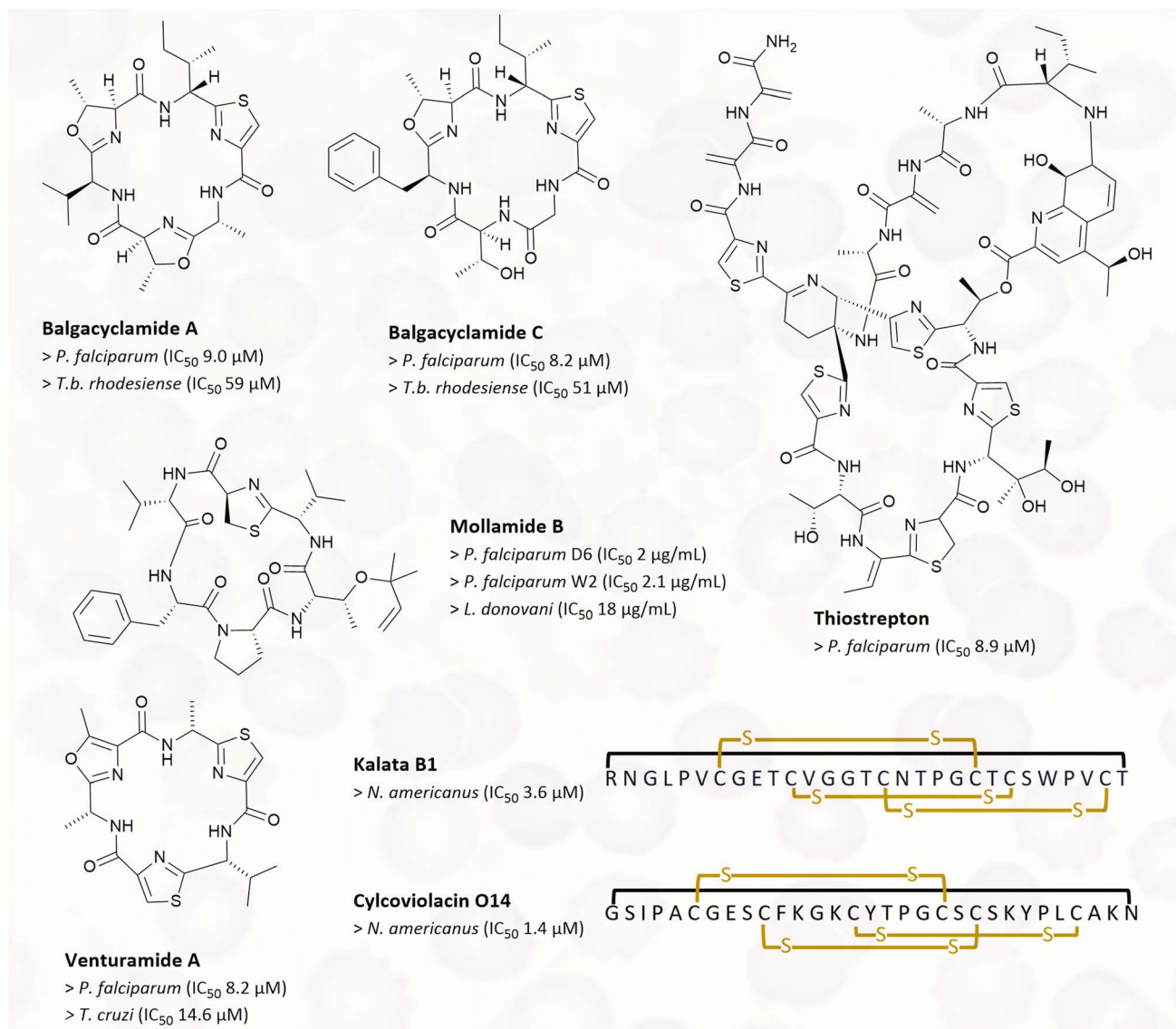


Fig. 6 RiPPs with antiparasitic properties: chemical structures of balgacyclamide A and C, thiostrepton, mollamide B, venturamide A and B, kalata B1, and cycloviolacin O14 are shown. For kalata B1 and cycloviolacin O14, the one-letter code is used. Yellow indicated crosslinks are disulphide bridges, the black link indicates a macrocyclisation. IC₅₀ values of affected parasites are listed below the respective compound names.

and death if left untreated. The best antiparasitic activity of balgacyclamides was detected against *P. falciparum* K1 with an IC₅₀ of 8.2–9 μM, depending on the derivative. Besides, moderate activity against *T.b. rhodesiense* with IC₅₀ values between 51–59 μM, and no activity against other parasites was reported.¹¹⁰ The cyanobactin mollamide B (Fig. 6), isolated from the tunicate *Didemnum molle*, was tested for its antiparasitic activity against the chloroquine-sensitive *P. falciparum* strain D6, the chloroquine-resistant *P. falciparum* strain W2, and *L. donovani*. It showed moderate activity with IC₅₀ values of 2 μg ml⁻¹, 2.1 μg ml⁻¹, and 18 μg ml⁻¹, respectively.¹¹¹ Other peptides isolated from cyanobacteria, venturamide A and B (Fig. 6), were also tested against tropical parasites. They were isolated from *Oscillatoria* sp., a marine cyanobacterium. In bioassays both compounds showed moderate activity against *P. falciparum* and *T. cruzi*.

Venturamide A and B showed IC₅₀ values of 8.2 μM and 5.2 μM for *P. falciparum*, and 14.6 μM and 15.8 μM for *T. cruzi*, respectively.¹¹²

Cyclotides are plant derived RiPPs, and some representatives of this family show anthelmintic activities. Kalata B1 was isolated from *Oldenlandia affinis*, and cycloviolacin O14 was isolated from *Viola odorata* (Fig. 6).^{113,114} Both peptides were screened for their anthelmintic properties against the human hookworm *Necator americanus*. Hookworm infections are a human health issue in the tropics and subtropics, and they are the leading cause of maternal and child morbidity. Infections in children can result in intellectual, cognitive and growth retardation.¹¹⁵ *In vitro* tests conducted on *N. americanus* larvae showed that kalata B1 has an IC₅₀ of 3.63 μM, while cycloviolacin O14 has an IC₅₀ of 1.40 μM.¹¹⁶





Table 5 Overview of RiPPs with anticancer bioactivities^a

Compound name	Bioactivity	Target (if known)	Activity metrics
6OTD	Proapoptotic, anti-proliferative	Telomerase	GI ₅₀ 21 nM (U251), 180 nM (SNB-78), 0.30 μ M (average of 39 human cancer cell lines) ¹⁵⁸
Amnosamides	Cytotoxic	Myosin	IC ₅₀ 320 μ M (HCT-116 cells) ¹⁴⁰
Amnosester A	Cytotoxic	n.d.	EC ₅₀ 56 μ M (MDA-MB-231), 21 μ M (SK-MEL-5), 140 μ M (SF-295), 15 μ M (NCI-H226), 17 μ M (OVCAR-3) ¹⁴⁶
Chaxapeptin	Inhibition of cell motility	n.d.	48% reduced motility speed at 50 μ M (ref. 162)
Comoramides and mayotamides	Cytotoxic	n.d.	IC ₅₀ 5–10 μ g ml ⁻¹ (A549, HT29, MEL-28) ¹³⁰
Defb14	Anticancer	n.d.	50 μ g per day (Lewis lung carcinoma) ¹⁵¹
Dendroamide A	Chemo-sensitisation	P-glycoprotein	0.6 μ M (MCF7-MDR) ¹⁶⁴
Felipeptin A1 and A2	Chemo-sensitisation	Downregulation of tumour suppressor Rb	Synergistic effects at 6.25 and 12.5 μ M ¹⁶⁶
Gymnopeptide A	Anticancer	n.d.	88.4 nM (HeLa), 66.4 nM (A431), 26.5 nM (MCF7)
Gymnopeptide B	Anticancer	n.d.	37.4 nM (MDA-MB-231), 18.0 nM (T47D) ¹⁵³
Hymenamide B	Cytotoxic	n.d.	42.5 nM (HeLa), 44.3 nM (A431), 18.5 nM (MCF7)
Kintamdin	Cytotoxic	n.d.	30.7 nM (MDA-MB-231), 14.0 nM (T47D) ¹⁵³
Lissoclinamide 7	Cytotoxic	n.d.	IC ₅₀ 3.5 μ g ml ⁻¹ (L1210)
Lymphostin	Cytotoxic	Metal chelating ¹⁶⁷	6.0 μ g ml ⁻¹ (epidermoid carcinoma KB) ⁷⁵
Microcin E492	Antitumour	mTOR inhibition	IC ₅₀ 0.6 (HTB-22), 2.4 μ M (CRL-11147), 12.0 μ M (HTB-38) ¹⁵²
Nisin A	Cytotoxic, proapoptotic	n.d.	IC ₅₀ 0.06 μ g ml ⁻¹ (T24), 0.04 μ g ml ⁻¹ (MRC5CV1) ¹²⁸
Nisin ZP	Antitumour	Induction of CHAC1 expression ¹³⁵	IC ₅₀ 1.7 nM (ref. 148)
Patellamide D	Chemo-sensitisation	Calpain	Induces apoptosis at \sim 20 μ g ml ⁻¹ in HeLa cells ¹³³
Patellin 6	Cytotoxic	n.d.	IC ₅₀ 600 μ M (SWA480) ¹³⁴
<i>P. vulgaris</i> defensins	Anticancer	n.d.	<i>In vivo</i> dosed at 200 mg kg ⁻¹ (UM-SCC-17B) ¹³⁵
Siomycin A	Anticancer, proapoptotic	n.d.	800 mg kg ⁻¹ (UM-SCC-17B) ¹³⁶
Sungsanpin	Inhibition of cell motility	Forkhead box M1 (ref. 168)	3.3 μ M (CEM/VLB ₁₀₀) ¹⁶⁵
Telomestatin	Proapoptotic, anti-proliferative	Induces expression of metalloproteinase 1 and 2 (ref. 160)	2 μ g ml ⁻¹ (P388, A549, HT29 and CV1) ¹²⁹
Thiostrepton	Anticancer, proapoptotic	Telomerase	4.1 μ M (HepG2), 8.3 μ M (MCF7) ¹⁵⁰
Ulleungdin	Inhibition of cell motility	Forkhead box M1	Inhibition <i>in vivo</i> shown at 20 μ M (ref. 124)
Wewakazole B	Anticancer	n.d.	53% reduced motility speed at 50 μ M (ref. 162)
Marketed (non-RiPP) anticancer drug			
5-Fluorouracil (5-FU)	Anticancer	Inhibition of thymidylate synthase ¹⁶⁹	GI ₅₀ 6.5 μ M (average of 39 human cancer cell lines) ¹⁵⁸
			IC ₅₀ 42.6 μ M (ref. 125)
			56% reduced motility speed at 50 μ M (ref. 162)
			IC ₅₀ 1.0 μ M (H460), IC ₅₀ 0.58 μ M (MCF7) ¹²⁷
			IC ₅₀ 45–5063 ng ml ⁻¹ (range across 19 cancer cell lines) ¹⁷⁰

^a n.d. = not determined, cancer cell type in brackets (if applicable).¹³⁵

2.5 Anticancer/antitumour RiPPs

Cancer is among the foremost causes of death around the globe, resulting in almost 10 million deaths in 2020.¹²¹ Consequently, there is a pressing need for improved clinical treatment options. Peptides and proteins are of particular interest, as they possess a significant advantage over small molecule drug candidates: they show a reduced probability of off-target interactions.¹²² In this section, we will first discuss thiostrepton and its analogues, followed by cyanobactins, bacteriocins, ammosamides and related pyrroloquinoline RiPPs, defensins, and RiPPs of an unknown class. The final section will discuss RiPP natural products that have no cytotoxic properties but inhibit lung cancer cell migration and RiPPs that have synergistic properties with chemotherapy drugs.

Siomycin A was originally described as an antibiotic similar in structure to thiostrepton isolated from *Streptomyces* sp. H-690.¹²³ In a study by Radhakrishnan *et al.* it was identified as a potential inhibitor of the transcription factor forkhead box M1 (FoxM1).^{124,125} Dysregulation of FoxM1 has been associated with lung cancer and basal cell breast carcinoma.¹²⁶ Siomycin A appears to be a negative regulator of FoxM1, which at least partly contributes to its anticancer and proapoptotic activities. It was demonstrated that siomycin A treatment induces apoptosis only in transformed lung fibroblasts in culture, leaving non-cancerous cells nearly unaffected.¹²⁴ Unsurprisingly, thiostrepton also has anticancer properties. It directly interacts with FoxM1 as shown in MCF7 breast cancer cell lines by Hegde *et al.*¹²⁵

Cyanobactins are better known for their antimicrobial activities, but some also display cytotoxicity against human cancer cell lines. For instance, wewakazole B (Fig. 8), isolated from *Moorea producens* has an IC_{50} of 1.0 μ M (Table 5) against human H460 lung and an IC_{50} of 0.58 μ M against human MCF7 breast cancer cells.¹²⁷ Lissoclinamides are a group of cyanobactins isolated from the ascidian *Lissoclinum patella*. Initial

cytotoxicity assays revealed lissoclinamide 7 to be the most potent compound, with an IC_{50} of 0.06 μ g ml^{-1} for urinary bladder carcinoma cells.¹²⁸ Another cyanobactin from *L. patella* is patellin 6. In cytotoxicity assays, an IC_{50} of 2 μ g ml^{-1} for P388 (leukaemia), A549 (lung carcinoma), HT29 (colon cancer) and CV1 (non-cancerous cells) cells was measured.¹²⁹ Further expanding the group of cytotoxic cyanobactins, hymenamamide B from *Hymeniacidon* sp. exhibited an IC_{50} of 3.2 and 6.0 μ g ml^{-1} against murine lymphoma L1210 cells and human epidermoid carcinoma KB cells, respectively.⁷⁵ Hymenamamide B also has antifungal properties (see Section 2.2 above). Similarly, several peptides from *Didemnum molle* symbionts, comoramides A and B, and mayotamides A and B (Fig. 5) show mild cytotoxicity against A549 lung cancer cells, HT29 colorectal cancer cells, and MEL-28 malignant melanoma cells.¹³⁰

Class I bacteriocins represent a group of RiPPs that were initially reported as antimicrobials, and were subsequently discovered to possess a range of properties, including anti-cancer effects.¹³¹ One example is microcin E492 (MccE492), which showed activity against SW620 human colorectal cancer. Cancer cell viability decreased in a dose-dependent manner when treated with 0, 30 and 60 μ g ml^{-1} MccE492 suspension in an *in vivo* model using zebrafish larvae. Additionally, MccE492 induces apoptosis in HeLa cells at about 20 μ g ml^{-1} (Table 5). It is produced by *Klebsiella pneumoniae* RYC492 and it is mainly active against Enterobacteriaceae.^{132,133} Similarly, nisin A, a bacteriocin well known for its antibacterial application in food preservation, also displays cytotoxic properties. It induces apoptosis in cells of colon cancer, breast cancer, and hepatic cancer, with an IC_{50} of around 600 μ M.¹³⁴ Furthermore nisin A was tested in a floor-of-mouth oral cancer xenograft mouse model. Treatment with 200 mg kg^{-1} per day of nisin A led to significant tumour volume decrease with no observed adverse effects.¹³⁵ Additionally Kamarajan *et al.*¹³⁶ tested nisin ZP, a natural occurring variant of nisin A. Both nisin A and nisin ZP are produced by *Lactococcus lactis*. Nisin ZP was tested in a similar oral cancer floor-of-mouth mouse model. It was found to decelerate tumour growth, inhibit cancer cell proliferation, prevent angiogenic processes, suppress orasphere formation and tumorigenesis *in vivo* at dosages up to 800 mg kg^{-1} per day. Converting this dose to human administration would entail giving 66.7 mg kg^{-1} to humans, which falls well within the no observed adverse effect level (NOAEL) range of 225 mg nisin A kg^{-1} bodyweight identified by the European Food and Safety Administration. It is theorized that nisin ZP activates calpain which then leads to apoptosis in cancerous cells (Fig. 7).^{136–138}

Ammosamides belong to the natural product class of pyrroloquinoline alkaloids, and their biosynthetic pathway contains cryptic RiPP genes.¹³⁹ Ammosamides were initially isolated from the *Streptomyces* strain CNR-698 and are compounds with intriguing structures and bright colours, but with low solubility.¹⁴⁰ Though they were originally thought to constitute a 16-member family, it has been shown that most ammosamides are artefacts of ammosamide C, forming derivatives when exposed to nucleophiles, air, and light.^{140,141} Ammosamides A and B (Fig. 8) display *in vitro* cytotoxicity towards the human colorectal cancer cell line HCT-116, with an

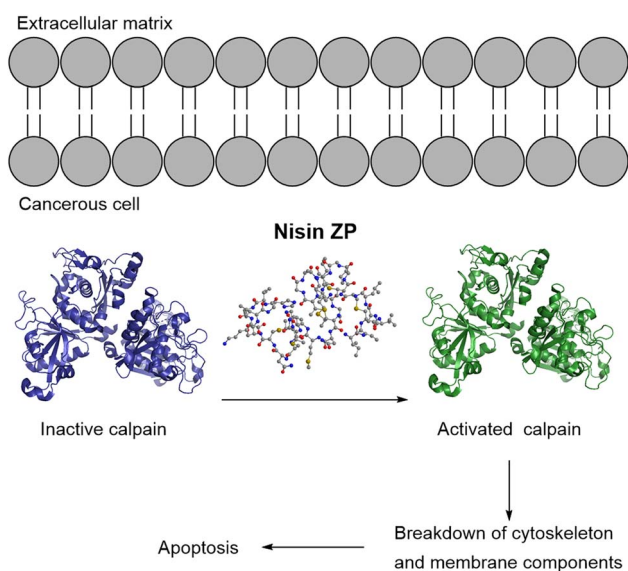


Fig. 7 Nisin ZP activates calpain in cancerous cells; this activation leads to the hydrolysis of different cytosolic and membrane proteins and induces apoptosis. Schematic is based on ref. 136 and 138.



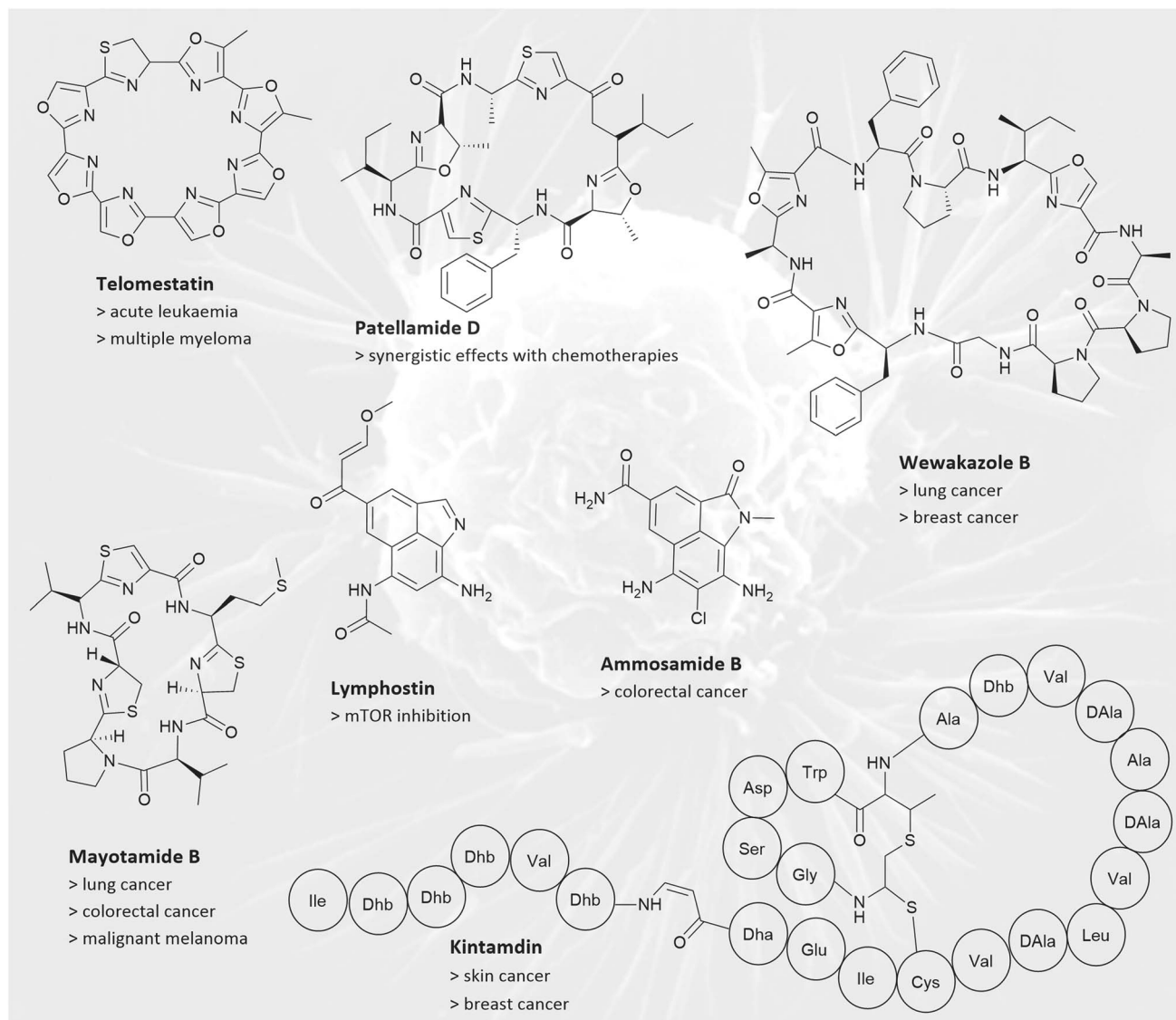


Fig. 8 RiPPs with anticancer activities: chemical structures of ammosamide B, kintamadin, lymphostin, wewakazole B, telomestatin, patellamide D, and mayotamide B are shown. Examples for affected cancer cell lines are given below the compound names. For kintamadin, the three-letter code is used.

IC₅₀ of 320 nM. Myosin appears to be the intracellular target.¹⁴² Ammosamide B derivatives demonstrate potent quinone reductase 2 (QR2) inhibition. QR2 has been identified as a potential target for the development of chemotherapeutic drugs.¹⁴³ Research by Li *et al.* indicates that ammosamides can be a highly effective scaffold for the development of targeted therapies.¹⁴⁴ Several derivatives were synthesised using the pyrroloquinoline base to optimise inhibition of BRD4. BRD4 is a transcriptional and epigenetic regulator, well-established as a target in cancer research.¹⁴⁵ Derivative 49 showed the highest inhibition against BRD4, when screened against MV4-11 and 22RV1, leukaemia and prostate cancer cell lines. Additionally, it also showed good anti-proliferative effects. Consequently, this compound displays potential as a new drug lead for chemotherapeutics.¹⁴⁴ The ammosester subfamily of ammosamides was discovered in the genome of *Streptomyces unicalis* DCA2648 and exhibits modest cytotoxicity against several human cancer

cell lines, including melanoma (SK-MEL-5), breast (MDA-MB-231), central nervous system (SF-295), non-small cell lung (NCI-H226), and ovarian (OVCAR-3) cancer.¹⁴⁶

Additional members of the pyrroloquinoline natural product family are the lymphostins (Fig. 8). Like ammosamides, they possess a non-canonical biosynthesis, involving genes associated with RiPP biosynthesis, the exact function of which require further investigation.¹³⁹ Originally isolated from *Streptomyces* sp. KY11783, lymphostins and their derivatives are potent inhibitors of mTOR (mechanistic Target of Rapamycin), exhibiting IC₅₀ values ranging between 0.8 and 1.8 nM.^{147,148} The serine/threonine kinase mTOR is involved in regulating cell survival, cell growth, cell metabolism, protein synthesis, and autophagy. As shown in animal models and clinical cancer patients, dysfunction of mTOR contributes to tumorigenesis.¹⁴⁹

Some defensins (see Section 2.4) show anticancer activity both *in vitro* and *in vivo*. For instance, a defensin from *Phaseolus*



vulgaris cv. “extra-long purple pole bean” displays potent antiproliferative properties against tumour cells. The IC₅₀ values for hepatoma (HepG2) cells and breast cancer (MCF7) cells are 4.1 μ M and 8.3 μ M, respectively.¹⁵⁰ Additionally, Defb14, a mouse defensin, was shown to restrain the growth of Lewis lung carcinoma in a mouse model without causing any observable damage to the surrounding tissue.¹⁵¹

Kintamdin is a novel, macrocyclic RiPP that does not fall into any known RiPP category (Fig. 8). Kintamdin was isolated from *Streptomyces* sp. RK44 and shows cytotoxic activity against skin and breast cancer cell lines, with IC₅₀ values of 2.4 μ M and 0.6 μ M respectively.¹⁵² Other macrocyclic peptides of an unknown class were obtained from the mushroom *Gymnopus fusipes*, the cyclic gymnopeptides A and B. They show antiproliferative properties against cervical, epidermoid, and breast cancer cell lines with IC₅₀ values ranging from 10–90 nM.¹⁵³ Telomestatin (Fig. 8), a macrocyclic RiPP isolated from *Streptomyces anulatus*, has demonstrated great potential as a chemotherapeutic agent.^{154,155} It interacts specifically with telomerase, resulting in proapoptotic and antiproliferative effects in acute leukaemia and multiple myeloma.^{156,157} Telomestatin has more than one mode of action, making it an ideal candidate for further development of anticancer agents. It causes telomere dysfunction, downregulates the proto-oncogene *c-Myc*, and induces a higher level of replication stress response in cancer cells compared to non-cancerous cells. The telomestatin analogue 6OTD shows activity in lower concentrations compared to the native peptide when exerting its anticancer effects on a human cancer cell line panel. On average 6OTD had an GI₅₀ value of 0.30 μ M for 39 cell lines, while telomestatin had an average GI₅₀ of 6.5 μ M.¹⁵⁸ Additionally, 6OTD showed potent antitumour activity against human glioblastoma U251 cells, with treated tumours only reaching 33% of the non-treated tumour sizes in an *in vivo* mouse model. Other analogues are under investigation.^{158,159}

Another approach to cancer treatment involves the use of drugs that inhibit cancer cell motility. Although not cytotoxic in nature, these compounds can help reduce the risk of cancer metastasis in patients. One group of compounds that can achieve this are certain lasso peptides, which have closely related sequences and structures. This cluster of lasso peptides is made up of sungsanpin, chaxapeptin, and ulleungdin. These compounds were isolated from *Streptomyces* sp. SNJ013, *Streptomyces leeuwenhoekii* C58 and *Streptomyces* sp. KCB13f003, respectively. Sungsanpin was first discovered in a deep-sea sediment sample, while chaxapeptin was found in the Atacama desert, and ulleungdin on a volcanic island.^{160–162} Despite their different origins, these natural products exhibit similar bioactivity. At a concentration of 50 μ M, all three compounds inhibit cell migration of A549 lung cancer cells by approximately 50% compared to untreated cells.^{160–162} These three RiPPs have the potential to inspire a derivative compound for clinical applications. Digal *et al.* have already demonstrated that the knot motif is not necessary for retaining function. Macrocyclic and head-to-side chain derivatives of sungsanpin and ulleungdin have retained comparable bioactivity.¹⁶³

In addition to classical cytotoxic antitumour medications, pharmaceuticals that work in synergy with chemotherapy drugs offer a promising approach, particularly for targeting multidrug-resistant tumours. One such compound is dendroamide A, which was isolated from *Stigonema dendroideum* and has been shown to increase the accumulation of vinblastine, a chemotherapy drug, in P-glycoprotein-overexpressing breast carcinoma cells (MCF7/ADR).¹⁶⁴ Similarly, patellamide D (Fig. 8) from the marine tunicate *Lissaculum patella*, reduced IC₅₀ values for the cytotoxins vinblastine, colchicine, and adriamycin in CEM/VLB₁₀₀ human leukaemic cells, reversing the cell line's multi drug resistance.¹⁶⁵ The findings suggest potential for the use of patellamide D as a therapeutic agent for drug-resistant cancers. Felipeptin A1 and A2 are lasso peptides isolated from *Amycolatopsis* sp. YIM10. They have been shown to promote cell proliferation in cancer cell lines MCF7, HCT-116 and A375 (malignant melanoma). When applied individually, only marginal effects were observed. However, when both feli-peptin A1 and A2 were combined, a synergistic pro-proliferation effect was observed in cell viability assays. Pre-treating A375 cell lines with 6.25 or 12.5 μ M of each feli-peptin increased the sensitivity to the cytotoxic chemotherapy drug doxorubicin. Pre-treating doxorubicin-resistant MCF7 cancer cells with feli-peptins A1 and A2 re-sensitised them to the chemotherapeutic drug. This study demonstrates that feli-peptin A1 and A2 have potential in future chemotherapy applications to reduce dosage and overcome resistance in clinical settings.¹⁶⁶

2.6 Analgesic RiPPs

Chronic pain is an immense burden on people and public health providers worldwide; according to the US Center for Disease Control and Prevention point prevalence is at 20.9% for chronic pain in the United States.¹⁷¹ In contrast to acute pain, chronic pain has little evolutionary benefit and becomes a disease in its own right. Pain management is therefore of the utmost importance for patients to mitigate not only the pain itself but other biological, social, and psychological factors that occur alongside this condition.¹⁷² There are several RiPPs with analgesic properties. This section will begin by exploring two lanthipeptides, followed by an examination of conotoxins and finally, other venom peptides that possess analgesic properties. Conotoxins are peptides isolated from the cone snails venom, and are used by the snails for immobilising prey, which can later be swallowed whole.¹⁷³

The lanthipeptide labyrinthopeptin A2 (LabyA2, see Section 2.3 above) was first isolated because of its antiviral activity in culture extracts of *Actinomadura namibiensis* DSM 6313, further investigation was prompted by its unusual post-translational modifications. LabyA2 (Fig. 6) contains labionin, an unprecedented non-canonical amino acid. LabyA2 showed *in vivo* efficacy in a mouse model of neuropathic pain. In a spared nerve injury mouse model, intravenous administration of LabyA2 led to significant attenuation of tactile allodynia over a 6 h observation period (Table 6).¹⁷⁴ Iorio *et al.* identified another lanthipeptide that contains labionin in *Actinoplanes* sp. DSM14059, known as NAI-112. In addition to labionin, NAI-112 has another



Table 6 Overview of RiPPs with analgesic properties^a

Compound name	Bioactivity	Target (if known)	Activity metrics
Crotalphine	Antinociceptive	κ and δ opioid receptors	Dosed at 1 $\mu\text{g kg}^{-1}$ (ref. 183)
CVIE and CVIF	Antiallodynic	Neuronal-type Ca^{2+} channels	Dosed at 1 nM (ref. 184)
Eu1.6	Analgesic	Neuronal-type Ca^{2+} channels	IC_{50} 1.1 nM (ref. 103)
Labyrinthopeptin A2	Antiallodynic	n.d.	ED_{50} 50 $\mu\text{g kg}^{-1}$ (ref. 174)
NAI-112	Antinociceptive	n.d.	Dosed at 30 mg kg^{-1} (ref. 175)
Psalmotoxin 1	Analgesic	ASIC1a	Dosed at 0.1 nmol per mouse ¹⁸¹
Tx3–5	Antinociceptive	n.d.	ID_{50} 16.6 fmol per site, max inhibition dose of 30 fmol per site ¹⁸²
Marketed (RiPP) analgesic drug			
Ziconotide	Antiallodynic	Neuronal-type Ca^{2+} channels	IC_{50} 2–29 pmol (binding to Ca^{2+} channel), <i>in vivo</i> models: ID_{50} 3–30 pmol (acute pain), ID_{50} 30–1000 ng (chronic pain) ¹⁸⁵

^a n.d. = not determined.

unusual modification: the indole nitrogen of Trp13 carries a 6-deoxyhexose moiety. This is the first example of an *N*-glycosylated lanthipeptide and the first natural product in which a tryptophan residue is *N*-glycosylated. After observing similarities with LabyA2, the researchers tested for antinociceptive activity. In these assays, NAI-112 proved to be effective in alleviating acute pain induced by formalin injections, by reducing hyperalgesia and allodynia in a dose-dependent manner. It also demonstrated efficacy on established chronic pains in a mouse model with chronic constriction injury, with full effect at a dose of 30 mg kg^{-1} of NAI-112. Furthermore, there was no indication of NAI-112 affecting motor coordination up to 60 minutes after administration and no signs of toxicity were observed.¹⁷⁵

The most prominent group of analgesic peptide drug leads comprise conotoxins. One example is the α -conopeptide Eu1.6

from *Conus eburneus* (Fig. 10), which shows significant analgesic activity in neuropathic pain models at low doses, surpassing the positive control of morphine and gabapentin. Eu1.6 represents the first conopeptide showing this effect with intravenous administration in contrast to intrathecal injections, which are injected into the spinal canal. Additionally, no significant impact on the motor, cardiac or respiratory function of mice was observed even at a dose that was 100 times higher than the effective dose. This result renders Eu1.6 a promising lead structure in treating neuropathic pain.¹⁷⁶

The ω -conotoxins CVIE and CVIF (Fig. 10) were isolated from *Conus catus* venom, and they are selective inhibitors of neuronal-type Ca^{2+} channel (Fig. 9). In a rat model of persistent pain, intrathecal administration of 1 nM led to a significant reversal of mechanical allodynia to the pre-injury baseline levels. Concurrently side effects of shakes, tail twitching, and serpentine tail movements were recorded.¹⁷⁷ Those side effects are typical, and they do not necessarily represent a hurdle in developing these compounds into pharmaceuticals. Ziconotide, a synthetic derivative of the conotoxin MVIIA also belongs to the ω -conotoxin family and is the first RiPP drug to be approved by the FDA (see Section 3.2 below).¹⁷⁸

The group of venom ribosomal peptide natural product is vast, with many valuable peptides found not only in cone snail venom but also spider venoms. Some examples are described below; for a more in-depth review see Wu *et al.*¹⁷⁹ Psalmotoxin 1 (PcTx1, Fig. 10) was isolated from the South American tarantula *Psalmopoeus cambridgei*. It is non-lethal and blocks ASIC1a neurons that are associated with a variety of pain sensations.¹⁸⁰ In different rodent models, PcTx1 showed similar pain relief as morphine, for both acute and neuropathic pain. In cases of chronic constriction injury (CCI) of the sciatic nerve in rats, thermal hyperalgesia and tactile allodynia were reversed by PcTx1. The peptide needs to be injected intrathecally or intracerebroventricularly, as intraperitoneal or subcutaneous injections had no effect.¹⁸¹ Another example is the peptide Tx3–5 (Fig. 10) from *Phoneutria nigriventer* venom. It has been proven that intrathecal injection of Tx3–5 in different mouse models can

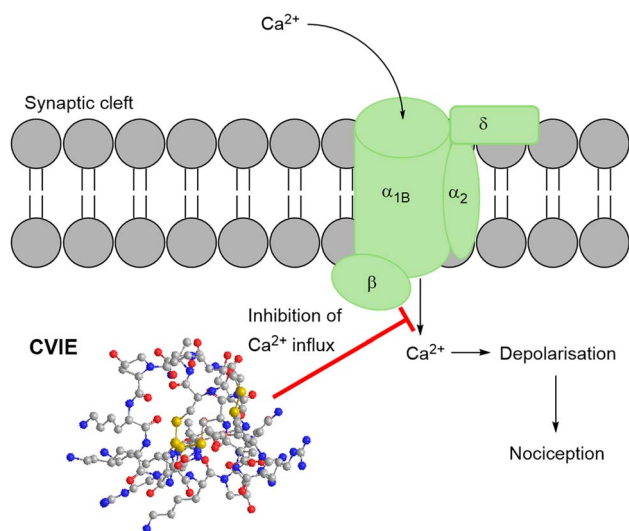


Fig. 9 The conotoxin CVIE selectively blocks N-type Cav2.2 calcium channels, thereby inhibiting nociception.



prevent or reverse postoperative nociception, show partial antinociceptive effects in a neuropathic pain model, and can nearly abolish cancer related nociception. However, it did not change noxious heat sensitivity or mechanical threshold, implying no effects on physiological pain. Additionally, no visible adverse effects were observed at the administered dose, with a TD_{50} about 50-fold higher than maximum inhibition dose.¹⁸²

Crotalphine (Fig. 10), a venom-derived peptide obtained from the South American rattlesnake *Crotalus durissus terrificus*, also exhibited antinociceptive properties. The oral administration of crotalphine effectively blocked hyperalgesia induced by PGE_2 , an inflammatory agent. The peptide demonstrated a dose-dependent, antinociceptive effect over a course of 5 days. Injecting crotalphine intravenously and intraplantarly (into the sole of the foot) also showed long lasting antinociceptive effects, both systematically

and locally, for PGE_2 -induced hyperalgesia. The effects are most likely mediated by activation of κ -opioid receptors. Oral availability and long lasting effects indicate the therapeutic potential of crotalphine and its derivatives for treating chronic pain.¹⁸³

2.7 Further bioactivities

Newly discovered natural products are usually tested for the most common or most urgently needed bioactivities, such as antibiotic, antiviral, antifungal, and anticancer activities. However, there are some other interesting bioactivities such as antiinflammatory, antidiabetic, antihypertensive, and anti-parkinsonian effects. Examples of RiPPs showing these bioactivities are described for different RiPP families, *i.e.*, lanthipeptides, lasso peptides, and cyanobactins.

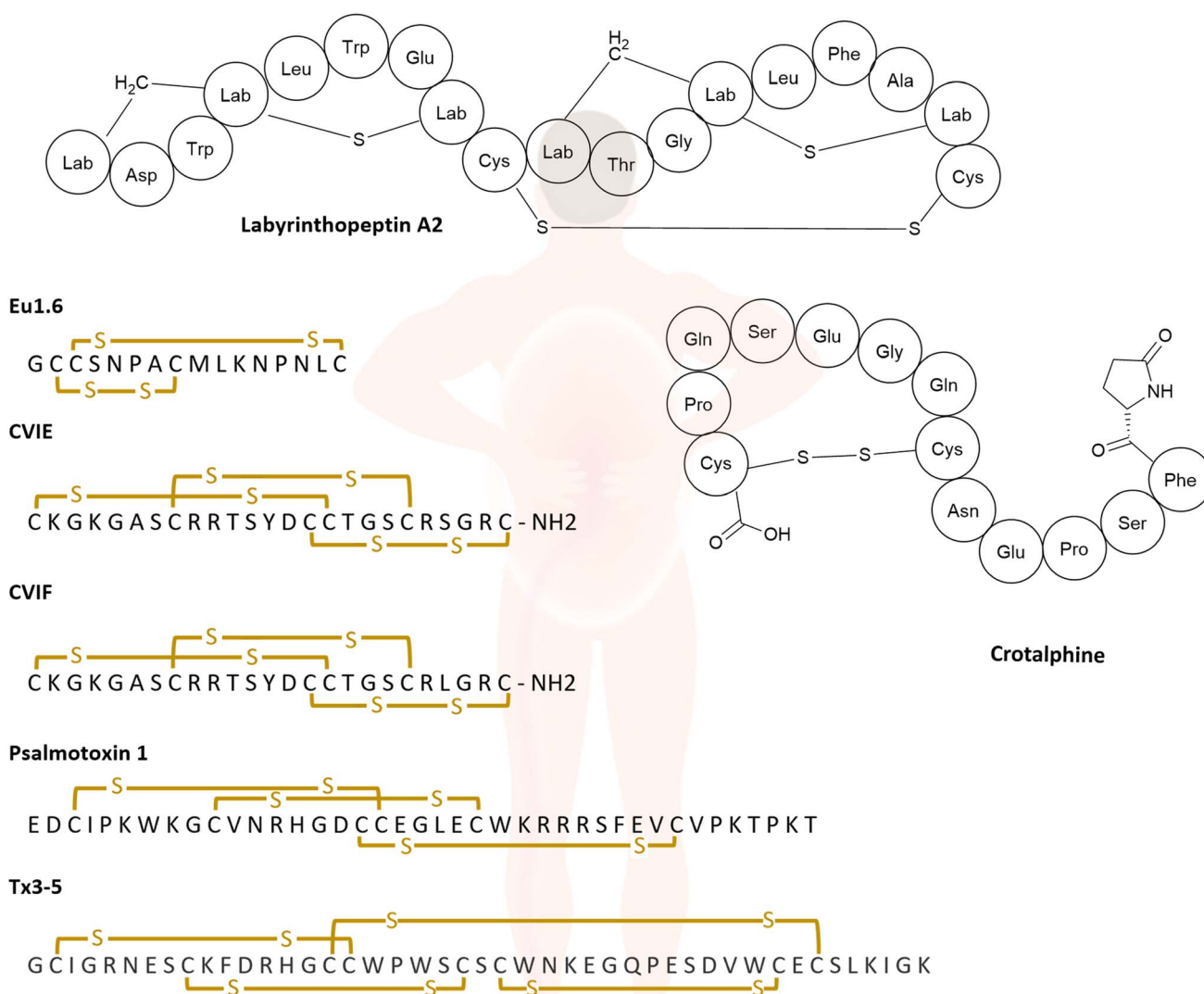


Fig. 10 Examples for RiPPs with analgesic activities. Structures of labyrinthopeptin A2, crotalphine, Eu1.6, CVIE, CVIF, psalmotoxin 1, and Tx3–5 are depicted. For Eu1.6, CVIE, CVIF, psalmotoxin 1, and Tx3–5, the one-letter code is used. Yellow indicated crosslinks are disulfide bridges.



The recently discovered lanthipeptides myxococin A and B from *Myxococcus fulvus* show antiinflammatory effects in lipopolysaccharide-induced mouse macrophages without any detectable cytotoxicity, which makes them interesting drug candidates (Table 7).¹⁸⁶ Ancovenin is a cinnamycin derivative isolated from *Streptomyces* sp. No. A647P-2 and belongs to the lanthipeptide subfamily as well. It shows antihypertensive activity by inhibiting the angiotensin 1-converting enzyme (ACE) with an IC_{50} of 0.87 μ M and was named after this activity (angiotensin converting enzyme inhibitor). ACE inhibition is clinically used for the treatment of high blood pressure.^{187–189}

In the class II of lasso peptides, two receptor antagonists are found: Anantin and RES-701-1. Anantin, a peptide isolated from *Streptomyces coeruleus*, binds to the atrial natriuretic peptide (ANP) receptor. ANP is the natural agonist of this membrane receptor, and it is involved in blood pressure homeostasis; Anantin is the first microbially produced antagonist of ANP.^{190,191} RES-701-1 was isolated from *Streptomyces* sp. RE-701 and binds strongly to the type B endothelin receptor with an IC_{50} of 10 nM. Endothelins are a group of potent vasoactive peptides and high endothelin levels are found in several pathological conditions such as systemic hypertension, cardiac ischemia, and asthmatic attacks.¹⁹² The endothelin type B receptor mediates vasoconstriction and vasodilatation, hence it is involved in blood pressure regulation. Several RES-701-1-related compounds have been isolated: the derivative RES-701-2 from the RES-701-1 producer and RES-701-3 and RES-701-4 from *Streptomyces* sp. RE-896. The derivatives differ in two positions; 1 and 2 contain an alanine in position 7, while 3 and 4 contain a serine in this position, and 1 and 3 contain a tryptophan in position 16, whereas 2 and 4 contain 7-hydroxytryptophan.¹⁹³ The biosynthetic gene cluster of RES-701-3 and -4 has recently been identified in the marine bacterium *Streptomyces caniferus* CA-271066.¹⁹⁴ RES-701-1 and its derivatives are selective endothelin antagonists with RES-701-3 > -1 > -2 > -4 (order of potency).¹⁹³ Moreover, synthetic derivatives of RES-701-1 have been developed: a hybrid peptide chemically synthesised from the RES-701-1 N-terminus and the endothelin-1 C-terminus exhibited an IC_{50} of 0.24 nM.¹⁹⁵ Further derivatives also showed antagonistic activity for the endothelin type A receptor.^{196,197} These results nicely exemplify that natural products are promising lead structures for drug development.

The compound BI-32169 belongs to class III of the lasso peptide subfamily.¹⁹⁸ BI-32169 is produced by *Streptomyces* sp. (DSM 14996)

and inhibits the human glucagon receptor with an IC_{50} of 440 nM. Its methyl ester derivative exhibits slightly more potent inhibitory activity with an IC_{50} of 320 nM.^{199,200} Glucagon antagonists are considered promising candidates for new antidiabetic therapies.²⁰¹

The macrocyclic peptide agardhiheptin A was isolated from the cyanobacterium *Oscillatoria agardhii* (NIES-204) and belongs to the RiPP family of cyanobactins. Agardhiheptin A is a weak plasmin inhibitor (IC_{50} of 65 μ g ml^{-1}); the protease plasmin is involved in the regulation of fibrinolysis and plasmin inhibitors are clinically used to treat hyperfibrinolysis-associated bleeding events.^{202,203}

The recently discovered fungal RiPP natural product acalitinide is a macrotricyclic compound with two disulphide bridges; disulphide bridges are a common motif in marketed peptide drugs. Acalitinide was isolated from the ascomycete *Acaulium album* and shows neuroprotective effects. It is a promising drug candidate to treat the Parkinson's disease.²⁰⁴

2.8 Imaging and diagnostic agents

Therapeutic applications not only encompass medication but also diagnostics such as radiocontrast agents, medical probes, or microscopic dyes. A few RiPPs, mainly analogues of peptide hormones, are used for such applications.

Phalloidin is a macrocyclic RiPP of the phalloxin group produced by the fungus *Amanita phalloides*.²⁰⁵ Functionalised with a fluorophore, phalloidin derivatives are commonly used as a stain in immunofluorescence microscopy because of their high affinity for actin filaments in cells and tissues.²⁰⁶

Bombesin is a peptide hormone isolated from the skin of the toad *Bombina bombina*.²⁰⁷ It binds with high affinity to receptors of certain cancer cells, *i.e.* gastrin-releasing peptide receptor (GRPR)-positive tumours such as human prostate and breast tumours, as well as small-cell lung, ovarian, and endometrial cancer. Fluorescently labelled bombesin derivatives have been investigated for the application in peptide receptor imaging.^{208–210} Later on, bombesin analogues for targeted tumour therapy approaches were developed, since they might be used as drug shuttles for the intracellular delivery of different cytotoxic compounds.²¹¹

Several other compounds derived from peptide hormones can be used in radiopharmaceutical applications. Examples include somatostatin analogues for the diagnosis of neuroendocrine tumours,^{212,213} glucagon-like peptide-1 analogues for imaging of beta cell function in diabetes patients,^{214,215} and

Table 7 Overview of RiPPs with non-canonical bioactivities^a

Compound name	Bioactivity	Target (if known)	Activity metrics
Acalitinide	Neuroprotective/anti-parkinsonian	Dopamine neurons ²⁰⁴	n.d.
Agardhiheptin A	Regulation of fibrinolysis	Plasmin	IC_{50} 65 μ g ml^{-1} (ref. 202)
Anantin	Antihypertensive	Atrial natriuretic peptide receptor	IC_{50} 1.0 μ M (ref. 190)
Ancovenin	Antihypertensive	Angiotensin converting enzyme	IC_{50} 0.87 μ M (ref. 188)
BI-32169	Antidiabetic	Glucagon receptor	IC_{50} 440 nM (ref. 199)
BI-32169-CH ₃	Antidiabetic	Glucagon receptor	IC_{50} 320 nM (ref. 199)
Myxococin A + B	Antiinflammatory ¹⁸⁶	n.d.	n.d.
RES-701-1	Antihypertensive	Type B endothelin receptor	IC_{50} 10 nM (ref. 192)

^a n.d. = not determined.



cholecystokinin analogues for the visualisation of medullary thyroid carcinomas, small-cell lung cancers, and stromal ovarian cancers.^{216,217}

2.9 Synthetic, engineered RiPP natural products

The previous sections have shown that RiPPs exhibit a broad range of bioactivities and are interesting drug candidates for treating a variety of diseases. In addition, the RiPP biosynthesis represents a possible biotechnological production route, and it is particularly well suited for engineering approaches to produce customised peptide products for pharmaceutical applications. For more comprehensive reviews on RiPP engineering, the reader is referred to ref. 17–19.

Several bioengineered RiPPs with new activities have been identified in different screening approaches. The protein–protein interaction (PPI) of the HIV p6 protein with the human TSG101 protein at its UEV domain is crucial for viral cell exit. Aiming to find an inhibitor of this PPI, a lanthipeptide library with the prochlorosin leader peptide and diversified core peptides was constructed. To generate bicyclic peptides, the promiscuous synthetase ProcM from *Prochlorococcus* was employed. One selective inhibitor, the peptide variant XY3-3 (Fig. 3), was obtained. Binding to UEV at a distinct site to p6 protein was proven. Interaction of p6-UEV was shown to be disrupted with an IC₅₀ of 3.6 μM. Moreover, the requirement of both thioether rings for binding and thus for the activity of XY3-3 was demonstrated.²¹⁸ XY3-3 is an interesting antiviral drug candidate. Further examples for bioengineered RiPPs are compounds containing the ‘RGD’ binding epitope. The human integrin α_vβ₃ is a potential drug target that recognises the ‘RGD’ motif. Compounds targeting the α_vβ₃ integrin receptor may be used as antiangiogenics in cancer diagnostics and therapy. By precursor engineering, this ‘RGD’ motif has been introduced into many different RiPPs, including lanthipeptides,^{219,220} lasso peptides,^{221–223} knottins,^{224,225} and cyclotides (θ-defensins).²²⁶

In general, RiPP pathways are relatively short and streamlined; the biosynthetic gene clusters consist of separate precursor peptide(s) and post-translationally acting enzymes that are often promiscuous. This modularity facilitates the gene cluster manipulation and allows to mix and match enzymes from different pathways to generate tailored designer peptides. The promiscuity of the maturases is founded on the separation of the substrate recognition site and the modification site; this spatial segregation allows the modification of various core sequences. Furthermore, the gene-encoded precursor peptides can be modified easily by simple mutagenesis and enable the rapid generation of novel peptide variants.¹⁹

Even though combinatorial biosynthesis of RiPP pathways is theoretically easy, the combination of enzymes from different pathways can be challenging. Many RiPP maturases contain so-called RiPP recognition elements (RRE) to recognise and bind the leader part of their cognate precursor peptides.²²⁷ Consequently, they do not accept precursor peptides with different or without leader peptides. For such naturally incompatible maturase-precursor combinations, where maturases are inactive towards designed core peptides, various engineering

strategies have been described, see ref. 18 for a comprehensive overview. Here, we will only briefly mention the most common strategies. The leader peptide part can be engineered to promote correct protein–protein interactions between the leader peptide and the maturase, e.g. rationally designed chimeric leader peptides harbouring RREs for different maturases can be employed.²²⁸ However, this approach is quite time-consuming, since the leader parts required for maturase activity need to be experimentally determined for every single maturase. In a second step, the chimeric leader needs to be produced and tested. Leader peptide complementation could be a simpler strategy. The required leader peptide can be supplied either *in cis*, meaning it is covalently linked to the maturase, or *in trans*, meaning it is expressed as a separate peptide molecule.^{229–232} However, this strategy is not generally applicable to all types of RiPP systems, but it works for specific RiPPs. Another option would be a sortase-based leader-peptide exchange approach,²³³ in which a sortase A recognition site is introduced between the leader peptide and the core peptide. The sortase exchanges the leader peptide of one maturase with the leader peptide of a second or third maturase. To incorporate unnatural amino acids that cannot be introduced by post-translational modifications of proteinogenic amino acids, the flexizyme technology can be applied.^{234,235} This technology enables the reprogramming of codons to accommodate unnatural amino acids using aptamers to charge tRNAs *in vitro*.

In summary, several options to circumvent compatibility issues have been described. The bioengineering of RiPPs is still in its early stages, and several obstacles must be overcome before RiPP processing enzymes from various subfamilies can be effectively combined for targeted peptide engineering. The RiPP technology represents a great chance for drug development by offering opportunities to generate a broad range of drug candidate derivatives in peptide libraries. Moreover, these engineered pathways could at the same time be suitable to produce peptide drugs at a large scale, superseding multi-step total synthesis of these compounds.

3 Advanced RiPP (candidate) drugs

3.1 RiPPs in clinical trials

Although many RiPPs have been discovered exhibiting pharmaceutically interesting bioactivities, only few entered clinical trials so far. Data about clinical trials is hard to access since most of it is not published in the peer-reviewed literature. Therefore, the following enumeration is not comprehensive, but contains the candidates we were able to identify with our means.

Cystic fibrosis is caused by a defective or missing cystic fibrosis transmembrane conductance regulator (CFTR) anion channel. Lancovutide (or Moli1901 or duramycin) was demonstrated to activate chloride channels, thereby having the potential to compensate for the dysfunctional CFTR. It entered phase IIb clinical trial for treatment of cystic fibrosis in 2007. Since the administration of lancovutide compared to placebo showed no significant positive effect on cystic fibrosis patients, no further clinical investigations were carried out to our knowledge.^{236,237}



Two semisynthetic derivatives of thiopeptide GE2270A, LFF571 and CB-06-01 (or NAI-003), completed both phase II clinical trials. Novartis Pharmaceuticals developed and tested LFF571 successfully against moderate *C. difficile* infections. Knowledge about the target of LFF571 is not published. By terminating the antibacterial research in 2018, the LFF571 project was stopped by Novartis.²³⁸

The company Cassiopea SpA made a proof-of-concept investigation of CB-06-01 for treatment of moderate-to-severe acne. CB-06-01 assumably binds the elongation factor Tu in *Propionibacterium acnes*, thereby abolishing protein biosynthesis. Derivatisation of GE2270A to CB-06-01 reduced the spectrum of activity but led to a lower minimal inhibitory concentration against *P. acnes*. Phase II dose-ranging studies of CB-06-01 were planned to be performed in the future according to Cassiopea SpA.^{239–241}

There are multiple conotoxins known that possess clinically relevant properties. Conotoxin Vc1.1 was isolated from *Conus victoriae* and discovered to act as a nicotinic acetylcholine receptor antagonist (nAChR). The synthetic version of Vc1.1, named ACV1, was found to be a potential therapeutic for the treatment of diabetic neuropathy, post herpetic neuralgia, and sciatic neuropathic pain.²⁴² Metabolic Pharmaceuticals Ltd tested ACV1 for treatment of neuropathic pain in a phase I clinical trial, aiming to assess the safety, tolerability, pharmacokinetics and -dynamics.²⁴³ Furthermore, they examined the application of ACV1 as medication for diabetic peripheral neuropathy and post-herpetic neuralgia in a phase II study.²⁴⁴ Due to remarkably lower efficiency of ACV1 in humans compared to rats and thus immensely higher doses necessary in human treatment, Metabolic Pharmaceuticals Ltd eventually terminated its ACV1 clinical research project.²⁴⁵

MrIA and MrIB, two closely related conopeptides produced by *Conus marmoreus*, founded the class of χ -conopeptides. They allosterically target the neuronal norepinephrine transporter (NET).²⁴⁶ To encounter the instability of natural MrIA peptide, Xenome Ltd synthesised a derivate of MrIA, Xen2174. Considering the enhanced stability, duration of analgesia and an equivalent efficacy vs. side effects window of Xen2174, clinical studies have been conducted.^{247–249} Two different clinical trials with healthy subjects were carried out under Xenome Ltd, investigating its therapeutic potential in pain management. Besides evaluating safety, tolerability, pharmacokinetics and -dynamics, pain modalities affected by Xen2174 and dose-dependent effects were investigated. The outcome was positively evaluated in both studies.^{250,251} However, a third trial comparing administration of Xen2174 alone and in interplay of Xen2174 with the well-established local anaesthetic bupivacaine, was rated negative after completion.^{252,253} Several other conopeptides are described as currently being in or having been in (pre-)clinical trials.²⁵⁴

3.2 RiPPs on the market

In the previous chapters, we presented an overview of potential drug candidates in the clinical trial pipeline of RiPPs. In this chapter, we will review those RiPPs that have been approved for a variety of indications.

Ziconotide (Fig. 11), known as Prialt®, is perhaps the most prominent example of a RiPP drug. It is a synthetic form of the ω -conotoxin MVIIA, which was originally discovered in *Conus magus*, and subsequently reported on by Olivera *et al.* in 1987.²⁵⁵ Ziconotide binds to N-type calcium channels in the spinal cord, and it is the only non-opioid analgesic approved for the treatment of chronic pain in Europe and the USA. It is administered intrathecally and recommended as first line treatment for neuropathic and nociceptive pain. Although adverse side effects continue to pose a concern, ziconotide has significant benefits, including the absence of bone marrow toxicity and respiratory depression, as well as displaying no signs of withdrawal symptoms post-treatment.²⁵⁶ Presently, investigation is underway to optimise treatment plans and mitigate adverse side effects (<http://ClinicalTrials.gov> Identifier NCT04321408), which underlines the importance of this drug.

Eptifibatide (Fig. 11) has been approved by both by the FDA and EMA for treating acute coronary syndrome since the late 1990s. It is derived from barbourin, a venom peptide found in the southeastern pygmy rattlesnake, *Sistrurus miliarius barbourin*.²⁵⁷ Eptifibatide is a cyclic heptapeptide and contains a modified version of the common 'KGD motif'. The 'RGD motif' allows it to bind to glycoprotein IIb/IIIa, the target protein. Additionally, its tertiary structure is essential to the specificity. Glycoprotein IIb/IIIa constitutes the final common pathway accountable for platelet aggregation. Inhibiting this pathway confers eptifibatide with its antithrombotic effect. More recently other possible indications have been investigated, *i.e.*, the treatment of ischemic stroke, carotid stenting, stenting of intracranial aneurysms, and septic shock. While initial studies have shown promising results more extensive investigations are required to confirm its therapeutic value.^{258,259}

Captopril (Fig. 11) is the first angiotensin-converting enzyme (ACE) inhibitor, a drug class used to treat high blood pressure. It is derived from a venom peptide, called bradykinin-potentiating factor, isolated from *Bothrops jararaca* venom.^{260,261} While intravenous applications showed antihypertensive properties, derivatising this peptide into an orally available drug while retaining antihypertensive properties paved the way for drug approval in the early eighties.^{262,263} Captopril is the prototype of all ACE inhibitor drugs, a class still used as first line treatment in cardiovascular diseases, and new indications are emerging. These include but are not limited to atherosclerosis, heart attacks, and diabetic nephropathy.²⁶⁴

Thiostrepton is prominently featured throughout this article. Although it lacks approval for human use as of yet, thiostrepton finds veterinary application. It is an approved antibiotic for skin and eye infections, such as in Animax Ointment (NDC 17033-122-75).²⁶⁵ Like thiostrepton, nosiheptide (Fig. 11) belongs to the thiopeptide class of antibiotics. It was isolated from *Streptomyces actinotus* 4003 and exhibits potent activity against Gram-positive bacteria.²⁶⁶ Additionally, nosiheptide possesses growth promoting properties, making it a beneficial feed additive for enhancing feed efficiency and weight gain in swine and chicken.^{267–269}

4 Challenges and limitations

Compared to the significant number of RiPP natural products that exhibit intriguing bioactivities, as discussed in the first



chapter 'Opportunities', only a small number of RiPPs are currently in clinical trials or on the market. The causes of this gap are diverse. The study of RiPPs and their biosynthetic machinery is much more recent compared to other natural product superfamilies, such as non-ribosomal peptides and polyketide natural products. Additional reasons include problems of stability and permeability, rapid renal elimination, and harmful side effects, among other factors. Traditionally, peptides have been considered as weak drug candidates due to their tendency to exhibit suboptimal ADME (Absorption, Distribution, Metabolism and Excretion) properties.²⁷⁰

4.1 Oral bioavailability/gastrointestinal stability

Oral drug administration is generally preferred for reasons of compliance and convenience. However, canonical peptides, which are composed of proteinogenic amino acids, are often labile compounds. The acidic pH in the stomach, as well as pancreatic and intestinal proteases, cause rapid degradation. Therefore, it is a significant challenge to deliver peptides orally, and research in this area has been ongoing for over 100 years. Since the discovery of insulin, attempts have been made by researchers to design an oral form, but without success to date.²⁷¹ Two prime examples of marketed, orally available peptides are cyclosporine A and desmopressin. The macrocyclic structure of both peptides enhances their protease and acid stability. Although RiPPs are not solely linear canonical peptides and certain post-translational modifications including epimerisations, backbone *N*-methylations, β -amino acids, and macrocyclisations can aid in improving their stability,¹⁹ acidic hydrolysis or proteases can result in partial degradation of the peptide compounds.

4.2 Absorption

Another challenge posed by RiPP candidate drugs relates to their ability to enter cells. Their large molecular weight (>500 g mol⁻¹) means that in most cases they do not satisfy the "Lipinski rule of five" criteria, and therefore have limited cell

permeability, particularly when administered orally where there are additional restrictions due to intestinal epithelial permeability. This is because peptides, which are often hydrophilic, are not able to diffuse passively through physiological barriers and cell membranes efficiently.²⁷²

One example of limited uptake is ziconotide (Prialt®), which is a drug available on the market (see Section 3.2). It cannot penetrate the blood–brain barrier and is therefore administered *via* intrathecal application. However, this approach carries the risk of severe infections, impeding the widespread use of ziconotide in clinical practice.¹⁸⁵

It should be noted that a small number of macrocyclic peptides have been approved as orally available peptide drugs. Backbone *N*-methylations can enhance their cell penetration: cyclosporine A, a nonribosomal peptide that is used as an immunosuppressant in clinical settings, serves as an example.²⁷³

4.3 Metabolic stability

Therapeutic peptides undergo rapid elimination. In particular, oral administration leads to a significant first pass effect that reduces the amount of active ingredient systemically available after absorption in the gastrointestinal tract. The peptides have a short half-life in blood plasma, and the liver and kidneys clear them from circulation in minutes.²⁷²

Glycosylation of ribosomal peptides can enhance the metabolic stability by reducing the clearance rate and/or preventing side-chain oxidation.^{274,275} Furthermore, the conjugation with lipids or synthetic polyethylene glycol (PEG) units has been shown to increase the serum half-life of peptides.^{276,277} Similar post-translational modifications, specifically glycosylation and lipidation, are naturally occurring in the families of glycosins,¹ lipolanthines,³⁴ and selidamides²⁷⁸ within the RiPP natural product class, among others. These modifications can improve the circulation of peptides in the bloodstream, which is a critical factor in their therapeutic application. Thus, these glyco- and lipopeptides are likely to exhibit enhanced metabolic stability, rendering them favourable candidates for therapeutic peptides.

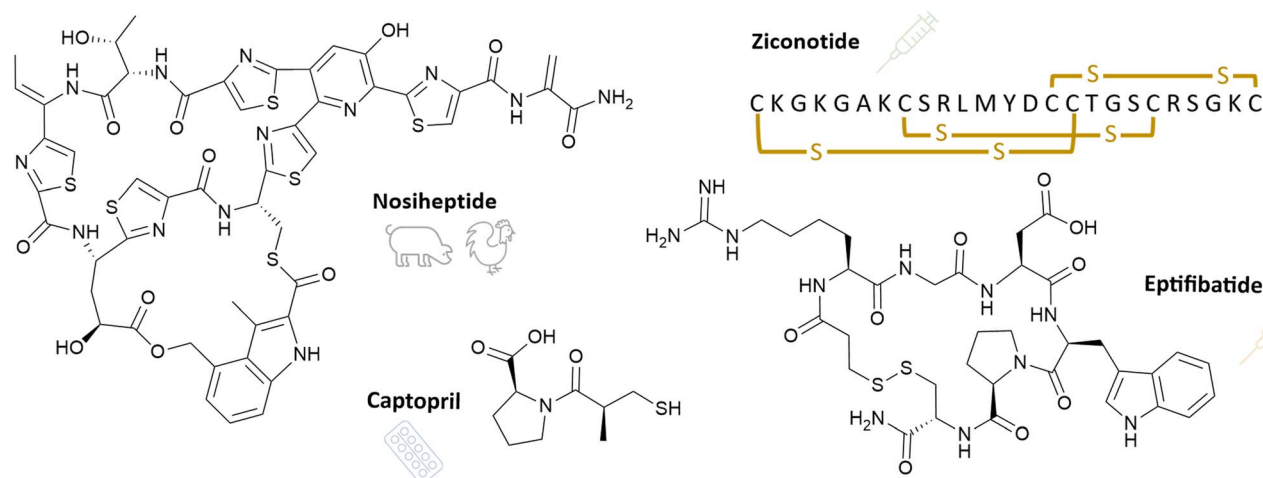


Fig. 11 RiPP(-)-based compounds, that are currently in clinical use. Chemical structures of nosiheptide, captopril, eptifibatide, and ziconotide are depicted. For ziconotide, the one-letter code is used. Yellow indicated crosslinks are disulphide bridges.



4.4 Toxicity

Canonical peptides generally possess high conformational flexibility, which can contribute to a lack of selectivity. This may cause activation of multiple target receptors, leading to undesired side effects.²⁷² In contrast, RiPPs exhibit greater specificity due to the imposition of conformational constraints *via* post-translational modifications.

Peptides have a predictable metabolism, and their metabolites are seldom toxic. Thus, ribosomal peptides have a lower incidence of undesired side effects compared to small molecules.²⁷⁶ However, like every small molecule or large biological drug, ribosomal peptides can also have toxic side effects. For instance, ziconotide shows systemic and central nervous system side effects in higher doses, including dizziness, blurred vision, nystagmus, and sedation, which ultimately subside after discontinuing the drug.¹⁸⁵

Although post-translational modifications can improve the physicochemical properties of ribosomal peptides, including oral bioavailability (*via* macrocyclisation), cell permeability (using *N*-methylamides) and metabolic stability (through glycosylation or lipidation), these structural characteristics must be combined and cannot be universally applied to every drug candidate. This leads to biosynthetic challenges and may decrease binding affinity at the target, thus diminishing the bioactivity.

5 Concluding remarks

The previous chapter highlights that RiPP(-based) therapies still have a considerable way to progress. The currently available strategies necessitate improvement, and new strategies must be developed to conquer the encountered obstacles. One of the concerns surrounding peptide medications available *via* oral administration is their stability, which can be enhanced *via* numerous post-translational modifications, such as macrocyclisation or conjugation with fatty acids. Some RiPPs exhibit inherent stability due to their specific structural features. One prominent example is the family of lasso peptides which are robust against high temperatures, acidic conditions, and proteases.^{279,280} Although oral administration may not be possible, beneficial bioactivities can result in therapeutic medicines, as different methods of treatment are attainable (for example, topical or parenteral, *e.g.*, IM, SC, IV, IT, ¶ *etc.*). In particular, for serious illnesses, patients frequently stay in the hospital, which enables drug administration beyond the oral route. Besides, a new form of drug administration addressing the buccal mucosa is currently under development: a suction patch inspired by octopus suckers that is loaded with a peptide drug and a permeation enhancer, and may serve as a viable alternative for parenteral drugs.²⁸¹

In addition to diverse methods for extending the half-life of peptide drugs, including PEGylation²⁸² and fusion proteins,²⁸³ various approaches for regulating drug release at specific body sites and over a certain period of time have been described. These

encompass the use of biodegradable polymer matrices,²⁸⁴ engineered living materials,²⁸⁵ pulmonary and intranasal drug delivery systems,^{286,287} vector-based methods and nanoparticles.^{288,289} Nanoparticle systems have already been employed in RiPPs: Nisin A, commonly used as a food preservative, is a narrow-spectrum lantibiotic that is effective against Gram-positive bacteria. However, when nisin A is combined with gold-coated nanoparticles forming so called nanocomposites, it can exhibit activity against Gram-negative bacteria.²⁹⁰ This demonstrates how new formulations can enhance the activity spectrum of antibiotics. In general, the application of new formulations represents a promising measure for unstable drug candidates. This has already been demonstrated by mRNA vaccines that utilise lipid-based vectors. A similar liposome encapsulation technique using liposomes has recently been applied to the anti-parkinsonian drug candidate acalite. To enhance the affinity to the transferrin receptor and enable brain targeting, the liposome carrying acalite was equipped with surface-bound transferrin. The experimental treatment was effectively administered to a mouse model, ultimately reaching the mouse brain and culminating six hours after being administered.²⁰⁴ This achievement underscores the significance of novel strategies for drug delivery.

RiPPs show potential as drug candidates and can also facilitate the construction of drug delivery systems. A stable conjugation method for protein therapeutics is the tetrazine ligation approach, named “TyrEx cycloaddition”, which relies on a post-translational modification installed by a RiPP maturase. This method facilitates the bonding of a drug molecule with an antibody that specifically targets tumour cells. The process of conjugation is based on an aminopyruvate unit that is introduced at a short tag through a post-translational splicing reaction catalysed by a splicease. This short tag can be genetically incorporated into a target protein or an antibody, which enables the visualisation and localisation of tumour cells as well as attacking them by releasing the drug after entering the tumour cell. Such a methodology is believed to be a significant advance towards more effective cancer therapies.²⁹¹

In the future, it is necessary to apply or adapt the drug delivery strategies mentioned earlier to promising RiPP candidate drugs. This step will enable the full realisation of RiPPs' or RiPP-based compounds' potential and widen their availability for various medical treatments.

6 Author contributions

All authors worked together on the manuscript, in detail: conceptualisation: S. M.; supervision: S. M.; visualisation: I. P., M.-P. S.; writing – original draft: I. P., M.-P. S., S. M.; writing – review and editing: S. M.

7 Conflicts of interest

There are no conflicts to declare.

8 Acknowledgements

The authors thank Harald Gross (Pharmaceutical Institute, University of Tübingen) for proofreading and critical

¶ IM = intramuscular; IT = intrathecal; IV = intravenous; SC = subcutaneous



discussions. Research in the Mordhorst lab is funded by the Baden-Württemberg Stiftung through the Postdoctoral fellowship for leading early career researchers and by the Federal Ministry of Education and Research (BMBF) and the Baden-Württemberg Ministry of Science as part of the Excellence Strategy of the German Federal and State Governments.

9 Notes and references

- 1 M. Montalbán-López, 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. Martínez, S. K. Nair, Y. Nicolet, S. Rebuffat, H.-G. Sahl, D. Sareen, E. W. Schmidt, L. Schmitt, K. Severinov, R. D. Süssmuth, 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.
- 2 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. Göransson, 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. Müller, S. K. Nair, I. F. Nes, G. E. Norris, B. M. Olivera, H. Onaka, M. L. Patchett, J. Piel, M. J. T. 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. Süssmuth, 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.
- 3 J. P. Klinman and F. Bonnot, *Chem. Rev.*, 2014, **114**, 4343–4365.
- 4 R. Ayikpoe, V. Govindarajan and J. A. Latham, *Appl. Microbiol. Biotechnol.*, 2019, **103**, 2903–2912.
- 5 R. S. Ayikpoe and J. A. Latham, *J. Am. Chem. Soc.*, 2019, **141**, 13582–13591.
- 6 G. E. Kenney and A. C. Rosenzweig, *J. Biol. Chem.*, 2018, **293**, 4606–4615.
- 7 X. Thomas, D. Destoumieux-Garzón, J. Peduzzi, C. Afonso, A. Blond, N. Birlirakis, C. Goulard, L. Dubost, R. Thai, J.-C. Tabet and S. Rebuffat, *J. Biol. Chem.*, 2004, **279**, 28233–28242.
- 8 G. Vassiliadis, J. Peduzzi, S. Zirah, X. Thomas, S. Rebuffat and D. Destoumieux-Garzón, *Antimicrob. Agents Chemother.*, 2007, **51**, 3546–3553.
- 9 Y. Li and S. Rebuffat, *J. Biol. Chem.*, 2020, **295**, 34–54.
- 10 P. Vlieghe, V. Lisowski, J. Martinez and M. Khrestchatisky, *Drug Discovery Today*, 2010, **15**, 40–56.
- 11 D. J. Craik, D. P. Fairlie, S. Liras and D. Price, *Chem. Biol. Drug Des.*, 2013, **81**, 136–147.
- 12 X. Jing and K. Jin, *Med. Res. Rev.*, 2020, **40**, 753–810.
- 13 A. Tapeinou, M.-T. Matsoukas, C. Simal and T. Tselios, *Pept. Sci.*, 2015, **104**, 453–461.
- 14 L. Cao, T. Do and A. J. Link, *J. Ind. Microbiol. Biotechnol.*, 2021, **48**, kuab005.
- 15 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.
- 16 Y. Fu, A. H. Jaarsma and O. P. Kuipers, *Cell. Mol. Life Sci.*, 2021, **78**, 3921–3940.
- 17 G. A. Hudson and D. A. Mitchell, *Curr. Opin. Microbiol.*, 2018, **45**, 61–69.
- 18 A. L. Vagstad, *Curr. Opin. Biotechnol.*, 2023, **80**, 102891.
- 19 S. Mordhorst, F. Ruijne, A. L. Vagstad, O. P. Kuipers and J. Piel, *RSC Chem. Biol.*, 2023, **4**, 7–36.
- 20 L. M. Repka, J. R. Chekan, S. K. Nair and W. A. van der Donk, *Chem. Rev.*, 2017, **117**, 5457–5520.
- 21 W. Gu, S.-H. Dong, S. Sarkar, S. K. Nair and E. W. Schmidt, in *Methods in Enzymology*, ed. B. S. Moore, Academic Press, 2018, vol. 604, pp. 113–163.
- 22 W. L. Cheung-Lee and A. J. Link, *J. Ind. Microbiol. Biotechnol.*, 2019, **46**, 1371–1379.
- 23 J. D. Hegemann, M. Zimmermann, X. Xie and M. A. Marahiel, *Acc. Chem. Res.*, 2015, **48**, 1909–1919.
- 24 J. Martins and V. Vasconcelos, *Mar. Drugs*, 2015, **13**, 6910–6946.
- 25 F. Hubrich, A. Lotti, T. A. Scott and J. Piel, *Chimia*, 2021, **75**, 543.
- 26 C. Kaletta, K.-D. Entian and G. Jung, *Eur. J. Biochem.*, 1991, **199**, 411–415.
- 27 L. A. Lindenfelser, T. G. Pridham and C. E. Kemp, *Antibiot. Chemother.*, 1959, **9**, 690–695.
- 28 A. Ökesli, L. E. Cooper, E. J. Fogle and W. A. Van Der Donk, *J. Am. Chem. Soc.*, 2011, **133**, 13753–13760.
- 29 L. Huo, A. Ökesli, M. Zhao and W. A. Van Der Donk, *Appl. Environ. Microbiol.*, 2017, **83**, e02698.
- 30 F. Märki, E. Hänni, A. Fredenhagen and J. Van Oostrum, *Biochem. Pharmacol.*, 1991, **42**, 2027–2035.
- 31 J. P. Deisinger, M. Arts, I. Kotsogianni, J.-S. Puls, F. Grein, F. J. Ortiz-López, N. I. Martin, A. Müller, O. Genilloud and T. Schneider, *iScience*, 2023, **26**, 106394.
- 32 M. Xu, F. Zhang, Z. Cheng, G. Bashiri, J. Wang, J. Hong, Y. Wang, L. Xu, X. Chen, S. Huang, S. Lin, Z. Deng and M. Tao, *Angew. Chem., Int. Ed.*, 2020, **59**, 18029–18035.
- 33 F. J. Ortiz-López, D. Carretero-Molina, M. Sánchez-Hidalgo, J. Martín, I. González, F. Román-Hurtado, M. Cruz, S. García-Fernández, F. Reyes, J. P. Deisinger, A. Müller, T. Schneider and O. Genilloud, *Angew. Chem., Int. Ed.*, 2020, **59**, 12654–12658.
- 34 V. Wiebach, A. Mainz, M.-A. J. Siegert, N. A. Jungmann, G. Lesquame, S. Tirat, A. Dreux-Zigha, J. Aszodi, D. Le Beller and R. D. Süssmuth, *Nat. Chem. Biol.*, 2018, **14**, 652–654.
- 35 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.



- 36 K. J. Molohon, J. O. Melby, J. Lee, B. S. Evans, K. L. Dunbar, S. B. Bumpus, N. L. Kelleher and D. A. Mitchell, *ACS Chem. Biol.*, 2011, **6**, 1307–1313.
- 37 F. Baquero and F. Moreno, *FEMS Microbiol. Lett.*, 1984, **23**, 117–124.
- 38 T. D. M. Pham, Z. M. Ziora and M. A. T. Blaskovich, *Med. Chem. Commun.*, 2019, **10**, 1719–1739.
- 39 J. Davagnino, M. Herrero, D. Furlong, F. Moreno and R. Kolter, *Proteins*, 1986, **1**, 230–238.
- 40 M. Herrero and F. Moreno, *Microbiology*, 1986, **132**, 393–402.
- 41 H. Onaka, H. Tabata, Y. Igarashi, Y. Sato and T. Furumai, *J. Antibiot.*, 2001, **54**, 1036–1044.
- 42 H. Onaka, M. Nakaho, K. Hayashi, Y. Igarashi and T. Furumai, *Microbiology*, 2005, **151**, 3923–3933.
- 43 H. Mathur, D. Field, M. Upton and P. D. Cotter, *Front. Microbiol.*, 2021, **12**, 695081.
- 44 J. Stepper, S. Shastri, T. S. Loo, J. C. Preston, P. Novak, P. Man, C. H. Moore, V. Havlíček, M. L. Patchett and G. E. Norris, *FEBS Lett.*, 2011, **585**, 645–650.
- 45 W. J. Kelly, R. V. Asmundson and C. M. Huang, *J. Appl. Bacteriol.*, 1996, **81**, 657–662.
- 46 A. A. Vinogradov and H. Suga, *Cell Chem. Biol.*, 2020, **27**, 1032–1051.
- 47 A. H. Kutscher, L. Seguin, E. V. Zegarelli and J. D. Piro, *J. Am. Dent. Assoc.*, 1959, **59**, 715–720.
- 48 S. B. Singh, J. Occi, H. Jayasuriya, K. Herath, M. Motyl, K. Dorso, C. Gill, E. Hickey, K. M. Overbye, J. F. Barrett and P. Masurekar, *J. Antibiot.*, 2007, **60**, 565–571.
- 49 M. J. Pucci, J. J. Bronson, J. F. Barrett, K. L. DenBleyker, L. F. Discotto, J. C. Fung-Tome and Y. Ueda, *Antimicrob. Agents Chemother.*, 2004, **48**, 3697–3701.
- 50 N. M. Haste, W. Thienphrapa, D. N. Tran, S. Loesgen, P. Sun, S.-J. Nam, P. R. Jensen, W. Fenical, G. Sakoulas, V. Nizet and M. E. Hensler, *J. Antibiot.*, 2012, **65**, 593–598.
- 51 E. Gavrish, C. S. Sit, S. Cao, O. Kandrór, A. Spoering, A. Peoples, L. Ling, A. Fetterman, D. Hughes, A. Bissell, H. Torrey, T. Akopian, A. Mueller, S. Epstein, A. Goldberg, J. Clardy and K. Lewis, *Chem. Biol.*, 2014, **21**, 509–518.
- 52 M. Iwatsuki, R. Uchida, Y. Takakusagi, A. Matsumoto, C.-L. Jiang, Y. Takahashi, M. Arai, S. Kobayashi, M. Matsumoto, J. Inokoshi, H. Tomoda and S. Ōmura, *J. Antibiot.*, 2007, **60**, 357–363.
- 53 M. Tsunakawa, S.-L. Hu, Y. Hoshino, D. J. Detlefson, S. E. Hill, T. Furumai, R. J. White, M. Nishio, K. Kawano, S. Yamamoto, Y. Fukagawa and T. Oki, *J. Antibiot.*, 1995, **48**, 433–434.
- 54 S. Tan, K. C. Ludwig, A. Müller, T. Schneider and J. R. Nodwell, *ACS Chem. Biol.*, 2019, **14**, 966–974.
- 55 D. V. Carson, Y. Zhang, L. So, W. L. Cheung-Lee, A. J. Cartagena, S. A. Darst and A. J. Link, *bioRxiv*, 2023, preprint, DOI: [10.1101/2023.06.21.545946](https://doi.org/10.1101/2023.06.21.545946).
- 56 R. A. Salomón and R. N. Fariás, *J. Bacteriol.*, 1992, **174**, 7428–7435.
- 57 T. A. Knappe, U. Linne, S. Zirah, S. Rebuffat, X. Xie and M. A. Marahiel, *J. Am. Chem. Soc.*, 2008, **130**, 11446–11454.
- 58 K. Kuznedelov, E. Semenova, T. A. Knappe, D. Mukhamedyarov, A. Srivastava, S. Chatterjee, R. H. Ebright, M. A. Marahiel and K. Severinov, *J. Mol. Biol.*, 2011, **412**, 842–848.
- 59 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. Böhringer, 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. Schäberle, A. D'Onofrio and K. Lewis, *Nature*, 2019, **576**, 459–464.
- 60 S. Groß, F. Panter, D. Pogorevc, C. E. Seyfert, S. Deckarm, C. D. Bader, J. Herrmann and R. Müller, *Chem. Sci.*, 2021, **12**, 11882–11893.
- 61 C. E. Seyfert, C. Porten, B. Yuan, S. Deckarm, F. Panter, C. D. Bader, J. Coetzee, F. Deschner, K. H. M. E. Tehrani, P. G. Higgins, H. Seifert, T. C. Marlovits, J. Herrmann and R. Müller, *Angew. Chem., Int. Ed.*, 2023, **62**, e202214094.
- 62 C. E. Seyfert, A. V. Müller, D. J. Walsh, J. Birkelbach, A. M. Kany, C. Porten, B. Yuan, D. Krug, J. Herrmann, T. C. Marlovits, A. K. H. Hirsch and R. Müller, *ChemRxiv*, 2023, preprint, DOI: [10.26434/chemrxiv-2023-24tmj](https://doi.org/10.26434/chemrxiv-2023-24tmj).
- 63 M. Meteleev, A. Arseniev, L. B. Bushin, K. Kuznedelov, T. O. Artamonova, R. Kondratenko, M. Khodorkovskii, M. R. Seyedsayamdost and K. Severinov, *ACS Chem. Biol.*, 2017, **12**, 814–824.
- 64 W. L. Cheung-Lee, M. E. Parry, A. Jaramillo Cartagena, S. A. Darst and A. J. Link, *J. Biol. Chem.*, 2019, **294**, 6822–6830.
- 65 D. V. Carson, M. Patiño, H. E. Elashal, A. J. Cartagena, Y. Zhang, M. E. Whitley, L. So, A. K. Kayser-Browne, A. M. Earl, R. P. Bhattacharyya and A. J. Link, *ACS Infect. Dis.*, 2023, **9**, 111–121.
- 66 C. Asensio, J. C. Pérez-Díaz, M. C. Martínez and F. Baquero, *Biochem. Biophys. Res. Commun.*, 1976, **69**, 7–14.
- 67 Y. Li, Y. Han, Z. Zeng, W. Li, S. Feng and W. Cao, *J. Agric. Food Chem.*, 2021, **69**, 8758–8767.
- 68 Y. Han, Y. Li, Z. Zeng, W. Li, S. Feng and W. Cao, *Microbiol. Spectrum*, 2022, **10**, e01859.
- 69 W. L. Cheung-Lee, M. E. Parry, C. Zong, A. J. Cartagena, S. A. Darst, N. D. Connell, R. Russo and A. J. Link, *ChemBioChem*, 2020, **21**, 1335–1340.
- 70 H. Knothe and G. A. Dette, *J. Antimicrob. Chemother.*, 1981, **8**, 33–41.
- 71 G. A. Pankuch, M. R. Jacobs and P. C. Appelbaum, *Antimicrob. Agents Chemother.*, 1994, **38**, 2065–2072.
- 72 G. D. Brown, D. W. Denning, N. A. R. Gow, S. M. Levitz, M. G. Netea and T. C. White, *Sci. Transl. Med.*, 2012, **4**, 165rv13.
- 73 K. I. Mohr, C. Volz, R. Jansen, V. Wray, J. Hoffmann, S. Bernecker, J. Wink, K. Gerth, M. Stadler and R. Müller, *Angew. Chem., Int. Ed.*, 2015, **54**, 11254–11258.
- 74 M. Tsuda, H. Shigemori, Y. Mikami and J. Kobayashi, *Tetrahedron*, 1993, **49**, 6785–6796.
- 75 J. Kobayashi, M. Tsuda, T. Nakamura, Y. Mikami and H. Shigemori, *Tetrahedron*, 1993, **49**, 2391–2402.



- 76 N. Mizuhara, M. Kuroda, A. Ogita, T. Tanaka, Y. Usuki and K. Fujita, *Bioorg. Med. Chem.*, 2011, **19**, 5300–5310.
- 77 V. Valiante, M. C. Monteiro, J. Martín, R. Altwasser, N. El Aouad, I. González, O. Knemeyer, E. Mellado, S. Palomo, N. de Pedro, I. Pérez-Victoria, J. R. Tormo, F. Vicente, F. Reyes, O. Genilloud and A. A. Brakhage, *Antimicrob. Agents Chemother.*, 2015, **59**, 5145–5153.
- 78 A. J. Sucher, E. B. Chahine and H. E. Balcer, *Ann. Pharmacother.*, 2009, **43**, 1647–1657.
- 79 F. Marco, M. A. Pfaller, S. A. Messer and R. N. Jones, *Diagn. Microbiol. Infect. Dis.*, 1998, **32**, 33–37.
- 80 S. Arikian, M. Lozano-Chiu, V. Paetznick and J. H. Rex, *Antimicrob. Agents Chemother.*, 2001, **45**, 327–330.
- 81 J. Piret and G. Boivin, *Front. Microbiol.*, 2021, **11**, 631736.
- 82 N. M. Bösch, M. Borsa, U. Greczmiel, B. I. Morinaka, M. Gugger, A. Oxenius, A. L. Vagstad and J. Piel, *Angew. Chem., Int. Ed.*, 2020, **59**, 11763–11768.
- 83 M. Shao, J. Ma, Q. Li and J. Ju, *Mar. Drugs*, 2019, **17**, 127.
- 84 G. Helynck, C. Dubertret, J.-F. Mayaux and J. Leboul, *J. Antibiot.*, 1993, **46**, 1756–1757.
- 85 D. Fréchet, J. D. Guillon, F. Herman, D. Faucher, G. Helynck, B. Monegier du Sorbier, J. P. Ridoux, E. James-Surcouf and M. Vuilhorgne, *Biochemistry*, 1994, **33**, 42–50.
- 86 M. Sánchez-Hidalgo, J. Martín and O. Genilloud, *Antibiotics*, 2020, **9**, 67.
- 87 G. Féfir, M. I. Petrova, G. Andrei, D. Huskens, B. Hoorelbeke, R. Snoeck, J. Vanderleyden, J. Balzarini, S. Bartoschek, M. Brönstrup, R. D. Süßmuth and D. Schols, *PLoS One*, 2013, **8**, e64010.
- 88 N. Naruse, O. Tenmyo, K. Tomita, M. Konishi, T. Miyaki, H. Kawaguchi, K. Fukase, T. Wakamiya and T. Shiba, *J. Antibiot.*, 1989, **42**, 837–845.
- 89 A. S. Richard, A. Zhang, S.-J. Park, M. Farzan, M. Zong and H. Choe, *Proc. Natl. Acad. Sci. U.S.A.*, 2015, **112**, 14682–14687.
- 90 T. E. Smith, C. D. Pond, E. Pierce, Z. P. Harmer, J. Kwan, M. M. Zachariah, M. K. Harper, T. P. Wyche, T. K. Matainaho, T. S. Bugni, L. R. Barrows, C. M. Ireland and E. W. Schmidt, *Nat. Chem. Biol.*, 2018, **14**, 179–185.
- 91 T. E. Smith, C. D. Pond, E. Pierce, Z. P. Harmer, J. Kwan, M. M. Zachariah, M. K. Harper, T. P. Wyche, T. K. Matainaho, T. S. Bugni, L. R. Barrows, C. M. Ireland and E. W. Schmidt, *Nat. Chem. Biol.*, 2018, **14**, 179–185.
- 92 G. Féfir, M. I. Petrova, G. Andrei, D. Huskens, B. Hoorelbeke, R. Snoeck, J. Vanderleyden, J. Balzarini, S. Bartoschek, M. Brönstrup, R. D. Süßmuth and D. Schols, *PLoS One*, 2013, **8**, e64010.
- 93 M. Tsunakawa, S.-L. Hu, Y. Hoshino, D. J. Detlefson, S. E. Hill, T. Furumai, R. J. White, M. Nishio, K. Kawano, S. Yamamoto, Y. Fukagawa and T. Oki, *J. Antibiot.*, 1995, **48**, 433–434.
- 94 D. H. Katz, J. F. Marcelletti, M. H. Khalil, L. E. Pope and L. R. Katz, *Proc. Natl. Acad. Sci. U.S.A.*, 1991, **88**, 10825–10829.
- 95 J. Barretina Ginesta, J. Castaner, J. Bozzo and M. Bayes, *Drugs Future*, 2005, **30**, 0469.
- 96 *Product Monograph 'Invirase'*, Hoffmann-La Roche Limited, 2020.
- 97 B. M. Watkins, *Trends Parasitol.*, 2003, **19**, 477–478.
- 98 Neglected tropical diseases, <https://www.who.int/news/item/16-07-2020-neglected-tropical-diseases-treating-more-than-one-billion-people-for-the-fifth-consecutive-year>, accessed 23 August 2023.
- 99 M. Papagianni, *Biotechnol. Adv.*, 2003, **21**, 465–499.
- 100 C. M. Scheidler, L. M. Kick and S. Schneider, *ChemBioChem*, 2019, **20**, 1479–1486.
- 101 M. N. Aminake, S. Schoof, L. Sologub, M. Leubner, M. Kirschner, H.-D. Arndt and G. Pradel, *Antimicrob. Agents Chemother.*, 2011, **55**, 1338–1348.
- 102 J. Lambert, E. Keppi, J. L. Dimarcq, C. Wicker, J. M. Reichhart, B. Dunbar, P. Lepage, A. Van Dorsselaer, J. Hoffmann and J. Fothergill, *Proc. Natl. Acad. Sci. U.S.A.*, 1989, **86**, 262–266.
- 103 S.-H. Liu, H.-F. Li, Y. Yang, D. Wei, H.-B. Jiang, W. Dou, G.-R. Yuan and J.-J. Wang, *AMB Express*, 2018, **8**, 5.
- 104 P. Bulet, S. Cociancich, M. Reuland, F. Sauber, R. Bischoff, G. Hegy, A. Van DORSSLAER, C. Hetru and J. A. Hoffmann, *Eur. J. Biochem.*, 1992, **209**, 977–984.
- 105 R. B. Williams, *Avian Pathol.*, 2005, **34**, 29–47.
- 106 R. Pigeault, J. Vézilier, S. Cornet, F. Zélé, A. Nicot, P. Perret, S. Gandon and A. Rivero, *Philos. Trans. R. Soc., B*, 2015, **370**, 20140300.
- 107 M. Shahabuddin, I. Fields, P. Bulet, J. A. Hoffmann and L. H. Miller, *Exp. Parasitol.*, 1998, **89**, 103–112.
- 108 N. Boulanger, C. Lowenberger, P. Volf, R. Ursic, L. Sigutova, L. Sabatier, M. Svobodova, S. M. Beverley, G. Späth, R. Brun, B. Pesson and P. Bulet, *Infect. Immun.*, 2004, **72**, 7140–7146.
- 109 E. Torres-Guerrero, M. R. Quintanilla-Cedillo, J. Ruiz-Esmenjaud and R. Arenas, *F1000Research*, 2017, **6**, 750.
- 110 C. Portmann, S. Sieber, S. Wirthensohn, J. F. Blom, L. Da Silva, E. Baudat, M. Kaiser, R. Brun and K. Gademann, *J. Nat. Prod.*, 2014, **77**, 557–562.
- 111 M. S. Donia, B. Wang, D. C. Dunbar, P. V. Desai, A. Patny, M. Avery and M. T. Hamann, *J. Nat. Prod.*, 2008, **71**, 941–945.
- 112 R. G. Linington, J. González, L.-D. Ureña, L. I. Romero, E. Ortega-Barria and W. H. Gerwick, *J. Nat. Prod.*, 2007, **70**, 397–401.
- 113 O. Saether, D. J. Craik, I. D. Campbell, K. Sletten, J. Juul and D. G. Norman, *Biochemistry*, 1995, **34**, 4147–4158.
- 114 D. C. Ireland, M. L. Colgrave and D. J. Craik, *Biochem. J.*, 2006, **400**, 1–12.
- 115 D. J. Diemert, J. M. Bethony and P. J. Hotez, *Clin. Infect. Dis.*, 2008, **46**, 282–288.
- 116 M. L. Colgrave, A. C. Kotze, S. Kopp, J. S. McCarthy, G. T. Coleman and D. J. Craik, *Acta Trop.*, 2009, **109**, 163–166.
- 117 R. Khositnithikul, P. Tan-ariya and M. Mungthin, *Malar. J.*, 2008, **7**, 23.
- 118 S. W. Page, in *Small Animal Clinical Pharmacology*, ed. J. E. Maddison, S. W. Page and D. B. Church, W. B. Saunders, Edinburgh, Second Edition, 2008, pp. 198–260.



- 119 M. V. Díaz, M. R. Miranda, C. Campos-Estrada, C. Reigada, J. D. Maya, C. A. Pereira and R. López-Muñoz, *Acta Trop.*, 2014, **134**, 1–9.
- 120 R. C. Zauli-Nascimento, D. C. Miguel, J. K. U. Yokoyama-Yasunaka, L. I. A. Pereira, M. A. Pelli de Oliveira, F. Ribeiro-Dias, M. L. Dorta and S. R. B. Uliana, *Trop. Med. Int. Health*, 2010, **15**, 68–76.
- 121 Cancer, <https://www.who.int/news-room/fact-sheets/detail/cancer>, accessed 29 August 2023.
- 122 A. Kamb, S. Wee and C. Lengauer, *Nat. Rev. Drug Discovery*, 2007, **6**, 115–120.
- 123 H. Nishimura, S. Okamoto, M. Mayama, H. Ohtsuka, K. Nakajima, K. Tawara, M. Shimohira and N. Shimaoka, *J. Antibiot.*, 1961, **14**, 255–263.
- 124 S. K. Radhakrishnan, U. G. Bhat, D. E. Hughes, I.-C. Wang, R. H. Costa and A. L. Gartel, *Cancer Res.*, 2006, **66**, 9731–9735.
- 125 N. S. Hegde, D. A. Sanders, R. Rodriguez and S. Balasubramanian, *Nat. Chem.*, 2011, **3**, 725–731.
- 126 S. S. Myatt and E. W.-F. Lam, *Nat. Rev. Cancer*, 2007, **7**, 847–859.
- 127 J. A. V. Lopez, S. S. Al-Lihaibi, W. M. Alarif, A. Abdel-Lateff, Y. Nogata, K. Washio, M. Morikawa and T. Okino, *J. Nat. Prod.*, 2016, **79**, 1213–1218.
- 128 C. J. Hawkins, M. F. Lavin, K. A. Marshall, A. L. Van den Brenk and D. J. Watters, *J. Med. Chem.*, 1990, **33**, 1634–1638.
- 129 A. R. Carroll, J. C. Coll, D. J. Bourne, J. K. Macleod, T. M. Zabriskie, C. M. Ireland and B. F. Bowden, *Aust. J. Chem.*, 1996, **49**, 659–667.
- 130 A. Rudi, M. Aknin, E. M. Gaydou and Y. Kashman, *Tetrahedron*, 1998, **54**, 13203–13210.
- 131 S. Soltani, R. Hammami, P. D. Cotter, S. Rebuffat, L. B. Said, H. Gaudreau, F. Bédard, E. Biron, D. Drider and I. Fliss, *FEMS Microbiol. Rev.*, 2021, **45**, fuaa039.
- 132 M. A. Varas, C. Muñoz-Montecinos, V. Kallens, V. Simon, M. L. Allende, A. E. Marcoleta and R. Lagos, *Front. Microbiol.*, 2020, **11**, 405.
- 133 C. Hetz, M. R. Bono, L. F. Barros and R. Lagos, *Proc. Natl. Acad. Sci. U.S.A.*, 2002, **99**, 2696–2701.
- 134 S. Ahmadi, M. Ghollasi and H. M. Hosseini, *Microb. Pathog.*, 2017, **111**, 193–197.
- 135 N. E. Joo, K. Ritchie, P. Kamarajan, D. Miao and Y. L. Kapila, *Cancer Med.*, 2012, **1**, 295–305.
- 136 P. Kamarajan, T. Hayami, B. Matte, Y. Liu, T. Danciu, A. Ramamoorthy, F. Worden, S. Kapila and Y. Kapila, *PLoS One*, 2015, **10**, e0131008.
- 137 EFSA Panel on Food Additives and Nutrient Sources added to Food (ANS), M. Younes, P. Aggett, F. Aguilar, R. Crebelli, B. Dusemund, M. Filipič, M. J. Frutos, P. Galtier, U. Gundert-Remy, G. G. Kuhnle, C. Lambré, J.-C. Leblanc, I. T. Lillegaard, P. Moldeus, A. Mortensen, A. Oskarsson, I. Stankovic, I. Waalkens-Berendsen, R. A. Woutersen, M. Wright, L. Herman, P. Tobback, F. Pizzo, C. Smeraldi, A. Tard, A. Papaioannou and D. Gott, *EFSA J.*, 2017, **15**, e05063.
- 138 H. R. Momeni, *Cell J.*, 2011, **13**, 65.
- 139 P. A. Jordan and B. S. Moore, *Cell Chem. Biol.*, 2016, **23**, 1504–1514.
- 140 C. C. Hughes, J. B. MacMillan, S. P. Gaudêncio, P. R. Jensen and W. Fenical, *Angew. Chem., Int. Ed.*, 2009, **48**, 725–727.
- 141 D. Reimer and C. C. Hughes, *J. Nat. Prod.*, 2017, **80**, 126–133.
- 142 C. C. Hughes, J. B. MacMillan, S. P. Gaudêncio, W. Fenical and J. J. La Clair, *Angew. Chem., Int. Ed.*, 2008, **121**, 742–746.
- 143 C. M. Celli, N. Tran, R. Knox and A. K. Jaiswal, *Biochem. Pharmacol.*, 2006, **72**, 366–376.
- 144 W. Li, C. Zhang, H. Zhang, R. Dong, J. Liu, C. Wang, M. Wang, Y. Wang, C. Wang, Y. Zhang, L. Shi, Y. Xu and L.-P. Sun, *Bioorg. Chem.*, 2022, **127**, 105917.
- 145 B. Donati, E. Lorenzini and A. Ciarrocchi, *Mol. Cancer*, 2018, **17**, 164.
- 146 J. Luo, D. Yang, Hindra, A. Adhikari, L.-B. Dong, F. Ye, X. Yan, C. Rader and B. Shen, *J. Ind. Microbiol. Biotechnol.*, 2021, **48**, kuab027.
- 147 H. Nagata, K. Ochiai, Y. Aotani, K. Ando, M. Yoshida, I. Takahashi and T. Tamaoki, *J. Antibiot.*, 1997, **50**, 537–542.
- 148 A. Miyana, J. E. Janso, L. McDonald, M. He, H. Liu, L. Barbieri, A. S. Eustáquio, E. N. Fielding, G. T. Carter, P. R. Jensen, X. Feng, M. Leighton, F. E. Koehn and B. S. Moore, *J. Am. Chem. Soc.*, 2011, **133**, 13311–13313.
- 149 T. Tian, X. Li and J. Zhang, *Int. J. Mol. Sci.*, 2019, **20**, 755.
- 150 P. Lin, J. H. Wong and T. B. Ng, *Biosci. Rep.*, 2009, **30**, 101–109.
- 151 Y. Hanaoka, Y. Yamaguchi, H. Yamamoto, M. Ishii, T. Nagase, H. Kurihara, M. Akishita and Y. Ouchi, *Anticancer Res.*, 2016, **36**, 5999–6004.
- 152 S. Wang, S. Lin, Q. Fang, R. Gyampoh, Z. Lu, Y. Gao, D. J. Clarke, K. Wu, L. Trembleau, Y. Yu, K. Kyeremeh, B. F. Milne, J. Tabudravu and H. Deng, *Nat. Commun.*, 2022, **13**, 5044.
- 153 A. Ványolós, M. Dékány, B. Kovács, B. Krámos, P. Bérdi, I. Zupkó, J. Hohmann and Z. Béni, *Org. Lett.*, 2016, **18**, 2688–2691.
- 154 K. Shin-ya, K. Wierzba, K. Matsuo, T. Ohtani, Y. Yamada, K. Furihata, Y. Hayakawa and H. Seto, *J. Am. Chem. Soc.*, 2001, **123**, 1262–1263.
- 155 K. Amagai, H. Ikeda, J. Hashimoto, I. Kozono, M. Izumikawa, F. Kudo, T. Eguchi, T. Nakamura, H. Osada, S. Takahashi and K. Shin-ya, *Sci. Rep.*, 2017, **7**, 3382.
- 156 A. Nakajima, T. Tauchi, G. Sashida, M. Sumi, K. Abe, K. Yamamoto, J. H. Ohyashiki and K. Ohyashiki, *Leukemia*, 2003, **17**, 560–567.
- 157 M. A. Shammas, R. J. S. Reis, C. Li, H. Koley, L. H. Hurley, K. C. Anderson and N. C. Munshi, *Clin. Cancer Res.*, 2004, **10**, 770–776.
- 158 T. Nakamura, S. Okabe, H. Yoshida, K. Iida, Y. Ma, S. Sasaki, T. Yamori, K. Shin-ya, I. Nakano, K. Nagasawa and H. Seimiya, *Sci. Rep.*, 2017, **7**, 3605.
- 159 M. Yasuda, Y. Ma, S. Okabe, Y. Wakabayashi, D. Su, Y.-T. Chang, H. Seimiya, M. Tera and K. Nagasawa, *Chem. Commun.*, 2020, **56**, 12905–12908.



- 160 S. Um, Y.-J. Kim, H. Kwon, H. Wen, S.-H. Kim, H. C. Kwon, S. Park, J. Shin and D.-C. Oh, *J. Nat. Prod.*, 2013, **76**, 873–879.
- 161 S. S. Elsayed, F. Trusch, H. Deng, A. Raab, I. Prokes, K. Busarakam, J. A. Asenjo, B. A. Andrews, P. van West, A. T. Bull, M. Goodfellow, Y. Yi, R. Ebel, M. Jaspars and M. E. Rateb, *J. Org. Chem.*, 2015, **80**, 10252–10260.
- 162 S. Son, M. Jang, B. Lee, Y.-S. Hong, S.-K. Ko, J.-H. Jang and J. S. Ahn, *J. Nat. Prod.*, 2018, **81**, 2205–2211.
- 163 L. Digal, S. C. Samson, M. A. Stevens, A. Ghorai, H. Kim, M. C. Mifflin, K. R. Carney, D. L. Williamson, S. Um, G. Nagy, D.-C. Oh, M. C. Mendoza and A. G. Roberts, *ACS Chem. Biol.*, 2024, **19**, 81–88.
- 164 J. Ogino, R. E. Moore, G. M. L. Patterson and C. D. Smith, *J. Nat. Prod.*, 1996, **59**, 581–586.
- 165 A. B. Williams and R. S. Jacobs, *Cancer Lett.*, 1993, **71**, 97–102.
- 166 J. F. Guerrero-Garzón, E. Madland, M. Zehl, M. Singh, S. Rezaei, F. L. Aachmann, G. Courtade, E. Urban, C. Rückert, T. Busche, J. Kalinowski, Y.-R. Cao, Y. Jiang, C. Jiang, G. Selivanova and S. B. Zotchev, *iScience*, 2020, **23**, 101785.
- 167 L. A. Salvador-Reyes and H. Luesch, *Nat. Prod. Rep.*, 2015, **32**, 478–503.
- 168 I. Nakano, K. Joshi, K. Visnyei, B. Hu, M. Watanabe, D. Lam, E. Wexler, K. Saigusa, Y. Nakamura, D. R. Laks, P. S. Mischel, M. Viapiano and H. I. Kornblum, *J. Neuro-Oncol.*, 2011, **13**, 622–634.
- 169 D. B. Longley, D. P. Harkin and P. G. Johnston, *Nat. Rev. Cancer*, 2003, **3**, 330–338.
- 170 A. Beck, M. C. Etienne, S. Chéradame, J. L. Fischel, P. Formento, N. Renée and G. Milano, *Eur. J. Cancer*, 1994, **30**, 1517–1522.
- 171 S. M. Rikard, *MMWR Morb. Mortal. Wkly. Rep.*, 2023, **72**, 379–385.
- 172 S. P. Cohen, L. Vase and W. M. Hooten, *Lancet*, 2021, **397**, 2082–2097.
- 173 A. J. Kohn, P. R. Saunders and S. Wiener, *Ann. N. Y. Acad. Sci.*, 1960, **90**, 706–725.
- 174 K. Meindl, T. Schmiederer, K. Schneider, A. Reicke, D. Butz, S. Keller, H. Gühring, L. Vértessy, J. Wink, H. Hoffmann, M. Brönstrup, G. M. Sheldrick and R. D. Süssmuth, *Angew. Chem., Int. Ed.*, 2010, **49**, 1151–1154.
- 175 M. Iorio, O. Sasso, S. I. Maffioli, R. Bertorelli, P. Monciardini, M. Sosio, F. Bonezzi, M. Summa, C. Brunati, R. Bordoni, G. Corti, G. Tarozzo, D. Piomelli, A. Reggiani and S. Donadio, *ACS Chem. Biol.*, 2014, **9**, 398–404.
- 176 Z. Liu, P. Bartels, M. Sadeghi, T. Du, Q. Dai, C. Zhu, S. Yu, S. Wang, M. Dong, T. Sun, J. Guo, S. Peng, L. Jiang, D. J. Adams and Q. Dai, *Sci. Rep.*, 2018, **8**, 1004.
- 177 G. Berecki, L. Motin, A. Haythornthwaite, S. Vink, P. Bansal, R. Drinkwater, C. I. Wang, M. Moretta, R. J. Lewis, P. F. Alewood, M. J. Christie and D. J. Adams, *Mol. Pharmacol.*, 2010, **77**, 139–148.
- 178 C. I. Schroeder, C. J. Doering, G. W. Zamponi and R. J. Lewis, *Med. Chem.*, 2006, **2**, 535–543.
- 179 T. Wu, M. Wang, W. Wu, Q. Luo, L. Jiang, H. Tao and M. Deng, *J. Venomous Anim. Toxins Incl. Trop. Dis.*, 2019, **25**, e146318.
- 180 P. Escoubas, J. R. De Weille, A. Lecoq, S. Diochot, R. Waldmann, G. Champigny, D. Moinier, A. Ménez and M. Lazdunski, *J. Biol. Chem.*, 2000, **275**, 25116–25121.
- 181 M. Mazzuca, C. Heurteaux, A. Alloui, S. Diochot, A. Baron, N. Voilley, N. Blondeau, P. Escoubas, A. Gélot, A. Cupo, A. Zimmer, A. M. Zimmer, A. Eschalier and M. Lazdunski, *Nat. Neurosci.*, 2007, **10**, 943–945.
- 182 S. M. Oliveira, C. R. Silva, G. Trevisan, J. G. Villarinho, M. N. Cordeiro, M. Richardson, M. H. Borges, C. J. Castro, M. V. Gomez and J. Ferreira, *Pflug. Arch. Eur. J. Physiol.*, 2016, **468**, 881–894.
- 183 K. Konno, G. Picolo, V. P. Gutierrez, P. Brigatte, V. O. Zambelli, A. C. M. Camargo and Y. Cury, *Peptides*, 2008, **29**, 1293–1304.
- 184 G. Berecki, L. Motin, A. Haythornthwaite, S. Vink, P. Bansal, R. Drinkwater, C. I. Wang, M. Moretta, R. J. Lewis, P. F. Alewood, M. J. Christie and D. J. Adams, *Mol. Pharmacol.*, 2010, **77**, 139–148.
- 185 J. G. McGivern, *Neuropsychiatr. Dis. Treat.*, 2007, **3**, 69–85.
- 186 X. Wang, X. Chen, Z.-J. Wang, M. Zhuang, L. Zhong, C. Fu, R. Garcia, R. Müller, Y. Zhang, J. Yan, D. Wu and L. Huo, *J. Am. Chem. Soc.*, 2023, **145**, 16924–16937.
- 187 Y. Kido, T. Hamakado, T. Yoshida, M. Anno, Y. Motoki, T. Wakamiya and T. Shiba, *J. Antibiot.*, 1983, **36**, 1295–1299.
- 188 T. Wakamiya, Y. Ueki, T. Shiba, Y. Kido and Y. Motoki, *Tetrahedron Lett.*, 1985, **26**, 665–668.
- 189 T. Wakamiya, Y. Ueki, T. Shiba, Y. Kido and Y. Motoki, *Bull. Chem. Soc. Jpn.*, 1990, **63**, 1032–1038.
- 190 W. Weber, W. Fischli, E. Hochuli, E. Kupfer and E. K. Weibel, *J. Antibiot.*, 1991, **44**, 164–171.
- 191 D. F. Wyss, H.-W. Lahm, M. Manneberg and A. M. Labhardt, *J. Antibiot.*, 1991, **44**, 172–180.
- 192 Y. Morishita, S. Chiba, E. Tsukuda, T. Tanaka, T. Ogawa, M. Yamasaki, M. Yoshida, I. Kawamoto and Y. Matsuda, *J. Antibiot.*, 1994, **47**, 269–275.
- 193 T. Ogawa, K. Ochiai, T. Tanaka, E. Tsukuda, S. Chiba, K. Yano, M. Yamasaki, M. Yoshida and Y. Matsuda, *J. Antibiot.*, 1995, **48**, 1213–1220.
- 194 D. Oves-Costales, M. Sánchez-Hidalgo, J. Martín and O. Genilloud, *Mar. Drugs*, 2020, **18**, 238.
- 195 T. Suzawa, K. Shibata, T. Tanaka, Y. Matsuda and M. Yamasaki, *Bioorg. Med. Chem. Lett.*, 1997, **7**, 1715–1720.
- 196 K. Shibata, T. Suzawa, T. Ohno, K. Yamada, T. Tanaka, E. Tsukuda, Y. Matsuda and M. Yamasaki, *Bioorg. Med. Chem.*, 1998, **6**, 2459–2467.
- 197 K. Shibata, K. Yano, T. Tanaka, Y. Matsuda and M. Yamasaki, *Lett. Pept. Sci.*, 1997, **4**, 167–170.
- 198 T. A. Knappe, U. Linne, X. Xie and M. A. Marahiel, *FEBS Lett.*, 2010, **584**, 785–789.
- 199 O. Potterat, K. Wagner, G. Gemmecker, J. Mack, C. Puder, R. Vettermann and R. Streicher, *J. Nat. Prod.*, 2004, **67**, 1528–1531.
- 200 M. Lotfy, H. Kalasz, G. Szalai, J. Singh and E. Adeghate, *Open Med. Chem. J.*, 2014, **8**, 28–35.



- 201 J. G. McCormack, N. Westergaard, M. Kristiansen, C. L. Brand and J. Lau, *Curr. Pharm. Des.*, 2001, **7**, 1451–1474.
- 202 H. J. Shin, H. Matsuda, M. Murakami and K. Yamaguchi, *Tetrahedron*, 1996, **52**, 13129–13136.
- 203 R. A. Al-Horani and U. R. Desai, *Med. Res. Rev.*, 2014, **34**, 1168–1216.
- 204 Z. Tong, X. Xie, H. Ge, R. Jiao, T. Wang, X. Wang, W. Zhuang, G. Hu and R. Tan, *Acta Pharm. Sin. B.*, 2024, **14**, 881–892.
- 205 J. D. Walton, H. E. Hallen-Adams and H. Luo, *Pept. Sci.*, 2010, **94**, 659–664.
- 206 M. Romani and J. Auwerx, *Bio-Protoc.*, 2021, **11**, e4183.
- 207 A. Anastasi, V. Erspamer and M. Bucci, *Experientia*, 1971, **27**, 166–167.
- 208 C. Schweinsberg, V. Maes, L. Brans, P. Bläuenstein, D. A. Tourwé, P. A. Schubiger, R. Schibli and E. G. Garayoa, *Bioconjugate Chem.*, 2008, **19**, 2432–2439.
- 209 S. M. Okarvi and I. A. Jammaz, *Nucl. Med. Biol.*, 2012, **39**, 795–804.
- 210 N. Sadeghzadeh, M. Ahmadzadeh and M. Erfani, *J. Radioanal. Nucl. Chem.*, 2013, **298**, 287–293.
- 211 P. Hoppenz, S. Els-Heindl and A. G. Beck-Sickinger, *J. Pept. Sci.*, 2019, **25**, e3224.
- 212 R. Eychenne, C. Bouvry, M. Bourgeois, P. Loyer, E. Benoist and N. Lepareur, *Molecules*, 2020, **25**, 4012.
- 213 V. Ambrosini, L. Zanoni, A. Filice, G. Lamberti, G. Argalia, E. Fortunati, D. Campana, A. Versari and S. Fanti, *Cancers*, 2022, **14**, 1055.
- 214 L. Joosten, M. Brom, H. Peeters, S. Heskamp, M. Béhé, O. Boerman and M. Gotthardt, *Mol. Pharm.*, 2018, **15**, 486–494.
- 215 I. Velikyan and O. Eriksson, *Theranostics*, 2020, **10**, 437–461.
- 216 L. Aloj, M. R. Panico, C. Caracó, A. Zannetti, S. Del Vecchio, C. Di Nuzzo, C. Arra, G. Morelli, D. Tesauero, S. De Luca, C. Pedone and M. Salvatore, *Biopolymers*, 2002, **66**, 370–380.
- 217 S. Roosenburg, P. Laverman, F. L. van Delft and O. C. Boerman, *Amino Acids*, 2011, **41**, 1049–1058.
- 218 X. Yang, K. R. Lennard, C. He, M. C. Walker, A. T. Ball, C. Doigneaux, A. Tavassoli and W. A. Van Der Donk, *Nat. Chem. Biol.*, 2018, **14**, 375–380.
- 219 K. J. Hetrick, M. C. Walker and W. A. van der Donk, *ACS Cent. Sci.*, 2018, **4**, 458–467.
- 220 J. D. Hegemann, S. C. Bobeica, M. C. Walker, I. R. Bothwell and W. A. van der Donk, *ACS Synth. Biol.*, 2019, **8**, 1204–1214.
- 221 T. A. Knappe, F. Manzenrieder, C. Mas-Moruno, U. Linne, F. Sasse, H. Kessler, X. Xie and M. A. Marahiel, *Angew. Chem., Int. Ed.*, 2011, **50**, 8714–8717.
- 222 J. D. Hegemann, M. De Simone, M. Zimmermann, T. A. Knappe, X. Xie, F. S. Di Leva, L. Marinelli, E. Novellino, S. Zahler, H. Kessler and M. A. Marahiel, *J. Med. Chem.*, 2014, **57**, 5829–5834.
- 223 K. Mohri, K. P. H. Nhat, M. Zouda, S. Warashina, Y. Wada, Y. Watanabe, S. Tagami and H. Mukai, *Eur. J. Pharm. Sci.*, 2023, **180**, 106339.
- 224 R. H. Kimura, A. M. Levin, F. V. Cochran and J. R. Cochran, *Proteins: Struct., Funct., Bioinf.*, 2009, **77**, 359–369.
- 225 A. P. Silverman, A. M. Levin, J. L. Lahti and J. R. Cochran, *J. Mol. Biol.*, 2009, **385**, 1064–1075.
- 226 A. C. Conibear, A. Bochen, K. J. Rosengren, P. Stupar, C. Wang, H. Kessler and D. J. Craik, *ChemBioChem*, 2014, **15**, 451–459.
- 227 B. J. Burkhardt, G. A. Hudson, K. L. Dunbar and D. A. Mitchell, *Nat. Chem. Biol.*, 2015, **11**, 564–570.
- 228 B. J. Burkhardt, N. Kakkar, G. A. Hudson, W. A. van der Donk and D. A. Mitchell, *ACS Cent. Sci.*, 2017, **3**, 629–638.
- 229 E. Reyna-González, B. Schmid, D. Petras, R. D. Süssmuth and E. Dittmann, *Angew. Chem., Int. Ed.*, 2016, **55**, 9398–9401.
- 230 T. J. Oman, P. J. Knerr, N. A. Bindman, J. E. Velásquez and W. A. van der Donk, *J. Am. Chem. Soc.*, 2012, **134**, 6952–6955.
- 231 Y. Goto, Y. Ito, Y. Kato, S. Tsunoda and H. Suga, *Chem. Biol.*, 2014, **21**, 766–774.
- 232 J. Koehnke, G. Mann, A. F. Bent, H. Ludewig, S. Shirran, C. Botting, T. Lebl, W. E. Houssen, M. Jaspars and J. H. Naismith, *Nat. Chem. Biol.*, 2015, **11**, 558–563.
- 233 L. Franz and J. Koehnke, *Chem. Commun.*, 2021, **57**, 6372–6375.
- 234 Y. Goto, T. Katoh and H. Suga, *Nat. Protoc.*, 2011, **6**, 779–790.
- 235 S. R. Fleming, T. E. Bartges, A. A. Vinogradov, C. L. Kirkpatrick, Y. Goto, H. Suga, L. M. Hicks and A. A. Bowers, *J. Am. Chem. Soc.*, 2019, **141**, 758–762.
- 236 E. Eber, M. Trawinska-Bartnicka, D. Sands, G. Bellon, U. Mellies, K. Bolbás, S. Quattrucci, H. Mazurek, R. Widmann, C. Schoergenhofer, B. Jilma and F. Ratjen, *J. Cystic Fibrosis*, 2021, **20**, 61–67.
- 237 E. Eber, *Lancovutide (Moli1901) Inhalation Solution Study in Adolescents and Adults With Cystic Fibrosis*, ClinicalTrials.gov, Identifier: NCT00671736, 2007.
- 238 *Safety and Efficacy of Multiple Daily Dosing of Oral LFF571 in Patients With Moderate Clostridium Difficile Infections*, ClinicalTrials.gov, Identifier: NCT01232595, 2010.
- 239 D. C. K. Chan and L. L. Burrows, *J. Antibiot.*, 2021, **74**, 161–175.
- 240 A. Fabbretti, C.-G. He, E. Gaspari, S. Maffioli, L. Brandi, R. Spurio, M. Sosio, D. Jabes and S. Donadio, *Antimicrob. Agents Chemother.*, 2015, **59**, 4560–4568.
- 241 *Clinical efficacy and safety of NAI-Acne gel 3% applied twice-a-day to patients with facial acne vulgaris*, European Union Clinical Trials Register, Identifier: 2014-001491-62, 2014.
- 242 B. G. Livett, D. W. Sandall, D. Keays, J. Down, K. R. Gayler, N. Satkunanathan and Z. Khalil, *Toxicol.*, 2006, **48**, 810–829.
- 243 C. Herd, *A randomised, placebo-controlled, double-blind, single and multiple ascending dose study to assess the safety, tolerability, pharmacokinetics and pharmacodynamics of subcutaneous doses of ACV1 in healthy adult male subjects*, Australian New Zealand Clinical Trials Registry, Identifier: ACTRN12605000408684, 2005.
- 244 C. Herd, *A randomised, double-blind, placebo-controlled study to assess the safety, tolerability, pharmacodynamics,*



- and pharmacokinetics of subcutaneous doses of ACV1 in patients with diabetic peripheral neuropathic pain or post-herpetic neuralgia, Australian New Zealand Clinical Trials Registry, Identifier: ACTRN12607000201471, 2007.
- 245 Metabolic discontinues clinical trial programme for neuropathic pain drug, ACV1, Metabolic Pharmaceuticals Ltd, 2007.
 - 246 I. A. Sharpe, J. Gehrmann, M. L. Loughnan, L. Thomas, D. A. Adams, A. Atkins, E. Palant, D. J. Craik, D. J. Adams, P. F. Alewood and R. J. Lewis, *Nat. Neurosci.*, 2001, **4**, 902–907.
 - 247 A. Brust, E. Palant, D. E. Croker, B. Colless, R. Drinkwater, B. Patterson, C. I. Schroeder, D. Wilson, C. K. Nielsen, M. T. Smith, D. Alewood, P. F. Alewood and R. J. Lewis, *J. Med. Chem.*, 2009, **52**, 6991–7002.
 - 248 A. Brust, Conopeptide to drug: the development, structure and activity correlation of Xen2174, *29th European Peptide Symposium*, September 3 – 8, 2006, Gdansk, Poland, 2006.
 - 249 P. Okkerse, J. L. Hay, E. Sitsen, A. Dahan, E. Klaassen, W. Houghton and G. J. Groeneveld, *Br. J. Clin. Pharmacol.*, 2017, **83**, 751–763.
 - 250 A randomized, double-blind, placebo-controlled, serial-cohort, single ascending dose of Xen2174 PK/PD study administered intrathecally in healthy volunteers, Centrale Commissie Mensgebonden Onderzoek, Identifier: NL38941.056.11, 2011.
 - 251 A randomized, double-blind, placebo-controlled Phase I study to evaluate the safety and tolerability of intrathecally administered single ascending bolus doses of Xen2174 in healthy subjects, Centrale Commissie Mensgebonden Onderzoek, Identifier: NL29372.040.09, 2009.
 - 252 A randomized, double-blind, placebo-controlled, serial-cohort, single ascending dose of Xen2174 with bupivacaine interaction study administered intrathecally in healthy volunteers, Centrale Commissie Mensgebonden Onderzoek, Identifier: NL37832.058.11, 2011.
 - 253 F. T. Shafiei, R. K. McAllister and J. Lopez, in *StatPearls*, StatPearls Publishing, Treasure Island (FL), 2023.
 - 254 C. Allerton and Royal Society of Chemistry, *Pain therapeutics: current and future treatment paradigms*, RSC Publishing, Cambridge, 2014.
 - 255 B. M. Olivera, L. J. Cruz, V. De Santos, G. LeCheminant, D. Griffin, R. Zeikus, J. M. McIntosh, R. Galyean and J. Varga, *Biochemistry*, 1987, **26**, 2086–2090.
 - 256 J. E. Pope and T. R. Deer, *Expert Opin. Pharmacother.*, 2013, **14**, 957–966.
 - 257 R. M. Scarborough, J. W. Rose, M. A. Hsu, D. R. Phillips, V. A. Fried, A. M. Campbell, L. Nannizzi and I. F. Charo, *J. Biol. Chem.*, 1991, **266**, 9359–9362.
 - 258 R. M. Scarborough, *Am. Heart J.*, 1999, **138**, 1093–1104.
 - 259 G. Tonin and J. Klen, *Int. J. Mol. Sci.*, 2023, **24**, 5446.
 - 260 S. H. Ferreira, *Br. J. Pharmacol.*, 1965, **24**, 163–169.
 - 261 Y. S. Bakhle, *Nature*, 1968, **220**, 919–921.
 - 262 H. Gavras, H. R. Brunner, G. A. Turini, G. R. Kershaw, C. P. Tifft, S. Cuttelod, I. Gavras, R. A. Vukovich and D. N. McKinstry, *N. Engl. J. Med.*, 1978, **298**, 991–995.
 - 263 A. B. Atkinson and J. I. S. Robertson, *Lancet*, 1979, **314**, 836–839.
 - 264 K. Hanif, H. K. Bid and R. Konwar, *Hypertens. Res.*, 2010, **33**, 11–21.
 - 265 C. Bailly, *Eur. J. Pharmacol.*, 2022, **914**, 174661.
 - 266 F. Benazet, M. Cartier, J. Florent, C. Godard, G. Jung, J. Lunel, D. Mancy, C. Pascal, J. Renaut, P. Tarridec, J. Theilleux, R. Tissier, M. Dubost and L. Ninet, *Experientia*, 1980, **36**, 414–416.
 - 267 C. H. McGinnis, C. A. Johnson and J. E. Fox, *Poult. Sci.*, 1978, **57**, 1641–1645.
 - 268 F. Benazet and J. R. Cartier, *Poult. Sci.*, 1980, **59**, 1405–1415.
 - 269 G. L. Cromwell, T. S. Stahly, V. C. Speer and R. O'Kelly, *J. Anim. Sci.*, 1984, **59**, 1125–1128.
 - 270 M. R. Naylor, A. T. Bockus, M.-J. Blanco and R. S. Lokey, *Curr. Opin. Chem. Biol.*, 2017, **38**, 141–147.
 - 271 D. J. Brayden and M.-J. Alonso, *Adv. Drug Delivery Rev.*, 2016, **106**, 193–195.
 - 272 Y. Haggag, S. El-Gizawy and M. Osman, *Biomed. J. Sci. Tech. Res.*, 2018, **8**, 6659–6662.
 - 273 M. Dreyfuss, E. Härrä, H. Hofmann, H. Kobel, W. Pache and H. Tschertter, *Eur. J. Appl. Microbiol. Biotechnol.*, 1976, **3**, 125–133.
 - 274 E. Uchida, K. Morimoto, N. Kawasaki, Y. Ahmed, A. Said and T. Hayakawa, *Free Radical Res.*, 1997, **27**, 311–323.
 - 275 A. R. Costa, M. E. Rodrigues, M. Henriques, R. Oliveira and J. Azeredo, *Crit. Rev. Biotechnol.*, 2014, **34**, 281–299.
 - 276 M. Erak, K. Bellmann-Sickert, S. Els-Heindl and A. G. Beck-Sickinger, *Bioorg. Med. Chem.*, 2018, **26**, 2759–2765.
 - 277 E. M. Bech, S. L. Pedersen and K. J. Jensen, *ACS Med. Chem. Lett.*, 2018, **9**, 577–580.
 - 278 F. Hubrich, N. M. Bösch, 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.
 - 279 S. Soltani, S. Zirah, S. Rebuffat, F. Couture, Y. Boutin, E. Biron, M. Subirade and I. Fliss, *Front. Microbiol.*, 2022, **12**, 780355.
 - 280 J. D. Hegemann, *ChemBioChem*, 2020, **21**, 7–18.
 - 281 Z. Luo, D. Klein Cerrejon, S. Römer, N. Zoratto and J.-C. Leroux, *Sci. Transl. Med.*, 2023, **15**, eabq1887.
 - 282 S. Jevsevar, M. Kunstelj and V. G. Porekar, *Biotechnol. J.*, 2010, **5**, 113–128.
 - 283 W. R. Strohl, *BioDrugs*, 2015, **29**, 215–239.
 - 284 W. R. Gombotz and D. K. Pettit, *Bioconjugate Chem.*, 1995, **6**, 332–351.
 - 285 P. Dhakane, V. S. Tadimarri and S. Sankaran, *Adv. Funct. Mater.*, 2023, **33**, 2212695.
 - 286 D. K. Chellappan, P. Prasher, V. Saravanan, V. S. Vern Yee, W. C. Wen Chi, J. W. Wong, J. K. Wong, J. T. Wong, W. Wan, J. Chellian, N. Molugulu, S. L. Prabu, R. Ibrahim, T. Darmarajan, M. Candasamy, P. K. Singh, V. Mishra, M. D. Shastri, F. C. Zacconi, A. Chakraborty, M. Mehta, P. K. Gupta, H. Dureja, M. Gulati, S. K. Singh, G. Gupta, N. K. Jha, B. G. George Oliver and K. Dua, *Chem.-Biol. Interact.*, 2022, **351**, 109706.
 - 287 H. R. Costantino, L. Illum, G. Brandt, P. H. Johnson and S. C. Quay, *Int. J. Pharm.*, 2007, **337**, 1–24.



- 288 O. Afzal, A. S. A. Altamimi, M. S. Nadeem, S. I. Alzarea, W. H. Almalki, A. Tariq, B. Mubeen, B. N. Murtaza, S. Iftikhar, N. Riaz and I. Kazmi, *Nanomaterials*, 2022, **12**, 4494.
- 289 L. Chen, W. Hong, W. Ren, T. Xu, Z. Qian and Z. He, *Signal Transduction Targeted Ther.*, 2021, **6**, 1–25.
- 290 M. Vukomanović, V. Žunić, Š. Kunej, B. Jančar, S. Jeverica, R. Podlipec and D. Suvorov, *Sci. Rep.*, 2017, **7**, 4324.
- 291 D. Richter, E. Lakis and J. Piel, *Nat. Chem.*, 2023, **15**, 1422–1430.

